

**The Thermodynamic Properties of the f-Elements and their Compounds. Part 2. The Lanthanide and Actinide Oxides**

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# The Thermodynamic Properties of the *f*-Elements and their Compounds. Part 2. The Lanthanide and Actinide Oxides

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A comprehensive review of the thermodynamic properties of the oxide compounds of the lanthanide and actinide elements is presented. The available literature data for the solid, liquid, and gaseous state have been analysed and recommended values are presented. In case experimental data are missing, estimates have been made based on the trends in the two series, which are extensively discussed. © 2014 Euratom. [<http://dx.doi.org/10.1063/1.4825256>]

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## 1. Introduction

The thermodynamic properties of the  $4f$  (lanthanides) and  $5f$  (actinides) elements and their compounds have been subject of many studies since the Second World War, strongly stimulated by the demands of the nuclear technology. The development of nuclear reactor fuels based on uranium, thorium or plutonium and the understanding of the effects of fission product accumulation in the fuel, a significant fraction of which belongs to the lanthanide group, required such fundamental data. At the same time, many studies of the (thermodynamic) properties of the  $f$ -elements were stimulated by the scientific interest in the role of the  $f$ -elements in the chemical bonding, and particularly the differences between the  $4f$  and  $5f$  series.

Pioneering work on the major actinides has been performed during the Manhattan Project in the USA. The researchers in this project started many systematic studies of uranium and plutonium and its compounds, the results of which became available in literature in the 1950s. At the meetings organised in the frame of the *Peaceful Uses of Atomic Energy* initiative and also at the early *Symposia on Thermodynamics of Nuclear Materials* organised by the *International Atomic Energy Agency* a rapid expansion of the knowledge of the thermodynamic properties of the actinide elements and their compounds was presented. During the same period, the separation methods for the lanthanides, which are difficult due to their chemical similarity, improved significantly to yield these elements in sufficient pure form that was needed for accurate thermochemical and thermophysical measurements.<sup>1,2</sup>

In the 1960s and 1970s a wealth of scientific information on the  $f$ -elements has been published, the lanthanides as well as uranium and thorium being available in pure form to many researchers, and the other actinides being produced in significant quantities for studies at nuclear research laboratories. As a result, the understanding of the trends and systematics of their properties has improved considerably, revealing the differences between the localised  $4f$  electrons in the lanthanides and the heavy actinides (Am-Lr) and the itinerant  $5f$  electrons of the light actinides (Th-Np).

In this work, we will present a comprehensive review of the thermodynamic properties of the oxides of the lanthanides and actinides, based on critical review of the available literature according to procedures described in Secs. 2.1–2.5.

## 2. Approach

### 2.1. Thermal functions of condensed phases

The approach adopted in this review is based on a critical evaluation of the thermal functions (heat capacity, entropy, enthalpy increment) and enthalpies of formation of the lanthanide and actinide oxides using, when possible, the primary experimental data as reported in literature. To describe these as a function of temperature it is necessary to include the data on the structural transformations (including melting point).

The reported transition and melting temperatures have been corrected to the International Temperature Scale ITS-90. Though generally no detailed information is given about the standards used and it is not specifically stated to which earlier temperature scales the data refer, it is assumed that the results between 1948 and 1968 refer to IPTS-48 and between 1969 and 1990 to IPTS-68. Especially in the transition years this may not always be correct. The differences between ITS-90 and IPTS-68 and IPTS-68 and IPTS-48 are shown in Fig. 1.

The low-temperature heat capacity data and the resulting entropies have not been corrected nor refitted. First, the changes in the temperature scales in the range  $T = (0 \text{ to } 300) \text{ K}$  are small, and, second, refitting would only marginally change the results. In case of more than one set of experimental data covering the temperature range from close to 0 K to room temperature, often a motivated choice for one of them is made, based on sample purity and/or calorimetric accuracy. In other cases a joint treatment has been made using overlapping polynomial equations.

The high-temperature heat capacity of the solid phases has been obtained by refitting of the experimental data reported. The following polynomial equation for the enthalpy increment  $\{H^\circ(T) - H^\circ(298.15 \text{ K})\}$  has been adopted

$$\{H^\circ(T) - H^\circ(298.15 \text{ K})\} = \sum A_n(T/K)^n \quad (1)$$

with  $n = -1$  to 2, but in case of anomalous behavior of the heat capacity with  $n$  up to 4. This corresponds to a heat capacity equation of the type

$$C_p(T) = a_{-2}(T/K)^{-2} + \sum a_n(T/K)^n. \quad (2)$$

A difficult question to be answered is that of the temperature correction of high-temperature heat capacity and enthalpy data. As can be seen in Fig. 1 the temperature corrections become significant ( $>2 \text{ K}$ ) above 2000 K. For heat capacity data, which are rare above this temperature, the correction is straightforward, and has been made. However, for enthalpy increment data the corrections are not trivial as they depend on the type of device used and the condition of the sample. In case of the device was calibrated with a known standard (for example, sapphire) the temperature correction should also be made for the standard, and hence not only the temperature but also the enthalpy conversion factor is affected. Also in case of encapsulated samples the temperature correction should lead to an enthalpy correction, as the contribution of the encapsulation material would change. Since the required details are

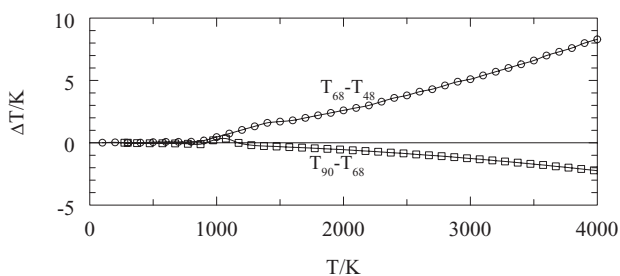


FIG. 1. The temperature corrections according to various temperature scales.

generally lacking to make the corrections properly, the experimental enthalpy data have not been corrected.

All literature data starting from 1945 have been collected systematically. When no experimental data were given in the papers, they have been extracted from digitalized graphs.

## 2.2. Enthalpies of formation of condensed phases

The derivation of the enthalpies of formation from calorimetric data was done by recalculation of the thermochemical reaction schemes (Hess cycles) using a consistent set of auxiliary data, and has been partially reported in earlier work<sup>3</sup> that was updated where necessary. For the solution calorimetric studies the partial molar enthalpy of formation of an acid solution (sln) at the concentration given, has been calculated from the enthalpy of formation of the infinitely dilute acid,<sup>4</sup> the enthalpy of formation of the HX solutions<sup>5</sup> and the densities of the HX solutions at 298.15 K,<sup>6</sup> neglecting the influence of the dissolved ions. Uncertainty limits of the measurements, as listed in the tables or text, are always the values given in the original paper, because in many cases they could not be recalculated due to lack of information. As a consequence they might refer to one standard deviation of the mean, twice the standard deviation of the mean, or the 95% confidence interval, which is not always clear. When combining data from different sources to a selected value, a weighted mean is therefore considered not justified and the uncertainty limit of the selected (mean) value has been estimated. Auxiliary data recommended by CODATA or values consistent with the CODATA selection<sup>4</sup> have been employed.

## 2.3. Thermal functions of gases

Thermal functions of the diatomic molecules were calculated in the present work using the approach developed by Gurvich *et al.*<sup>7</sup> The vibrational-rotational partition functions were calculated by direct summation over vibrational levels and by integration with respect to rotational levels. The upper limit of integration was assumed linearly decreasing with vibrational quantum number.

The electronic partition functions were calculated taking into account all experimentally known and estimated data on excited states. The value of  $Q_{int}$  and its derivatives were evaluated assuming that  $Q_{vib,rot}^{(i)} = (p_i/p_X)^{(X)}_{vib,rot}$ . This approach is well justified in the case of the molecules under consideration though the most of these molecules have numerous low-lying states, which contribute considerably to the thermal functions. The point is that these states as a rule belong to the same electron configuration as the ground state. It is well known that the states of the same configuration have close values of vibrational and rotational constants. Therefore, the tables of molecular constants present the vibrational and rotational constants only for the ground state. The only exception is the YbO molecule for which the ground state configuration gives only one state, namely, the  $X^1\Sigma$  state, while the low-lying states belong to the other configurations with quite different molecular constants. The excited states are presented



in the tables in two blocks for experimentally known and estimated states with corresponding statistical weights. The uncertainties in the energies of experimental states are usually small, but for estimated states they can amount to 10%–20% of the listed values.

For simplification of introducing a huge number of high-energy excited electronic states for molecules under consideration, containing  $f$  and  $d$  open shells, in the present work the approach of the density of states estimation was applied. In this approach, a group of high-energy states with close energies are united in one state with fixed (mostly rounded) energy and average statistical weight. This division into groups of states is being done in a way that does not interfere with the accuracy of the thermal functions calculation. Errors of the calculated thermal functions depend mostly on the accuracy of the data on molecular constants. At room temperature, the uncertainties in heat capacity as a rule do not exceed  $0.3\text{--}0.5\text{ J K}^{-1}\text{ mol}^{-1}$ . At higher temperatures, the uncertainties become larger because of the increasing contribution of excited states and errors of these estimations. In heat capacity the uncertainties can reach  $3\text{--}5\text{ J K}^{-1}\text{ mol}^{-1}$  at 4000 K, or even larger for molecules that entirely lack experimental data, such as PaO, NpO, AmO, CmO, or PuO.

The thermal function of the polyatomic molecules were calculated using the rigid-rotor harmonic oscillator approach,<sup>7,8</sup> that includes general approximations for the translational, rotation and vibrational contributions, and a direct summation of the electronic partition function.

The heat capacity values were approximated by two conjugated equations of the form:

$$C_p^\circ(T) = a_{-2}(T/\text{K})^{-2} + \sum a_n(T/\text{K})^n. \quad (3)$$

The accuracy of the approximation is around  $0.1\text{ J K}^{-1}\text{ mol}^{-1}$  over full temperature range from 298.15 to 4000 K.

The nomenclature used in the tables of molecular properties of the gaseous species is summarized in Table 1. The fundamental constants as recommended by CODATA are used in this work.

## 2.4. Enthalpies of formation of gases

The aim of this review is to select the most reliable enthalpies of formation for MO, MO<sub>2</sub>, and MO<sub>3</sub> molecules

TABLE 1. Nomenclature of the molecular properties for the di- and polyatomic species

Symbol	Name	Symbol	Name
$T_e$	Electronic energy level	$\sigma$	Symmetry number
$p$	Degeneracy (of $T_e$ )	$I_A I_B I_C$	Product of moments of inertia
$\omega_e$	Fundamental harmonic vibrational frequency	$\nu_i$	Fundamental vibrational frequency
$\omega_e x_e$	Anharmonicity correction		
$\beta_e$	Rotational constant		
$\alpha_e$	Rotational-vibration interaction constant		
$D_e$	Centrifugal distortion constant		

based on calculations with new thermal functions of these molecules, and to take into consideration new experimental data and quantum chemical calculations not included in previous assessments.

The recommended enthalpies of formation at the standard temperature  $T = 298.15\text{ K}$  have been selected in this review after critical analysis of all accessible published experimental data on high-temperature equilibria for reactions in the gas phase or in condensed and gas phases. The enthalpies of reactions were calculated from the equilibrium constants by the “second-law” and the “third-law” methods.<sup>7,9</sup> By the former, the enthalpy of reaction at the mean temperature of experiments  $T_{mean}$  is obtained from the temperature dependence of equilibrium constant,  $K_p$ , measured in some temperature interval:

$$\frac{\Delta_r H^\circ(T_{mean})}{R} = -\frac{d(\ln K_p)}{d(1/T)}. \quad (4)$$

The value  $T_{mean}$  is calculated using equation  $T_{mean} = (n^{-1} \sum T_i^{-1})^{-1}$ , where  $n$  is the number of experimental points. The  $\Delta_r H^\circ(T_{mean})$  value so obtained is reduced to the reference temperature 298.15 K:

$$\Delta_r H^\circ(298.15\text{ K}) = \Delta_r H^\circ(T_{mean}) - \Delta_r \{ \Delta_{298.15}^{T_{mean}} H^\circ \}. \quad (5)$$

In the “third-law” method,  $\Delta_r H^\circ(298.15\text{ K})$  is obtained from every experimental  $K_p$  value

$$\Delta_r H^\circ(298.15\text{ K}) = T\Delta_r \phi - RT \ln K_p, \quad (6)$$

using the free energy functions (or reduced Gibbs energies)  $\phi$ , which is defined as

$$\phi = -\{G^\circ(T) - H^\circ(298.15\text{ K})\}/T, \quad (7)$$

$$= S^\circ(T) - \frac{\Delta_{298.15}^T H^\circ}{T}, \quad (8)$$

for each reactant. The enthalpy of formation for the molecule under study,  $\Delta_f H^\circ(298.15\text{ K})$  is calculated from the enthalpy of a reaction using known enthalpies of formation for all other reactants.

The uncertainties ascribed to selected  $\Delta_f H^\circ(298.15\text{ K})$  values reflect statistical errors, uncertainties in the thermal functions, and uncertainties in enthalpies of formation for all other reaction participants. Most experimental measurements for equilibria involving the considered molecules were carried out by Knudsen effusion (KE) and mass spectrometric methods (MS), or by combination of both. In the case of MS measurements, the term  $RT \ln(1.5)$  was added to reflect uncertainties of equilibrium constants calculated from ion currents, due to uncertainties in ionization cross sections. Agreement of the second- and third-law values was regarded as an indication of reliability of experimental data.

## 2.5. Consistency and completeness

We have tried to maintain the internal consistency of the recommended data as much as possible, but in view of the complex interrelationships in some of the analysed systems (e.g., Ce-O, U-O or Pu-O) and the fact that data from other sources have been used this cannot be fully guaranteed. The review has been performed progressively during a period of several years. New information that became available during and just after this period has been incorporated as far as possible, but in some cases the implications of adopting new (more accurate) values were too far-reaching to be implemented. These cases are clearly identified in the text.

## 3. The Lanthanide Oxides in Solid and Liquid State

### 3.1. La<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.1.1. Polymorphism and melting point

Lanthanum(III) oxide has a A-type hexagonal sesquioxide structure (space group  $P\bar{3}m1$ ) at room temperature. It transforms to a H-type hexagonal structure upon heating. Foex and Traverse<sup>10</sup> suggest that this transformation is a simple displacement rearrangement of the lattice,<sup>10</sup> as in  $\alpha \rightarrow \beta$  quartz. Foex and Traverse,<sup>10</sup> Lopato *et al.*<sup>11</sup> and Wehner *et al.*<sup>12</sup> all reported the A  $\rightarrow$  H transition at  $T = 2313$  K, Shevthenko and Lopato<sup>13</sup> at  $T = 2303$  K, and<sup>14</sup> at  $(2319 \pm 5)$  K. The H phase subsequently transforms into a cubic X-type structure (space group  $Im\bar{3}m$ ) and the transformation temperatures were reported as  $T = 2383, 2413, 2363,$  and  $2373$  K,  $(2383 \pm 5)$  K, respectively. Except for the recent work by Ushakov and Navrotsky,<sup>14</sup> all other measurements must be converted to ITS-90. The measurements of Foex and Traverse<sup>10</sup> must be corrected by +14 K, following the procedure outlined by Coutures and Rand.<sup>15</sup> Lopato *et al.*,<sup>11</sup> Wehner *et al.*,<sup>12</sup> and Shevthenko and Lopato<sup>13</sup> reported no (detailed) information on the calibration of their measurements, but assuming the data refer to IPTS-68, a correction of  $-1$  K needs to be applied. We select  $T_{rs} = (2313 \pm 30)$  K for the A  $\rightarrow$  H transformation,  $T_{rs} = (2383 \pm 30)$  K for the H  $\rightarrow$  X transformation.

Various measurements of the melting temperature of solid La<sub>2</sub>O<sub>3</sub> have been reported as summarized in Table 2, which is based on the IUPAC review by Coutures and Rand;<sup>15</sup> the results being corrected to ITS-90. The recent value for the melting by temperature by Ushakov and Navrotsky<sup>14</sup> is in excellent agreement with the selected values by Coutures and Rand,<sup>15</sup> and the latter is retained,  $T_{fus} = (2577 \pm 15)$  K.

#### 3.1.2. Heat capacity and entropy

Low-temperature heat capacity measurements for La<sub>2</sub>O<sub>3</sub> have been reported by three different research groups: Goldstein *et al.*<sup>26</sup> from 16 to 300 K, King *et al.*<sup>27</sup> from 50 to 300 K, and Justice and Westrum, Jr.<sup>28</sup> from 5 to 350 K. The measurements reasonably agree to about  $T = 200$  K; above this

TABLE 2. Temperature of melting of lanthanum sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Wartenberg and Reusch <sup>16</sup>	2588	2581
Lambertson and Gunzel <sup>17</sup>	$2483 \pm 20$	$2489 \pm 20$
Sata and Kiyoura <sup>18</sup>	$2577 \pm 2$	$2583 \pm 2$
Foex <sup>19</sup>	2573	2587
Mordovin <i>et al.</i> <sup>20</sup>	$2493 \pm 30$	$2496 \pm 30$
Noguchi and Mizuno <sup>21</sup>	$2530 \pm 20$	$2532 \pm 20$
Treswjatskii <i>et al.</i> <sup>22a</sup>	$2583 \pm 20$	$2582 \pm 20$
Coutures <i>et al.</i> <sup>23</sup>	$2593 \pm 10$	$2592 \pm 10$
Wehner <i>et al.</i> <sup>12</sup>	$2563 \pm 30$	
Mizuno <i>et al.</i> <sup>24</sup>	$2569 \pm 20$	$2555 \pm 20$
Yoshimura <i>et al.</i> <sup>25</sup>	$2573 \pm 5$	$2576 \pm 5$
Shevthenko and Lopato <sup>13</sup>	2583	2582
Ushakov and Navrotsky <sup>14</sup>		$2574 \pm 10$
Selected value:		$2577 \pm 15$

<sup>a</sup>Also reported by Lopato *et al.*<sup>11</sup>

temperature the differences significantly increase. The selected standard entropy of La<sub>2</sub>O<sub>3</sub> has been derived from the measurement by Justice and Westrum, Jr.,<sup>28</sup> which is considered to be the most accurate

$$S^\circ(298.15 \text{ K}) = (127.32 \pm 0.84) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of La<sub>2</sub>O<sub>3</sub>(cr) has been measured by Blomeke and Ziegler<sup>29</sup> from 380 to 1170 K, King *et al.*<sup>27</sup> from 399 K to 1797 K, Yashvili *et al.*<sup>30</sup> from 380 to 1650 K and Sedmidubský *et al.*<sup>31</sup> from 689 to 1291 K, which are in perfect agreement, as shown in Fig. 2. The measurements smoothly join the low-temperature heat capacity measurements by Justice and Westrum, Jr.<sup>28</sup> Basili *et al.*<sup>32</sup> measured the heat capacity of La<sub>2</sub>O<sub>3</sub> from 400 to 850 K. Their results, only presented in graphical form, reasonably agree with the enthalpy measurements to about 550 K. The heat capacity above 298.15 K for A-type La<sub>2</sub>O<sub>3</sub> can be represented by the equation (298.15 to 1800 K):

$$C_p^\circ(T)/\text{J K}^{-1} \text{ mol}^{-1} = 120.6805 + 13.42414 \times 10^{-3}(T/\text{K}) - 14.13668 \times 10^5(T/\text{K})^{-2}$$

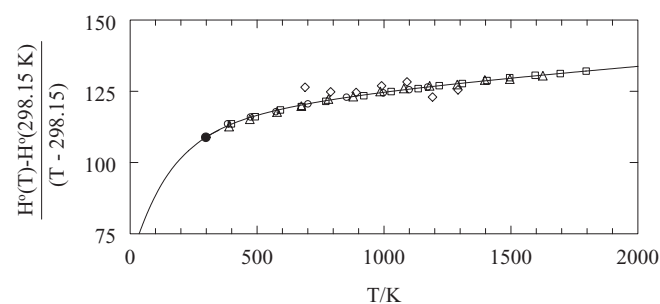


FIG. 2. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of La<sub>2</sub>O<sub>3</sub>;  $\circ$ , Blomeke and Ziegler<sup>29</sup>;  $\square$ , Yashvili *et al.*<sup>30</sup>;  $\triangle$ , King *et al.*<sup>27</sup>;  $\diamond$ , Sedmidubský *et al.*<sup>31</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice and Westrum, Jr.,<sup>28</sup> the curve shows the recommended equation.

derived from a fit of the combined enthalpy results of Bloemeke and Ziegler,<sup>29</sup> King *et al.*<sup>27</sup> and Yashvili *et al.*,<sup>30</sup> which are considered the most accurate. The boundary condition  $C_p^\circ(298.15) = 108.78 \text{ J K}^{-1} \text{ mol}^{-1}$  was applied, as derived from the low-temperature heat capacity measurements.<sup>28</sup>

Heat-capacity or enthalpy measurements have not been reported for the H, X, and liquid phases of  $\text{La}_2\text{O}_3$  and we estimate

$$C_p^\circ(\text{H}, T) = C_p^\circ(\text{X}, T) = 150 \text{ J K}^{-1} \text{ mol}^{-1},$$

$$C_p^\circ(\text{liq}, T) = 162 \text{ J K}^{-1} \text{ mol}^{-1}.$$

The transition enthalpies of  $\text{La}_2\text{O}_3$  have been measured by Ushakov and Navrotsky<sup>14</sup> recently, using high temperature thermal analysis

$$\Delta_{\text{trs}}H^\circ(\text{A} \rightarrow \text{H}) = (23 \pm 5) \text{ kJ mol}^{-1},$$

$$\Delta_{\text{trs}}H^\circ(\text{H} \rightarrow \text{X}) = (17 \pm 5) \text{ kJ mol}^{-1},$$

$$\Delta_{\text{fus}}H^\circ = (78 \pm 10) \text{ kJ mol}^{-1}.$$

These values have been selected here. They are only partially corresponding to the observation by Foex and Traverse<sup>10</sup> who noted a moderate thermal effect for the  $\text{A} \rightarrow \text{H}$  transformation by DTA (differential thermal analysis), and a significant one for the  $\text{H} \rightarrow \text{X}$ . Wu and Pelton<sup>33</sup> concluded from the fact that the liquidus at the  $\text{La}_2\text{O}_3$  side of the phase  $\text{La}_2\text{O}_3\text{-Al}_2\text{O}_3$  phase diagram does not show clear discontinuities, that the entropy changes of the  $\text{A} \rightarrow \text{H}$  and  $\text{H} \rightarrow \text{X}$  transformations are very small. They also concluded that the limiting slope of the liquid line in this phase diagram suggests an entropy of fusion

of  $25.1 \text{ J K}^{-1} \text{ mol}^{-1}$ , in fair agreement with the value found by Ushakov and Navrotsky,<sup>14</sup>  $30.3 \text{ J K}^{-1} \text{ mol}^{-1}$ .

### 3.1.3. Enthalpy of formation

The enthalpy of formation of  $\text{La}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected value from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{La}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1791.6 \pm 2.0) \text{ kJ mol}^{-1}.$$

All values relevant to the derivation of the standard enthalpy of formation of lanthanum sesquioxide are summarized in Table 3. Huber, Jr. and Holley, Jr.<sup>35</sup> determined the enthalpy of formation by combustion of a very pure sample of metal. This value has been confirmed by several authors using solution calorimetry.<sup>36–38</sup> However, the values for the enthalpy of solution of  $\text{La}(\text{cr})$  differ significantly.<sup>37–39</sup> As discussed by Cordfunke and Konings,<sup>34</sup> the results of Merli *et al.*<sup>39</sup> can be considered as the most accurate since they made their measurements on a well-defined sample. Therefore, the results of the other studies were recalculated using the values from this study, some obtained by inter- or extrapolation. The resulting enthalpies of formation are in excellent agreement with the combustion value and the selected value is the mean of the combustion value by Huber, Jr. and Holley, Jr.,<sup>35</sup> and the recalculated values obtained from and the enthalpy of solution measurements Montgomery and Hubert,<sup>36</sup> Fitzgibbon *et al.*,<sup>37</sup> and Gvelesiani and Yashvili.<sup>38</sup>

TABLE 3. The enthalpy of formation of  $\text{La}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  is the enthalpy of solution of  $\text{La}(\text{cr})$ ,  $\Delta H_2^\circ$  of  $\text{La}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Muthmann and Weis <sup>40</sup>	C			-1857.7
Matignon <sup>41</sup>	S			-1789.0
Kremers and Stevens <sup>42</sup>	C			-1912.1
Moose and Parr <sup>43</sup>	C			-1907.1
Beck <sup>44</sup>	S		-439.3	
Roth <i>et al.</i> <sup>45</sup>	C			-2255 ± 17
Huber, Jr. and Holley, Jr. <sup>35</sup>	C			-1793.1 ± 0.8
von Wartenberg <sup>46</sup>	S (0.1)		-468.6 ± 6.3	
Montgomery and Hubert <sup>36</sup>	S (0.51)	[-704.1 ± 1.2] <sup>b</sup>	-474.4 ± 1.6	-1791.3 ± 2.5
Fitzgibbon <i>et al.</i> <sup>37</sup>	S (1.0)	-705.5 ± 1.3	-474.4 ± 0.4	-1794.2 ± 2.7
		[-704.4 ± 1.2] <sup>c</sup>		-1792.0 ± 2.7
	S(1.0)	-705.6 ± 1.3	-473.8 ± 0.4	-1794.8 ± 2.7
		[-704.4 ± 1.2] <sup>c</sup>		-1792.5 ± 2.7
Gvelesiani and Yashvili <sup>38</sup>	S (1.0)	-708.0 ± 2.0	-475.3 ± 3.3	-1798.2 ± 5.2
		[-704.4 ± 1.2] <sup>c</sup>		-1791.0 ± 4.1
	S (1.5)	-708.8 ± 2.9	-475.3 ± 1.8	-1799.9 ± 6.1
		[-704.7 ± 1.2] <sup>b</sup>		-1791.7 ± 3.0
Oppermann <i>et al.</i> <sup>47</sup>	S (4.0)	[-706.2 ± 1.1] <sup>b</sup>	-472.6 ± 0.3	-1798.2 ± 2.4
Selected value:				-1791.6 ± 2.0

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ .

<sup>b</sup>Estimated/interpolated from the results of Merli *et al.*<sup>39</sup>

<sup>c</sup>Merli *et al.*<sup>39</sup>

### 3.2. CeO<sub>2</sub>(cr,l)

#### 3.2.1. Melting point

Cerium dioxide has a cubic fluorite structure (space group Fm $\bar{3}$ m) up to the melting point. The reported values for the temperature of melting are very dissimilar (Table 4), which is due to the fact that this compound starts to lose oxygen at elevated temperatures to form a substoichiometric CeO<sub>2-x</sub> phase. The melting point for CeO<sub>2</sub> strongly depends on the atmosphere under which the liquid phase is produced.<sup>48</sup> For example, Mordovin *et al.*<sup>20</sup> detected the liquid phase already at 2670 K, when heating ceria in an argon atmosphere. On the other hand many other authors<sup>10,49-51</sup> observed higher values of the melting point while heating CeO<sub>2</sub> under a strongly oxidising atmosphere (pure pressurised O<sub>2</sub>, air, or a mixture of oxygen and an inert gas at high pressure): between 2753 and 3073 K. Also Manara *et al.*,<sup>52</sup> in a recent unpublished investigation of the melting behavior of CeO<sub>2</sub>, performed measurements both under oxidising and reducing atmospheres, obtaining 2743 and 2675 K, respectively. In this last study, however, heating under high oxygen pressures could not be realised. Since the highest values are the most likely to correspond to quasistoichiometric CeO<sub>2</sub>, we select  $T_{fus} = (3083 \pm 50)$  K as the best melting point for stoichiometric cerium dioxide, the uncertainty being assigned by us. This is in line with the suggestion of Du *et al.*<sup>53</sup> that the melting point of CeO<sub>2</sub> must be between those of the group IVB dioxides and the actinide dioxides.

#### 3.2.2. Heat capacity and entropy

The low-temperature heat capacity of CeO<sub>2</sub> has been measured by Westrum, Jr. and Beale, Jr.<sup>56</sup> from 5 to 300 K. The entropy, as derived from these measurements, is adopted here as

$$S^\circ(298.15 \text{ K}) = (62.29 \pm 0.07) \text{ J K}^{-1} \text{ mol}^{-1}.$$

High-temperature enthalpy increments have been reported by Kuznetsov *et al.*<sup>57</sup> in the temperature range from 608 to 1172 K, King *et al.*<sup>27</sup> from 400 to 1800 K (only smoothed values are given in the paper), Pears *et al.*<sup>58</sup> from 640 to 2044 K, Mezaki *et al.*<sup>59</sup> from 490 to 1140 K, and Yashvili *et al.*<sup>60</sup>

TABLE 4. Temperature of melting of cerium dioxide

Authors	$T_{fus}/\text{K}$	
	Reported	ITS-90
Ruff <sup>54</sup>	2246	
von Wartenberg and Gurr <sup>55</sup>	2873	
Trombe <sup>50a</sup>	3073	3077
Tshierpanov and Trjesvyatsky <sup>49b</sup>	3083	3087
Foex and Traverse <sup>10</sup>	2753	2768
Mordovin <i>et al.</i> <sup>20</sup>	2670 ± 30	2673 ± 30
Watson <sup>51</sup>	2873	2872
Selected value:		3083 ± 50

<sup>a</sup>Paper not available to us, cited from Noguchi and Mizuno.<sup>21</sup>

<sup>b</sup>Paper not available to us, cited from Du *et al.*<sup>53</sup>

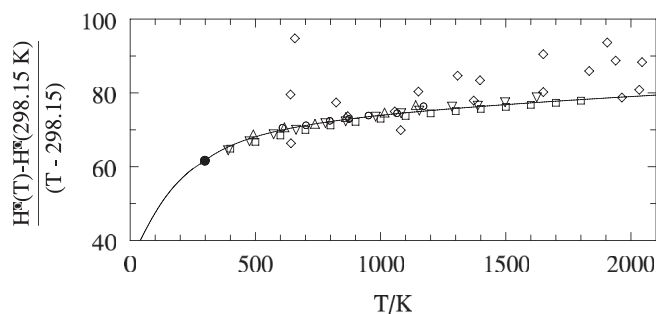


FIG. 3. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of CeO<sub>2</sub>;  $\circ$ , Kuznetsov *et al.*<sup>57</sup>;  $\square$ , King *et al.*<sup>27</sup>;  $\triangle$ , Mezaki *et al.*<sup>59</sup>;  $\nabla$ , Yashvili *et al.*<sup>60</sup>;  $\diamond$ , Pears *et al.*<sup>58</sup>;  $\bullet$ , value derived from the low-temperature measurements by Westrum, Jr. and Beale, Jr.,<sup>56</sup> the curve shows the recommended equation.

from 391 to 1624 K. These data are in reasonable agreement, as shown in Fig. 3.

High-temperature heat capacities have been measured by Riess *et al.*<sup>61</sup> by adiabatic scanning calorimetry (350–900 K). The results, presented as an equation only, are in excellent agreement with the values derived from the enthalpy increment measurements. Gallagher and Dworzak<sup>62</sup> determined the heat capacity between 418 and 758 K by DSC (differential scanning calorimetry). These data are, however, much too low to join the low-temperature data by Westrum, Jr. and Beale, Jr.<sup>56</sup>

The enthalpy data have been combined, and constrained  $C_p^\circ(298.15 \text{ K}) = 61.63 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low temperature data,<sup>56</sup> resulting in the following heat capacity equation:

$$C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = 74.4814 + 5.83682 \times 10^{-3} (T/\text{K}) - 1.29710 \times 10^6 (T/\text{K})^{-2}.$$

This equation is extrapolated to the melting point, which might neglect possible anomalous increase in the heat capacity as observed in other fcc dioxides of *f*-elements such as ThO<sub>2</sub> and UO<sub>2</sub>. The heat capacity of the liquid has been estimated as

$$C_p^\circ(\text{CeO}_2, \text{liq}, T) = 120 \text{ J K}^{-1} \text{ mol}^{-1}.$$

The entropy of fusion is assumed to be the same as that of the isostructural UO<sub>2</sub> phase ( $22.4 \text{ J K}^{-1} \text{ mol}^{-1}$ ), which is the only fcc dioxide for which this quantity is well defined. We thus obtain for the enthalpy of fusion

$$\Delta_{fus} H^\circ(\text{CeO}_2) = (69 \pm 5) \text{ kJ mol}^{-1}.$$

#### 3.2.3. Enthalpy of formation

The standard molar enthalpy of formation of CeO<sub>2</sub>(cr) has been determined by Huber, Jr. and Holley, Jr.<sup>35</sup> by oxygen-bomb combustion calorimetry using a well-analyzed sample of cerium metal, giving  $\Delta_f H^\circ(298.15 \text{ K}) = -(1088.6 \pm 1.4) \text{ kJ mol}^{-1}$ . This value, which was carefully corrected for impurities, is in excellent agreement with the result by Baker *et al.*<sup>63</sup> of a later oxygen-bomb combustion calorimetric investigation carried out in the same laboratory,



$\Delta_f H^\circ(298.15 \text{ K}) = -(1090.4 \pm 0.8) \text{ kJ mol}^{-1}$ . We have selected the latter value because it is based on a cerium metal sample of significantly higher purity

$$\Delta_f H^\circ(\text{CeO}_2, \text{cr}, 298.15 \text{ K}) = -(1090.4 \pm 1.0) \text{ kJ mol}^{-1}.$$

The results of early investigations<sup>40,43,64</sup> are mainly of historical interest due to a poor quality of materials and experimental techniques available at that time.

### 3.3. Ce<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.3.1. Polymorphism and melting point

At room temperature, cerium sesquioxide has a hexagonal A-type structure (space group P $\bar{3}m1$ ). Pankratz and Kelly<sup>65</sup> observed a transition at 1050 K from the A-type sesquioxide to a high-temperature modification but it should be noted that their sample had the hyperstoichiometric composition (Ce<sub>2</sub>O<sub>3.33</sub>). This transition has not been reported by other investigators, and does not fit into the general structure diagram on the lanthanide sesquioxides (Fig. 4). We therefore attribute it to this specific composition and not to the sesquioxide phase, and according to the phase diagram of the Ce-O system it could represent the phase boundary of the {Ce<sub>2</sub>O<sub>3</sub>+Ce<sub>6</sub>O<sub>10</sub>} two-phase field.

Lopato *et al.*<sup>11</sup> and Shevthenko and Lopato<sup>13</sup> reported that A-type Ce<sub>2</sub>O<sub>3</sub> transforms to a hexagonal H-type structure at 2393 K. They also found that the H-type phase transforms to a cubic X-type structure at 2407 K. No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of  $-1 \text{ K}$  needs to be applied for ITS-90. We select  $T_{irs} = (2392 \pm 30) \text{ K}$  for the A  $\rightarrow$  H transformation,  $T_{irs} = (2406 \pm 30) \text{ K}$  for the H  $\rightarrow$  X transformation.

The various data for the melting temperature have been summarized in Table 5, which is based on the IUPAC review by Coutures and Rand;<sup>15</sup> the results being corrected to ITS-90. The selected melting point is  $(2512 \pm 50) \text{ K}$ .

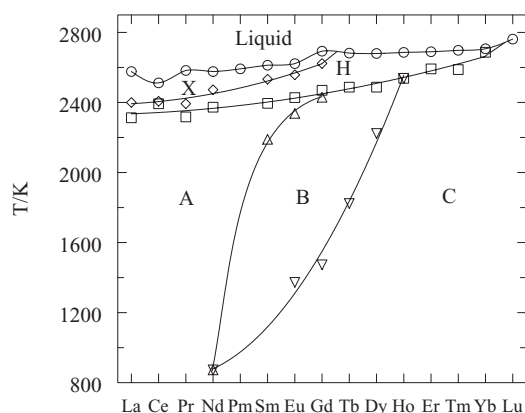


FIG. 4. The polymorphism in the Ln<sub>2</sub>O<sub>3</sub> series as a function of temperature.

TABLE 5. Temperature of melting of cerium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/\text{K}$	
	Reported	ITS-90
Sata and Kiyoura <sup>18</sup>	2483 $\pm$ 10	2489 $\pm$ 2
Mordovin <i>et al.</i> <sup>20</sup>	2415 $\pm$ 30	2429 $\pm$ 30
Treswjatskii <i>et al.</i> <sup>22</sup>	2513 $\pm$ 20	2512 $\pm$ 20
Shevthenko and Lopato <sup>13</sup>	2513	2512
Selected value:		2512 $\pm$ 15

#### 3.3.2. Heat capacity and entropy

Low-temperature heat capacity measurements on cerium sesquioxide have been reported by Weller and King<sup>66</sup> in the range from 53 to  $-296 \text{ K}$ , Justice and Westrum, Jr.<sup>67</sup> in the range from 5 to 345 K, and Huntelaar *et al.*<sup>68</sup> from 5 to 400 K. The data by Weller and King cover a smaller temperature range and, moreover, were obtained using a sample with composition Ce<sub>2</sub>O<sub>3.33</sub>. The sample of Justice and Westrum, Jr.<sup>67</sup> was reported to refer to the composition (Ce<sub>2</sub>O<sub>3.02</sub>) but this was later doubted by the authors<sup>69</sup> who suggested that the true composition was likely Ce<sub>2</sub>O<sub>3.33</sub> also. For that reason the data by Huntelaar *et al.*<sup>68</sup> are preferred. Hence, the standard entropy, as derived by the latter authors, has been adopted here as

$$S^\circ(298.15 \text{ K}) = (148.1 \pm 0.4) \text{ J K}^{-1} \text{ mol}^{-1}.$$

High-temperature enthalpy increment measurements have been reported by Pankratz and Kelly<sup>65</sup> from 398 to 1501 K, Kuznetsov *et al.*<sup>70</sup> from 578 to 1116 K and Huntelaar *et al.*<sup>68</sup> from 473 to 883 K. The data of Pankratz and Kelly<sup>65</sup> refer to the nonstoichiometric composition (Ce<sub>2</sub>O<sub>3.33</sub>). The authors corrected them for the presence of CeO<sub>2</sub>, but in view of the unexplained phase transition observed in the results (see above), the correctness of this approach may be doubted. The results of Kuznetsov *et al.*<sup>70</sup> and Huntelaar *et al.*<sup>68</sup> are in reasonable agreement and fit the low-temperature data. However, the former results also indicate an anomalous increase near 1000 K, which might indicate that their sample had not the stoichiometric composition and that the agreement is fortuitous. For that reason our selected heat capacity equation is based on the results of Huntelaar *et al.*<sup>68</sup> only, which was constrained to  $C_p^\circ(298.15) = 115.0$ , as derived from the low-temperature heat capacity measurements by the same authors,

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 113.736 + 28.4344 \times 10^{-3}(T/\text{K}) - 0.641205 \times 10^6(T/\text{K})^{-2}.$$

Basili *et al.*<sup>32</sup> determined the heat capacity of Ce<sub>2</sub>O<sub>3</sub> between 440 and 1100 K by means of a plane temperature waves method. Venkata Krishnan and Nagarajan<sup>71</sup> measured the heat capacity from 280 to 820 K by DSC. Their data, read from a graph, are in reasonable agreement with the heat capacity derived from the enthalpy increments.

The enthalpies of transition between the high-temperature modifications have been estimated, in absence of experimental

data. For the (A → H) we have assumed that the entropy of transition varies regularly between the known values for La<sub>2</sub>O<sub>3</sub> and Gd<sub>2</sub>O<sub>3</sub>. For the (H → X) and (X → liquid) transitions we have assumed the entropies to be the same as for La<sub>2</sub>O<sub>3</sub>:

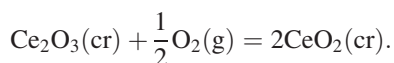
$$\begin{aligned}\Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (28 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (19 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (85 \pm 10) \text{ kJ mol}^{-1}.\end{aligned}$$

For the heat capacity of the high temperature phases we estimate

$$\begin{aligned}C_p^\circ(\text{H}, \text{T}) &= C_p^\circ(\text{X}, \text{T}) = 145 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{liq}, \text{T}) &= 157 \text{ J K}^{-1} \text{ mol}^{-1}.\end{aligned}$$

### 3.3.3. Enthalpy of formation

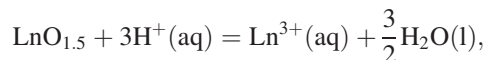
Several combustion calorimetric studies have been reported for the reaction



The results for the enthalpy of reaction are discordant, as summarized in Table 6. As suggested by Cordfunke and Konings,<sup>34</sup> this variation may be due to (i) differences in the O/M ratio of the starting material Ce<sub>2</sub>O<sub>3</sub>(cr), (ii) impurities in Ce<sub>2</sub>O<sub>3</sub>(cr) resulting from the fabrication by reduction of the dioxide (e.g., residual carbon has a big impact on the combustion values), and (iii) differences in the final state of the reaction product CeO<sub>2</sub> that is known to have a large range of substoichiometric compositions.

Huntelaar *et al.*<sup>68</sup> measured the enthalpy of solution of a well-defined sample of Ce<sub>2</sub>O<sub>3</sub>(cr) in 0.25 mol dm<sup>-3</sup> HCl(aq) from which the enthalpy of formation is derived as Δ<sub>f</sub>H<sup>o</sup>(298.15 K) = -(1813.1 ± 0.8) kJ mol<sup>-1</sup>. Putnam *et al.*<sup>74</sup> measured the enthalpy of formation by high-temperature

oxide-melt solution calorimetry. Recalculating their measurements with the selected value for the enthalpy of formation of CeO<sub>2</sub>, gives Δ<sub>f</sub>H<sup>o</sup>(298.15 K) = -(1809.2 ± 5.2) kJ mol<sup>-1</sup>. As noted by Morss and Konings<sup>76</sup> these results give an anomalous value for the hypothetical solution enthalpy of the reaction,



compared to the other lanthanide sesquioxides. For that reason we select the enthalpy of formation from the combustion study by Baker and Holley, Jr.<sup>73</sup> that was performed with CeO<sub>1.5+x</sub> samples that were carefully analysed and were extrapolated to CeO<sub>1.500</sub>:

$$\Delta_f H^\circ(\text{Ce}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1799.8 \pm 1.8) \text{ kJ mol}^{-1}.$$

This value is different from the one selected by Cordfunke and Konings.<sup>34</sup>

## 3.4. PrO<sub>2</sub>(cr,l)

### 3.4.1. Structure

PrO<sub>2</sub> has a face-centered cubic structure (space group Fm $\bar{3}$ m). Hyde *et al.*<sup>77</sup> showed that PrO<sub>2</sub> is stable up to 587 K and Ushakov and Navtrosky<sup>78</sup> up to 663 K in oxygen gas. Above this temperature it decomposes to Pr<sub>6</sub>O<sub>11</sub>.

### 3.4.2. Heat capacity and entropy

The heat capacity of PrO<sub>2</sub> has been measured by Gardiner *et al.*<sup>79</sup> from 2.4 to 23 K, showing a lambda anomaly due to the antiferromagnetic ordering at ~13.5 K. No measurements of the heat capacity of PrO<sub>2</sub> in a wider temperature range have been made from which the standard entropy could be derived. For that reason the standard entropy has been estimated from the systematics in the lanthanide and actinide oxides as

$$S^\circ(298.15 \text{ K}) = (80.8 \pm 2.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

TABLE 6. The enthalpy of formation of Ce<sub>2</sub>O<sub>3</sub>(cr) at 298.15 K; ΔH<sub>1</sub><sup>o</sup> and ΔH<sub>2</sub><sup>o</sup> are the enthalpies of solution of Ce(cr) and Ce<sub>2</sub>O<sub>3</sub>(cr) in HCl(aq), respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	ΔH <sub>1</sub> <sup>o</sup> /kJ mol <sup>-1</sup>	ΔH <sub>2</sub> <sup>o</sup> /kJ mol <sup>-1</sup>	Δ <sub>f</sub> H <sup>o</sup> /kJ mol <sup>-1</sup>
Kuznetsov <i>et al.</i> <sup>57</sup>	C			-1823.4 ± 1.8 <sup>b</sup>
Mah <sup>72</sup>	C			-1790.2 ± 1.5 <sup>c</sup>
Baker and Holley, Jr. <sup>73</sup>	C			-1799.8 ± 1.8 <sup>d</sup>
Huntelaar <i>et al.</i> <sup>68</sup>	S (0.25)	[-699.2 ± 0.2] <sup>e</sup>	-442.7 ± 0.6	-1813.1 ± 0.8 -1813.2 ± 3.2 <sup>f</sup>
Putnam <i>et al.</i> <sup>74</sup>	H			-1809.2 ± 5.2
Selected value:				-1813.0 ± 2.0

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in mol dm<sup>-3</sup>; H = high temperature oxide melt solution calorimetry.

<sup>b</sup>For the enthalpy of the reaction Ce<sub>2</sub>O<sub>3</sub>(cr) + ½ O<sub>2</sub>(g) = 2CeO<sub>2</sub>(cr) the following value was used: -357.4 ± 1.1 kJ mol<sup>-1</sup>.

<sup>c</sup>*idem*, -390.6 ± 0.4 kJ mol<sup>-1</sup>.

<sup>d</sup>*idem*, -381.0 ± 0.7 kJ mol<sup>-1</sup>.

<sup>e</sup>Spedding and Miller.<sup>75</sup>

<sup>f</sup>Cycle based on CeCl<sub>3</sub>.

In a similar manner the high temperature heat capacity for the temperature range from 298.15 to 600 K was estimated as

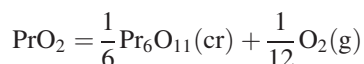
$$C_p^{\circ}/(\text{JK}^{-1}\text{mol}^{-1}) = 72.981 + 16.628 \times 10^{-3}(\text{T/K}) - 0.9990 \times 10^6(\text{T/K})^{-2}.$$

### 3.4.3. Enthalpy of formation

The enthalpy of formation of  $\text{PrO}_2$  was determined by Eyring *et al.*<sup>80</sup> and Gramsch and Morss<sup>81</sup> using solution calorimetry in nitric acid. Recalculation of the results yields  $-(949.3 \pm 4.3)$  kJ mol<sup>-1</sup> and  $-(959.1 \pm 2.3)$  kJ mol<sup>-1</sup>, respectively, being significantly different, entirely due to differences in the solution enthalpies. The selected enthalpy of formation is derived from the work by Gramsch and Morss,<sup>81</sup> as it is based on an excellently characterised sample

$$\Delta_f H^{\circ}(\text{PrO}_2, \text{cr}, 298.15 \text{ K}) = -(959.1 \pm 2.3) \text{ kJ mol}^{-1}.$$

Ushakov and Navtrosky<sup>78</sup> measured the enthalpy of the reaction



by DSC obtaining  $\Delta_r H^{\circ}(298.15 \text{ K}) = (10 \pm 2)$  kJ mol<sup>-1</sup> at 663 K. Using the enthalpies of formation of  $\text{PrO}_2$  and  $\text{PrO}_{1.833}$  selected in this work, we calculate  $\Delta_r H^{\circ}(298.15 \text{ K}) = (9.6 \pm 5.8)$  kJ mol<sup>-1</sup>, in excellent agreement.

## 3.5. $\text{PrO}_{1.833}(\text{cr,l})$

### 3.5.1. Melting point

$\text{PrO}_{1.833}$  (also designated as  $\text{Pr}_6\text{O}_{11}$ ) has a triclinic crystal structure (space group  $\text{P}\bar{1}$ ). Pankratz<sup>82</sup> found a phase transition around 760 K. However, it can be concluded from the  $\text{PrO}_{1.5}$ – $\text{PrO}_2$  phase diagram proposed by Turcotte *et al.*<sup>83</sup> that the upper limit of stability of  $\text{PrO}_{1.833}$  is about 750 K; above this temperature the  $\text{PrO}_{2-x}$  phase is stable in equilibrium with  $\text{O}_2(\text{g})$ .

Mordovin *et al.*<sup>20</sup> reported the melting point of  $\text{PrO}_{1.833}$  as  $T = (2315 \pm 30)$  K in Ar atmosphere, but mentioned that

substantial dissociation of the sample occurred. The relevance of this measurement must thus be doubted.

### 3.5.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{PrO}_{1.833}$  has not been measured. The selected standard entropy is interpolated between  $\text{PrO}_2(\text{cr})$  and  $\text{PrO}_{1.5}$ :

$$S^{\circ}(298.15 \text{ K}) = (79.2 \pm 2.0) \text{ JK}^{-1} \text{ mol}^{-1}$$

The high-temperature enthalpy increment of  $\text{PrO}_{1.833}$  was measured by Pankratz<sup>82</sup> from 398 to 1052 K and by Blomeke and Ziegler<sup>29</sup> from 383 to 1172 K. The data are in good agreement at low temperatures, but the difference increases steadily up to 1033 K where Pankratz<sup>82</sup> observed a phase transformation, whereas Blomeke and Ziegler<sup>29</sup> did not. As discussed above this phase transformation is very likely the peritectic decomposition. Since the sample of Blomeke and Ziegler<sup>29</sup> was heated in a capsule above this temperature before the measurement, (partial) decomposition probably has occurred, and their results might refer to the  $\text{PrO}_{2-x}$  phase. For that reason the data of Pankratz<sup>82</sup> below 760 K have been fitted to a polynomial equation, yielding for the heat capacity:

$$C_p^{\circ}/(\text{JK}^{-1}\text{mol}^{-1}) = 68.4932 + 15.9207 \times 10^{-3}(\text{T/K}) - 0.80968 \times 10^6(\text{T/K})^{-2}.$$

### 3.5.3. Enthalpy of formation

The enthalpy of formation of  $\text{PrO}_{1.833}$  was measured by Stubblefield *et al.*<sup>84</sup> by solution calorimetry and by Fitzgibbon *et al.*<sup>85</sup> using both solution and combustion calorimetry. The recalculated values are summarized in Table 7, based on a Hess cycle with  $\text{Pr}_2\text{O}_3$ . The results are in reasonable agreement and we select

$$\Delta_f H^{\circ}(\text{PrO}_{1.833}, \text{cr}, 298.15 \text{ K}) = -(944.6 \pm 2.5) \text{ kJ mol}^{-1}.$$

TABLE 7. The enthalpy of formation of  $\text{PrO}_{1.833}$  and  $\text{Pr}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^{\circ}$  and  $\Delta H_2^{\circ}$  are the enthalpies of solution of  $\text{Pr}(\text{cr})$  and  $\text{PrO}_{1.833}(\text{cr})$  in  $\text{HNO}_3(\text{aq})$ ,  $\text{Pr}(\text{cr})$  and  $\text{Pr}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$  or  $\text{HNO}_3(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^{\circ}/\text{kJ mol}^{-1}$	$\Delta H_2^{\circ}/\text{kJ mol}^{-1}$	$\Delta_f H^{\circ}/\text{kJ mol}^{-1}$
$\text{PrO}_{1.833}$				
Stubblefield <i>et al.</i> <sup>84</sup>	S (6.0)		$-183.3 \pm 0.4$	$-943.5 \pm 2.1$
Fitzgibbon <i>et al.</i> <sup>85</sup>	S (6.0)		$-179.6 \pm 2.0$	$-947.0 \pm 2.3$
	S (6.0)		$-179.9 \pm 2.0$	$-947.0 \pm 3.2$
	C			$-943.2 \pm 0.8$
Selected value:				$-944.6 \pm 2.5$
$\text{Pr}_2\text{O}_3$				
Stubblefield <i>et al.</i> <sup>84</sup>	S (6.0) <sup>b</sup>	$[-1020.9 \pm 3.4]^c$	$-447.7 \pm 0.8$	$-1831.6 \pm 3.5$
Fitzgibbon <i>et al.</i> <sup>85</sup>	S (2.0)	$-692.2 \pm 1.3$	$-432.0 \pm 1.4$	$-1809.9 \pm 3.0$
Selected value:				$-1809.9 \pm 3.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in mol dm<sup>-3</sup>

<sup>b</sup>The enthalpy of solution in  $\text{HNO}_3(\text{aq})$ .

<sup>c</sup>The enthalpy of solution of Pr in 6.0 mol dm<sup>-3</sup>  $\text{HNO}_3(\text{aq})$  is from Ref. 86.

TABLE 8. The enthalpy of formation of phases in the PrO<sub>2</sub>-PrO<sub>1.5</sub> system

Compound	$\Delta_f H^\circ(298.15 \text{ K})/\text{kJ mol}^{-1}$	Authors
PrO <sub>1.703</sub>	$-(926.3 \pm 2.2)$	Stubblefield <i>et al.</i> <sup>84</sup>
PrO <sub>1.717</sub>	$-(928.4 \pm 2.6)$	Fitzgibbon <i>et al.</i> <sup>85</sup>
PrO <sub>1.804</sub>	$-(938.9 \pm 2.0)$	Fitzgibbon <i>et al.</i> <sup>85</sup>

The enthalpies of formation of several other PrO<sub>2-x</sub> compositions were measured by Stubblefield *et al.*<sup>84</sup> and Fitzgibbon *et al.*<sup>85</sup> by solution calorimetry, the recalculated results are given in Table 8. These values show a regular trend between PrO<sub>1.5</sub> and PrO<sub>2</sub>, as shown in Fig. 5.

### 3.6. Pr<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.6.1. Polymorphism and melting point

At room temperature praseodymium sesquioxide has the rare earth hexagonal A-type structure (space group P $\bar{3}$ m1).

Lopato *et al.*<sup>11</sup> reported that A-type Pr<sub>2</sub>O<sub>3</sub> transforms to a hexagonal H-type structure at T = 2318 K, and Shevthenko and Lopato<sup>13</sup> at T = 2303 K. They also found that the H-type phase transforms to a cubic X-type structure at T = 2393 K<sup>11</sup> and T = 2403 K.<sup>13</sup> No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of -1 K needs to be applied for ITS-90. The results by Kravchonskaya *et al.*<sup>87</sup> are in agreement, but these authors only presented them in graphical form (A → H at about 2314 K and H → X at about 2417 K, as extracted from digitized graphs). We select T<sub>trs</sub> = (2310 ± 30) K for the A → H transformation, T<sub>trs</sub> = (2397 ± 30) K for the H → X transformation.

The measurements of the melting temperature of Pr<sub>2</sub>O<sub>3</sub> have been summarized in Table 9, which is based on the IUPAC review by Coutures and Rand,<sup>15</sup> the results being corrected to ITS-90. The selected melting point is (2583 ± 25) K.

#### 3.6.2. Heat capacity and entropy

The low-temperature heat capacity of A-Pr<sub>2</sub>O<sub>3</sub> has been measured by Lyutsareva *et al.*, unpublished results as cited by

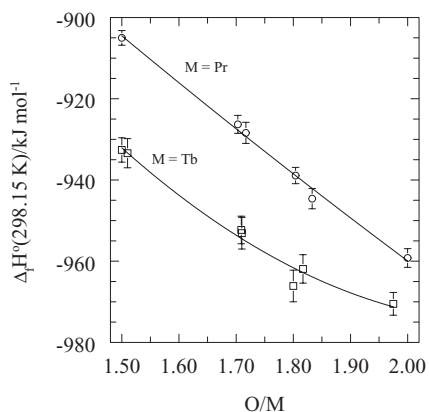


FIG. 5. The enthalpy of formation of compositions in the PrO<sub>2</sub>-PrO<sub>1.5</sub> (□) and TbO<sub>2</sub>-TbO<sub>1.5</sub> (□) systems.

TABLE 9. Temperature of melting of praseodymium sesquioxide

Authors	T <sub>fus</sub> /K	
	Reported	ITS-90
Foex <sup>19</sup>	2668	2582
Mordovin <i>et al.</i> <sup>20</sup>	2390 ± 30	2493 ± 30
Treswjatskii <i>et al.</i> <sup>22</sup>	2533 ± 20	2532 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2585 ± 10	2584 ± 10
Kravchonskaya <i>et al.</i> <sup>87</sup>	2573 <sup>a</sup>	
Mizuno <i>et al.</i> <sup>24</sup>	2549 ± 20	2535 ± 20
Shevthenko and Lopato <sup>13</sup>	2553	2552
Selected value:		2583 ± 25

<sup>a</sup>Derived digitally from the graph for the HfO<sub>2</sub>-Pr<sub>2</sub>O<sub>3</sub> diagram.

Gruber *et al.*<sup>69</sup> Their results are somewhat high near room temperature, and were smoothed together with the high-temperature data by Gruber *et al.*<sup>69</sup> giving no weight to the results above 185 K. The derived standard entropy at T = 298.15 K is selected here

$$S^\circ(298.15 \text{ K}) = (152.7 \pm 0.3) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of A-Pr<sub>2</sub>O<sub>3</sub> has been measured by Pankratz<sup>82</sup> and Kuznetsov and Rezukhina.<sup>88</sup> The former study covers the temperature range from 298.15 to 1800 K. The latter the temperature range from 293 to 1200 K and have only been reported in the form of an equation. Our recommended heat capacity equation is solely based on the results of Pankratz,<sup>82</sup> constrained to C<sub>p</sub><sup>°</sup>(298.15 K) = 118.15 J K<sup>-1</sup> mol<sup>-1</sup> from the low-temperature measurements:

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 121.6594 + 25.5611 \times 10^{-3}(\text{T/K}) - 0.98942 \times 10^6(\text{T/K})^{-2}.$$

This equation is extrapolated to the transition temperature. No heat-capacity or enthalpy measurements have been reported for the H, X, and liquid phases of Pr<sub>2</sub>O<sub>3</sub>. The properties of these modifications have been estimated in a similar way as for La<sub>2</sub>O<sub>3</sub> and Ce<sub>2</sub>O<sub>3</sub>. We thus obtain for the transition enthalpies

$$\Delta_{trs} H^\circ(\text{A} \rightarrow \text{H}) = (28 \pm 8) \text{ kJ mol}^{-1},$$

$$\Delta_{trs} H^\circ(\text{H} \rightarrow \text{X}) = (19 \pm 5) \text{ kJ mol}^{-1},$$

$$\Delta_{fus} H^\circ = (88 \pm 10) \text{ kJ mol}^{-1}.$$

For the heat capacity of the high temperature modifications of Pr<sub>2</sub>O<sub>3</sub> we estimate

$$C_p^\circ(\text{H}, \text{T}) = C_p^\circ(\text{X}, \text{T}) = 145 \text{ J K}^{-1} \text{ mol}^{-1},$$

$$C_p^\circ(\text{liq}, \text{T}) = 157 \text{ J K}^{-1} \text{ mol}^{-1}.$$

#### 3.6.3. Enthalpy of formation

The enthalpy of formation of Pr<sub>2</sub>O<sub>3</sub> has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected value from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Pr}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1809.9 \pm 3.0) \text{ kJ mol}^{-1}.$$



This value is based on the solution calorimetric study by Fitzgibbon *et al.*,<sup>85</sup> who measured the enthalpy of solution of both Pr(cr) and Pr<sub>2</sub>O<sub>3</sub>(cr) in 2.0 mol dm<sup>-3</sup> HCl(aq) (see Table 7). Stubblefield *et al.*<sup>84</sup> determined the enthalpy of reaction of Pr<sub>2</sub>O<sub>3</sub>(cr) with 6.0 mol dm<sup>-3</sup> HNO<sub>3</sub>(aq), but this approach is considered less reliable since some of the hydrogen that is produced during dissolution of the metal, might reduce the nitric acid.

### 3.7. Nd<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.7.1. Polymorphism and melting point

Nd<sub>2</sub>O<sub>3</sub> has a peculiar place in the lanthanide sesquioxide series. It commonly has a A-type hexagonal sesquioxide structure (space group P $\bar{3}$ m1) at room temperature, but also the C-type structure occurs at room temperature. Systematic studies<sup>89</sup> suggest that the C-form is the thermodynamically stable modification from room temperature to about T = 873–923 K. Impurities may affect this phase stability, but it is now generally accepted that the boundary between the A and C rare earth structures occurs near or at Nd<sub>2</sub>O<sub>3</sub> (see Fig. 4). However, as most of the thermodynamic measurements reported in literature have been made for the A-type phase, our recommended values refer to this phase.

Pankratz *et al.*<sup>90</sup> found a small thermal anomaly at ~1345 K by drop calorimetry whose origin is unknown (see below).

At high temperatures Nd<sub>2</sub>O<sub>3</sub> exhibits the common complex polymorphism of the light rare earth oxides. The phase transition to a H-type hexagonal structure (space group P6<sub>3</sub>/mmc) is observed at T = 2375 K (Refs. 10,11 and 91) and T = 2333 K.<sup>13</sup> This phase subsequently transforms to a cubic structure (space group Im $\bar{3}$ m) at T = 2473 K.<sup>10,11,91</sup> These measurements must all be converted to ITS-90. The measurements of Foex and Traverse<sup>10</sup> must be corrected by +14 K, following the procedure outlined by Coutures and Rand.<sup>15</sup> Lopato *et al.*,<sup>11</sup> Wehner *et al.*,<sup>12</sup> and Shevthenko and Lopato<sup>13</sup> reported no (detailed) information on the calibration of their measurements, but assuming the data refer to IPTS-68, a correction of -1 K needs to be applied. We select T<sub>trs</sub> = (2379 ± 30) K for the A → H transformation, T<sub>trs</sub> = (2477 ± 30) K for the H → X transformation.

The measurements of the melting temperature of Nd<sub>2</sub>O<sub>3</sub> have been summarized in Table 10, which is based on the IUPAC review by Coutures and Rand.<sup>15</sup> The selected melting point is (2577 ± 15) K.

#### 3.7.2. Heat capacity and entropy

The low-temperature heat capacity of A-type Nd<sub>2</sub>O<sub>3</sub> has been measured by Goldstein *et al.*<sup>26</sup> from 16 to 300 K, and by Justice and Westrum Jr.<sup>28</sup> from 5 to 350 K. Above 100 K both data sets agree within 0.5%, below 100 K the deviations significantly increase. The recommended standard entropy of Nd<sub>2</sub>O<sub>3</sub> is derived from the low-temperature heat capacity data of Justice and Westrum Jr.<sup>28</sup> who obtained S°(298.15 K) - S°(5 K) = 146.65 J K<sup>-1</sup> mol<sup>-1</sup>. Justice and Westrum estimated S°(5 K) - S°(0 K) = 11.80 J K<sup>-1</sup> mol<sup>-1</sup>, accounting for the 2R

TABLE 10. Temperature of melting of neodymium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	T <sub>fus</sub> /K	
	Reported	ITS-90
Lambertson and Gunzel <sup>17</sup>	2545 ± 20	2551 ± 20
Foex <sup>19</sup>	2583	2598
Mordovin <i>et al.</i> <sup>20</sup>	2486 ± 30	2489 ± 30
Noguchi and Mizuno <sup>21</sup>	2506 ± 20	2510 ± 20
Gibby <i>et al.</i> <sup>92</sup>	2573 ± 25	2572 ± 25
Treswjatskii <i>et al.</i> <sup>22</sup>	2593 ± 20	2592 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2598 ± 10	2597 ± 10
Bober <i>et al.</i> <sup>93</sup>	2540 ± 40	2539 ± 40
Mizuno <i>et al.</i> <sup>24</sup>	2563 ± 20	2550 ± 20
Shevthenko and Lopato <sup>13</sup>	2573	2572
Coutures <sup>94</sup>	2613 ± 10	2612 ± 10
Salikhov and Kan <sup>95</sup>		2564 ± 10
Selected value:		2577 ± 15

In (2) contribution of the ground state doublet of the <sup>4</sup>I<sub>9/2</sub> multiplet, resulting in

$$S^\circ(298.15 \text{ K}) = (158.7 \pm 1.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment has been measured by Blomeke and Ziegler<sup>29</sup> from 384 to 1172 K and Pankratz *et al.*<sup>90</sup> from 400 to 1795 K. The data of the two studies are in excellent agreement and smoothly join the low-temperature data (Fig. 6). Pankratz *et al.*<sup>90</sup> observed a minor thermal anomaly near 1395 K. We think this is an experimental artefact, as high temperature X-ray diffraction studies<sup>10,96</sup> do not reveal any evidence for such a transformation. Our recommended heat capacity equation is based on a polynomial fit of the results of both studies. The equation is constrained to C<sub>p</sub><sup>o</sup> = 111.34 J K<sup>-1</sup> mol<sup>-1</sup> from the low-temperature measurements.<sup>28</sup> We thus obtain

$$C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = 117.1079 + 28.13655 \times 10^{-3} (T/\text{K}) - 1.25845 \times 10^6 (T/\text{K})^{-2}.$$

This equation is extrapolated to the transition temperature. No heat-capacity or enthalpy measurements have been reported for the H, X, and liquid phases of Nd<sub>2</sub>O<sub>3</sub>. The properties of these modifications have been estimated in a similar way as for La<sub>2</sub>O<sub>3</sub> and La<sub>2</sub>O<sub>3</sub>. We thus obtain for the transition enthalpies

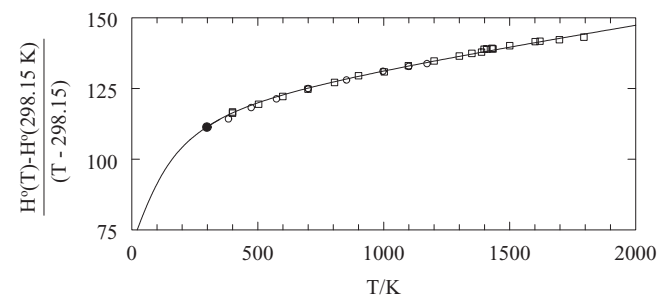


FIG. 6. The reduced enthalpy increment (in J K<sup>-1</sup> mol<sup>-1</sup>) of Nd<sub>2</sub>O<sub>3</sub>; ○, Blomeke and Ziegler<sup>29</sup>; □, King *et al.*<sup>27</sup>; ●, value derived from the low-temperature measurements by Justice and Westrum, Jr.,<sup>28</sup> the curve shows the recommended equation.

$$\begin{aligned}\Delta_{trs}H^\circ(A \rightarrow H) &= (29 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(H \rightarrow X) &= (20 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (88 \pm 10) \text{ kJ mol}^{-1}.\end{aligned}$$

For the heat capacity of the high temperature modifications of  $\text{Nd}_2\text{O}_3$  we estimate

$$\begin{aligned}C_p^\circ(\text{H}, T) &= C_p^\circ(\text{X}, T) = 145 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{liq}, T) &= 160 \text{ J K}^{-1} \text{ mol}^{-1}.\end{aligned}$$

### 3.7.3. Enthalpy of formation

The enthalpy of formation of hexagonal  $\text{Nd}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected value from that work, because no new information has been published since

$$\Delta_f H^\circ(\text{Nd}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1806.9 \pm 3.0) \text{ kJ mol}^{-1}.$$

This value has been derived from the studies listed in Table 11 (corrected for some errors), that have been employing solution as well as combustion calorimetry. The three combustion calorimetric measurements<sup>40,98,99</sup> are in reasonable agreement, but the value by Huber Jr. and Holley Jr.,<sup>98</sup>  $\Delta_f H^\circ(298.15 \text{ K}) = -(1808.1 \pm 1.0) \text{ kJ mol}^{-1}$ , was considered to be far more accurate since the starting materials were of rather high purity and the combustion was complete. The solution calorimetric studies heavily rely on the value for the enthalpy of solution of Nd metal used in the Hess cycle to derive the enthalpy of formation of  $\text{Nd}_2\text{O}_3$ . Cordfunke and Konings<sup>34</sup> therefore used the enthalpy of solution of Nd(cr) in  $\text{HCl}(\text{aq})$  by Merli *et al.*,<sup>39</sup> Tiflova,<sup>106</sup> and Stuve,<sup>105</sup> which are consistent with each other and measured on high-purity Nd(cr) samples, to recalculate the reported measurements in a systematic manner. The selected enthalpy of formation is the mean of the results of Huber Jr. and Holley Jr.<sup>98</sup> and the values derived from the results of Fitzgibbon *et al.*<sup>100</sup> and Popova and Monaenkova.<sup>103</sup>

TABLE 11. The enthalpy of formation of  $\text{Nd}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of Nd(cr) and  $\text{Nd}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Muthmann and Weis <sup>40</sup>	C			-1820
Matignon <sup>97</sup>	S (0.5)		-441.4	
Huber, Jr. and Holley, Jr. <sup>98</sup>	C			-1808.1 ± 1.0
Spedding <i>et al.</i> <sup>99</sup>	C			-1798.2
				-1789.2
Fitzgibbon <i>et al.</i> <sup>100</sup>	S (2.0)	$[-691.7 \pm 1.5]^b$	$-434.0 \pm 0.6$	$-1807.1 \pm 3.1$
	S (4.0)	$[-693.6 \pm 1.5]^c$	$-438.3 \pm 1.3$	$-1807.3 \pm 2.2$
Yashvili and Gvelesiani <sup>101</sup>	S (1.0)	$[-689.6 \pm 2.0]^d$	$-434.7 \pm 2.1$	$-1799.6 \pm 4.5$
Morss <i>et al.</i> <sup>102</sup>	S (6.0)	$[-695.7 \pm 1.8]^c$	$-419.6 \pm 6.0$	$-1831.7 \pm 7.0$
Popova and Monaenkova <sup>103</sup>	S (2.19)	$-686.8 \pm 1.0$	$-434.2 \pm 0.7$	$-1797.1 \pm 2.1$
		$[-690.8 \pm 1.6]^e$		$-1805.1 \pm 3.3$
		$[-691.9 \pm 2.0]^d$		$-1807.3 \pm 4.1$
Hennig and Oppermann <sup>104</sup>	S (4.0)	$[-691.7 \pm 1.5]^b$	$-419.5 \pm 0.4$	$-1816.1 \pm 1.9$
Selected value:				$-1806.9 \pm 3.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ ;

<sup>b</sup>Estimated from the data of Merli *et al.*<sup>39</sup>

<sup>c</sup>Stuve<sup>105</sup>

<sup>d</sup>Merli *et al.*<sup>39</sup>

<sup>e</sup>Tiflova.<sup>106</sup>

## 3.8. $\text{Pm}_2\text{O}_3(\text{cr,l})$

### 3.8.1. Polymorphism and melting point

The monoclinic B-type (space group  $C2/m$ ) and the cubic C-type (space group  $Fm\bar{3}m$ ) rare earth sesquioxides structure of  $\text{Pm}_2\text{O}_3$  have been found at room temperature, like is the case for its neighboring sesquioxides. According to the generally accepted stability diagram of the lanthanide sesquioxides, the C-type structure is probably the thermodynamic stable modification at room temperature. Chikalla *et al.*<sup>107</sup> found that the B-type form is stable above about  $T = 1073 \text{ K}$ , which is in reasonable agreement with the trend in the lanthanide sesquioxides as shown in Fig. 4, but claimed that this transition is irreversible. They determined the transition temperatures and the melting point, which we select here after a correction of  $-1 \text{ K}$ , assuming the data refer to IPTS-68:  $T = (2013 \pm 20) \text{ K}$  for the  $B \rightarrow A$  transformation,  $T = (2407 \pm 20) \text{ K}$  for the  $A \rightarrow H$  transformation, and  $T = (2497 \pm 20) \text{ K}$  for the  $H \rightarrow X$  transformation.

The melting point of  $\text{Pm}_2\text{O}_3$  was measured by Gibby *et al.*<sup>108</sup> as  $T = (2408 \pm 30) \text{ K}$ , and by Chikalla *et al.*<sup>107</sup> as  $T = (2593 \pm 30) \text{ K}$ . The former value is significantly lower than those of the neighboring  $\text{Nd}_2\text{O}_3$  and  $\text{Sm}_2\text{O}_3$ , and must be in error. We therefore select the value of Chikalla *et al.*<sup>107</sup> which becomes  $T = (2592 \pm 30) \text{ K}$  on ITS-90.

### 3.8.2. Heat capacity and entropy

Konings<sup>109</sup> estimated the standard entropy of  $\text{Pm}_2\text{O}_3$  from a systematic analysis of the trend in the lanthanide sesquioxides, describing the entropy as the sum of the lattice and excess components. The lattice component was obtained by interpolation of the values for  $\text{La}_2\text{O}_3$ ,  $\text{Gd}_2\text{O}_3$ , and  $\text{Lu}_2\text{O}_3$ , the excess component from the ground state degeneracy of the  ${}^2F_{7/2}$  multiplet. This value is selected here

$$S^\circ(298.15 \text{ K}) = (158.0 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high temperature heat capacity equations for C-Pm<sub>2</sub>O<sub>3</sub> (298–1073 K) and B-Pm<sub>2</sub>O<sub>3</sub> (1073–2013 K) have been estimated from the values for the other lanthanide sesquioxides as

$$\begin{aligned} C_p^\circ(\text{C}, T)/(\text{JK}^{-1} \text{mol}^{-1}) &= 122.9493 + 30.0141 \times 10^{-3}(T/\text{K}) \\ &\quad - 1.85217 \times 10^6(T/\text{K})^{-2} \\ C_p^\circ(\text{B}, T)/(\text{JK}^{-1} \text{mol}^{-1}) &= 129.454 + 19.960 \times 10^{-3}(T/\text{K}). \end{aligned}$$

For the heat capacity high temperature modifications we have assumed the same values as estimated for Sm<sub>2</sub>O<sub>3</sub> (see below)

$$\begin{aligned} C_p^\circ(\text{H}, T) = C_p^\circ(\text{X}, T) &= 165 \text{ JK}^{-1} \text{mol}^{-1}, \\ C_p^\circ(\text{liq}, T) &= 179 \text{ J} \cdot \text{K}^{-1} \text{mol}^{-1}. \end{aligned}$$

The transition enthalpies have been estimated as outlined for Ce<sub>2</sub>O<sub>3</sub>:

$$\begin{aligned} \Delta_{trs}H^\circ(\text{C} \rightarrow \text{B}) &= (7 \pm 3) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{B} \rightarrow \text{A}) &= (6 \pm 3) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (31 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (20 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (88 \pm 10) \text{ mol}^{-1}. \end{aligned}$$

### 3.8.3. Enthalpy of formation

Experimental values for the standard molar enthalpy of formation of promethium sesquioxide are not available in the literature. The selected  $\Delta_f H^\circ(298.15 \text{ K})$  value has been estimated by Cordfunke and Konings<sup>34</sup> using the dependence of  $\Delta_f H^\circ(\text{Ln}_2\text{O}_3, \text{cr}) - 2\Delta_f H^\circ(\text{Ln}^{3+}, \text{aq})$  on the atomic radii of the trivalent lanthanides and the enthalpy of formation of Pm<sup>3+</sup>(aq). Pm<sub>2</sub>O<sub>3</sub>(cr) is a boundary compound of the domains of existence of the hexagonal and monoclinic rare earth sesquioxides, both crystal structures are acceptable for promethium sesquioxide with equal degree of probability.<sup>110</sup> Assuming the hexagonal crystal structure of Pm<sub>2</sub>O<sub>3</sub>(cr) to be the stable form, the enthalpy of formation was calculated as

$$\Delta_f H^\circ(\text{Pm}_2\text{O}_3, \text{cr}) = -(1811 \pm 21) \text{ kJ mol}^{-1}.$$

## 3.9. Sm<sub>2</sub>O<sub>3</sub>(cr,l)

### 3.9.1. Polymorphism and melting point

The monoclinic B-type (space group C2/m) as well as the cubic C-type (space group Fm $\bar{3}$ m) structures of samarium sesquioxide have been reported to be stable at room temperature. According to the generally accepted stability diagram of the lanthanide sesquioxides (see Fig. 4), the C-type structure is probably the thermodynamic stable modification from room temperature to about 900 K. However, the energetics of the two modifications are probably very close and the transformation kinetics very sluggish. Since most of the thermodynamic measurements have been made for the B modification, our recommended values refer to this phase only.

TABLE 12. Temperature of melting of samarium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	T <sub>fus</sub> /K	
	Reported	ITS-90
Wisnyi and Pijanowski <sup>111</sup>	2573 ± 50	2590 ± 50
Curtis and Johnson <sup>112</sup>	2623 ± 50	2626 ± 50
Foex <sup>19</sup>	2593	2607
Mordovin <i>et al.</i> <sup>20</sup>	2536 ± 30	2539 ± 30
Noguchi and Mizuno <sup>21</sup>	2506 ± 20	2509 ± 20
Gibby <i>et al.</i> <sup>92</sup>	2623 ± 25	2622 ± 25
Tresvjatskii <i>et al.</i> <sup>22</sup>	2613 ± 20	2612 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2598 ± 10	2598 ± 10
Mizuno <i>et al.</i> <sup>24</sup>	2583	2570
Shevthenko and Lopato <sup>13</sup>	2583	2582
Selected value:		2613 ± 15

Foex and Traverse<sup>10</sup> reported that B-type Sm<sub>2</sub>O<sub>3</sub> transforms to the A-type Ln<sub>2</sub>O<sub>3</sub> structure at ~2173 K. They also found that the A → H and H → X transformation occurs at ~2373 and 2473–2523 K, respectively. These values must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +14 K. Lopato *et al.*<sup>11</sup> reported the B → A transformation at T = 2193 K and the A → H and H → X transformations at T = 2403 K and T = 2553 K, respectively. Shevthenko and Lopato<sup>13</sup> reported the B → A transformation at T = 2143 K, the A → H at T = 2343 K, and the H → X transformations at T = 2498 K. No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of –1 K needs to be applied to convert to ITS-90. We select T<sub>trs</sub> = (2190 ± 20) K for the B → A transformation, T<sub>trs</sub> = (2395 ± 20) K for the A → H transformation and T<sub>trs</sub> = (2533 ± 30) K for the H → X transformation.

The measurements of the melting temperature of Sm<sub>2</sub>O<sub>3</sub> have been summarized in Table 12, which is based on the IUPAC review by Coutures and Rand.<sup>15</sup> The selected melting point is (2613 ± 15) K.

### 3.9.2. Heat capacity and entropy

The low-temperature heat capacity of B-type Sm<sub>2</sub>O<sub>3</sub> has been measured by Justice and Westrum Jr.<sup>113</sup> from 10 to 350 K. They obtained S°(298.15 K) – S°(10 K) = 138.99 J K<sup>–1</sup> mol<sup>–1</sup>. Extrapolation of these results to T = 0 K, and accounting for the 2Rln(2) contribution of the ground state doublet of the split <sup>6</sup>H<sub>7/2</sub> multiplet, gives for the entropy at T = 298.15 K:

$$S^\circ(298.15 \text{ K}) = (150.6 \pm 0.3) \text{ J K}^{-1} \text{mol}^{-1}.$$

The high-temperature enthalpy increment of Sm<sub>2</sub>O<sub>3</sub> has been measured by Curtis and Johnson,<sup>112</sup> Pankratz *et al.*,<sup>114</sup> and Gvelesiani *et al.*<sup>115</sup>. The latter two groups measured the B-type hexagonal and the C-type cubic modifications, the results being in good agreement (Fig. 7). Curtis and Johnson<sup>112</sup> reported data on a sample consisting of a mixture of B and C, their results being in poor agreement with the other results. Pankratz *et al.*<sup>114</sup> reported a transition (at about 1195 K) with a small enthalpy effect (1.045 kJ mol<sup>–1</sup>) for B-type Sm<sub>2</sub>O<sub>3</sub> that was not observed by Gvelesiani *et al.*<sup>115</sup>. Also DTA analysis by Curtis and Johnson<sup>112</sup> did not reveal phase transformations up

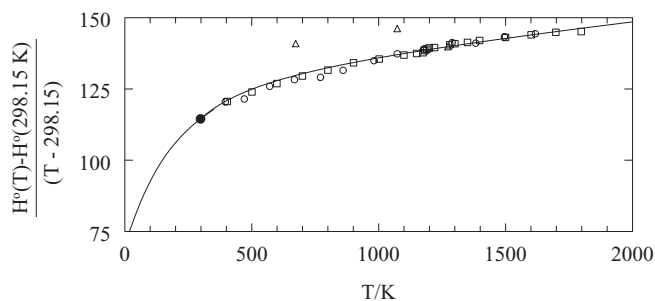


Fig. 7. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{mol}^{-1}$ ) of B- $\text{Sm}_2\text{O}_3$ ;  $\circ$ , Gvelesiani *et al.*<sup>115</sup>;  $\square$ , Pankratz *et al.*<sup>114</sup>;  $\triangle$ , Curtis and Johnson<sup>112</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice and Westrum Jr.<sup>113</sup>; the curve shows the recommended equation.

to 1403 K. Since there is no additional confirmation of this effect, we have neglected it in our analysis. Our recommended heat capacity equation for B- $\text{Sm}_2\text{O}_3$  is based on the combined results of Pankratz *et al.*<sup>114</sup> and Gvelesiani *et al.*<sup>115</sup>:

$$C_p^\circ / (\text{J K}^{-1} \text{mol}^{-1}) = 129.7953 + 19.03114 \times 10^{-3} (T/\text{K}) - 1.86227 \times 10^6 (T/\text{K})^{-2}.$$

This equation is constrained to  $C_p^\circ(298.15 \text{ K}) = 114.52 \text{ J K}^{-1} \text{mol}^{-1}$ , as derived from the low-temperature measurements Justice and Westrum Jr.<sup>113</sup> For C- $\text{Sm}_2\text{O}_3$  we obtain (without constraint)

$$C_p^\circ / (\text{J K}^{-1} \text{mol}^{-1}) = 132.4358 + 18.7799 \times 10^{-3} (T/\text{K}) - 2.40860 \times 10^6 (T/\text{K})^{-2}.$$

The entropy change of the C  $\rightarrow$  B transformation in the lanthanide sesquioxides was estimated from high pressure studies by Hoekstra<sup>116</sup> as  $6.3 \text{ J K}^{-1} \text{mol}^{-1}$  using the Clausius-Clapeyron equation, which is selected here. This quantity can also be estimated from the differences of the enthalpy equations at the transition temperature ( $-0.3 \text{ kJ mol}^{-1}$  at 900 K), plus the difference in the enthalpies of formation at 298.15 K ( $3.8 \pm 6.2 \text{ kJ mol}^{-1}$ ), giving  $\Delta_{trs}H^\circ = (3.5 \pm 6.2) \text{ kJ mol}^{-1}$  corresponding to  $\Delta_{trs}S^\circ = 3.9 \text{ J K}^{-1} \text{mol}^{-1}$ , in fair agreement with Hoekstra's estimate.<sup>116</sup>

No heat-capacity or enthalpy measurements have been reported for the H, X, and liquid phases of  $\text{Sm}_2\text{O}_3$ . The properties of these modifications have been estimated in a similar way as for  $\text{La}_2\text{O}_3$  and  $\text{Ce}_2\text{O}_3$ . We thus obtain for the transition enthalpies

$$\begin{aligned} \Delta_{trs}H^\circ(\text{C} \rightarrow \text{B}) &= (6 \pm 3) \text{ kJ mol}^{-1} \\ \Delta_{trs}H^\circ(\text{B} \rightarrow \text{A}) &= (7 \pm 3) \text{ kJ mol}^{-1} \\ \Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (32 \pm 8) \text{ kJ mol}^{-1} \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (20 \pm 5) \text{ kJ mol}^{-1} \\ \Delta_{fus}H^\circ &= (89 \pm 10) \text{ kJ mol}^{-1} \end{aligned}$$

For the heat capacity of the high temperature modifications of  $\text{Sm}_2\text{O}_3$  we estimate

$$\begin{aligned} C_p^\circ(\text{A}, T) &= 140 \text{ J K}^{-1} \text{mol}^{-1} \\ C_p^\circ(\text{H}, T) &= C_p^\circ(\text{X}, T) = 165 \text{ J K}^{-1} \text{mol}^{-1} \\ C_p^\circ(\text{liq}, T) &= 179 \text{ J K}^{-1} \text{mol}^{-1}. \end{aligned}$$

### 3.9.3. Enthalpy of formation

The enthalpy of formation of monoclinic and cubic  $\text{Sm}_2\text{O}_3$  have been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected value from that work, since no new information has been published since

$$\begin{aligned} \Delta_f H^\circ(\text{Sm}_2\text{O}_3, \text{monoclinic}, 298.15 \text{ K}) &= -(1823.0 \pm 4.0) \text{ kJ mol}^{-1} \\ \Delta_f H^\circ(\text{Sm}_2\text{O}_3, \text{cubic}, 298.15 \text{ K}) &= -(1826.8 \pm 4.8) \text{ kJ mol}^{-1}. \end{aligned}$$

The values for the enthalpy of formation of monoclinic  $\text{Sm}_2\text{O}_3$  obtained by Huber Jr. *et al.*<sup>117</sup> and Baker *et al.*<sup>118</sup> by oxygen-bomb combustion calorimetry and by Baker *et al.*<sup>118</sup> using solution calorimetry are in reasonable agreement (see Table 13), and are the basis for the selected value. Later results by Gvelesiani and Yashvili<sup>38</sup> significantly deviate from this value, but the nonmetallic impurities in their samples are not reported, which could be an explanation for the difference. The measurements by Hennig and Oppermann<sup>119</sup> of the enthalpy of solution of  $\text{Sm}_2\text{O}_3$  in  $\text{HCl}(\text{aq})$  is in good agreement with the

TABLE 13. The enthalpy of formation of  $\text{Sm}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Sm}(\text{cr})$  and  $\text{Sm}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ / \text{kJ mol}^{-1}$	$\Delta H_2^\circ / \text{kJ mol}^{-1}$	$\Delta_f H^\circ / \text{kJ mol}^{-1}$
Huber Jr. <i>et al.</i> <sup>117</sup>	C			$-1815.4 \pm 2.0$
Montgomery and Hubert <sup>36</sup>	S (0.48)		$-408.8 \pm 1.4$	
Spedding <i>et al.</i> <sup>99</sup>	C			$-1777.3$
Gvelesiani and Yashvili <sup>38</sup>	S (0.7)	$-683.7 \pm 5.4$	$-389.5 \pm 0.4$	$-1835.4 \pm 10.8$
Baker <i>et al.</i> <sup>118</sup>	S (1.0)	$-682.6 \pm 2.2$	$-391.2 \pm 3.6$	$-1831.5 \pm 5.7$
	C			$-1824.2 \pm 2.6$
	S (2.0)	$-690.1 \pm 1.3$	$-417.1 \pm 1.2$	$-1820.8 \pm 2.9$
	S (3.99)	$-689.5 \pm 3.8$	$-406.7 \pm 4.6$	$-1830.7 \pm 8.9$
Hennig and Oppermann <sup>119</sup>	S (4.0)		$-412.8 \pm 0.5$	$-1824.6 \pm 7.6^b$
Selected value:				$-1823.0 \pm 4.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$

<sup>b</sup>Using  $\Delta H_1^\circ$  from Baker *et al.*<sup>118</sup>



results by Baker *et al.*<sup>118</sup> However, because of the poor characterisation of the Sm<sub>2</sub>O<sub>3</sub> sample and the fact that the measurements by Hennig and Opperman deviate significantly for most of the lanthanide sesquioxides (see La<sub>2</sub>O<sub>3</sub>, Nd<sub>2</sub>O<sub>3</sub>, and Eu<sub>2</sub>O<sub>3</sub>) this value was not taken into account by Cordfunke and Konings.<sup>34</sup>

The enthalpy of formation of cubic Sm<sub>2</sub>O<sub>3</sub> was also derived from the work of Baker *et al.*<sup>118</sup> and Gvelesiani and Yashvili,<sup>38</sup> who both determined the value for  $\Delta_{trs}H^\circ$  (monoclinic/cubic) by solution calorimetric measurements of the two crystallographic modifications. As discussed by Cordfunke and Konings<sup>34</sup> the results are in reasonable agreement,  $-(3.7 \pm 2.6)$  kJ mol<sup>-1</sup> and  $-(5.5 \pm 4.0)$  kJ mol<sup>-1</sup>, respectively, but the value of Baker *et al.*<sup>118</sup> is preferred for reasons given in the preceding paragraph.

### 3.10. Eu<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.10.1. Polymorphism and melting point

Europium sesquioxide has a complex polymorphism: both the monoclinic (B-structure) and the cubic (C-structure) are found to coexist at room temperature. At standard pressure conditions the C form is the most stable form, the B form thus being metastable. The C-Eu<sub>2</sub>O<sub>3</sub> phase has a fluorite-type cubic structure (space group Fm $\bar{3}$ m). The B-Eu<sub>2</sub>O<sub>3</sub> form is the monoclinic modification of europium sesquioxide (space group C2/m).

The C  $\rightarrow$  B transformation has been studied extensively. Stecura<sup>120</sup> reported that this transition is irreversible, but most other studies have found that the transformation kinetics are sluggish but reversible. The temperature was found at 1348 K (Ref. 121) and 1373 K.<sup>89</sup> Ainscough *et al.*<sup>122</sup> observed significant differences in the transformation temperature and rate for air and hydrogen atmospheres, the transformation taking place faster and at lower temperature (75 K) in air. This was confirmed by Suzuki *et al.*<sup>123</sup> for air and vacuum. More recently Sukhushina *et al.*<sup>124</sup> measured the oxygen potential of stoichiometric B-Eu<sub>2</sub>O<sub>3</sub> and C-Eu<sub>2</sub>O<sub>3</sub> between 1150 and 1450 K. From the results they derive  $T_{trs} = 1350.6$  K from the intersection of the curves. The phase transformations at high temperatures have been studied by Foex and Traverse.<sup>10</sup> They reported the B  $\rightarrow$  A transformation at  $T = 2313$  K, the A  $\rightarrow$  H transformation at  $T = 2413$  K, and the H  $\rightarrow$  X transformation at  $T = 2543$  K. These results must be converted to ITS-90 by +14 K, following the procedure outlined by Coutures and Rand.<sup>15</sup>

We select  $T_{trs} = (1350 \pm 15)$  K for the C  $\rightarrow$  B transformation,  $T_{trs} = (2327 \pm 30)$  K for the B  $\rightarrow$  A transformation,  $T_{trs} = (2427 \pm 30)$  K for the A  $\rightarrow$  H transformation, and  $T_{trs} = (2557 \pm 30)$  K for the H  $\rightarrow$  X transformation.

The measurements of the melting temperature of Eu<sub>2</sub>O<sub>3</sub> have been summarized in Table 14, which is based on the IUPAC review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2622 \pm 20)$  K.

#### 3.10.2. Heat capacity and entropy

The low-temperature heat capacity of C-Eu<sub>2</sub>O<sub>3</sub> has been measured by Lyutsareva *et al.*<sup>126</sup> from 7 to 319 K and these

TABLE 14. Temperature of melting of europium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Wisnyi and Pijanowski <sup>111</sup>	2323 $\pm$ 30	2339 $\pm$ 30
Schneider <sup>125</sup>	2513 $\pm$ 10	2519 $\pm$ 10
Foex <sup>19</sup>	2603	2617
Mordovin <i>et al.</i> <sup>20</sup>	2276 $\pm$ 30	2278 $\pm$ 30
Noguchi and Mizuno <sup>21</sup>	2564 $\pm$ 20	2567 $\pm$ 20
Coutures <i>et al.</i> <sup>23</sup>	2633 $\pm$ 10	2632 $\pm$ 10
Mizuno <i>et al.</i> <sup>24</sup>	2618 $\pm$ 20	2605 $\pm$ 20
Selected value:		2622 $\pm$ 20

authors also measured the low-temperature heat capacity of B-Eu<sub>2</sub>O<sub>3</sub> from 8 to 311 K. The heat capacity and standard entropy values for B-Eu<sub>2</sub>O<sub>3</sub> are consistent with the results for the other A- and B-type Ln<sub>2</sub>O<sub>3</sub> compounds considering the lattice and excess electronic components.<sup>127</sup> For that reason we recommend the entropy value for B-Eu<sub>2</sub>O<sub>3</sub> from this work, with an increased uncertainty:

$$S^\circ(298.15 \text{ K}) = (143.5 \pm 0.5) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The heat capacity and entropy values for C-Eu<sub>2</sub>O<sub>3</sub>, on the other hand, only poorly agree with the results for the C-type Ln<sub>2</sub>O<sub>3</sub> compounds, giving a too high lattice component above 200 K after subtraction of the excess heat capacity using crystal field data for Eu<sup>3+</sup> in a cubic (C) environment. A similar observation was made for Pr<sub>2</sub>O<sub>3</sub> measured by the same authors (see above). We therefore reject the standard entropy  $S^\circ(298.15 \text{ K}) = (142.24 \pm 0.14) \text{ J K}^{-1} \text{ mol}^{-1}$  derived from that work, and estimate for C-Eu<sub>2</sub>O<sub>3</sub>:

$$S^\circ(298.15 \text{ K}) = (136.4 \pm 2.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of C-Eu<sub>2</sub>O<sub>3</sub> has been reported by Pankratz and King,<sup>128</sup> and Tsagareishvili and Gvelesiani.<sup>129</sup> The two data sets are in reasonable agreement. These results also show a slight misfit with the low-temperature data (see Fig. 8), confirming these are probably too high. The combined results have been fitted to the polynomial equation, to yield for the heat capacity

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 136.2978 + 14.9877 \times 10^{-3}(T/K) - 1.4993 \times 10^6(T/K)^{-2}.$$

This equation is constrained to  $C_p^\circ(298.15 \text{ K}) = 123.9 \text{ J K}^{-1} \text{ mol}^{-1}$  as estimated by us, and not to  $C_p^\circ(298.15 \text{ K}) = 127.09 \text{ J K}^{-1} \text{ mol}^{-1}$  as derived from the low-temperature heat capacity measurements.

The high-temperature enthalpy increment of B-Eu<sub>2</sub>O<sub>3</sub> has also been determined by Curtis and Tharp,<sup>130</sup> Pankratz and King,<sup>128</sup> and Gvelesiani *et al.*<sup>115</sup>; the latter article also includes some numerical results from previously reported measurements by Tsagareishvili and Gvelesiani.<sup>129</sup> The results of Pankratz and King<sup>128</sup> and Gvelesiani *et al.*<sup>115</sup> are in poorer agreement, the former being up to 2.5% lower, in contrast to the results for C-Eu<sub>2</sub>O<sub>3</sub>. An unexplained transition at  $\sim 900$  K

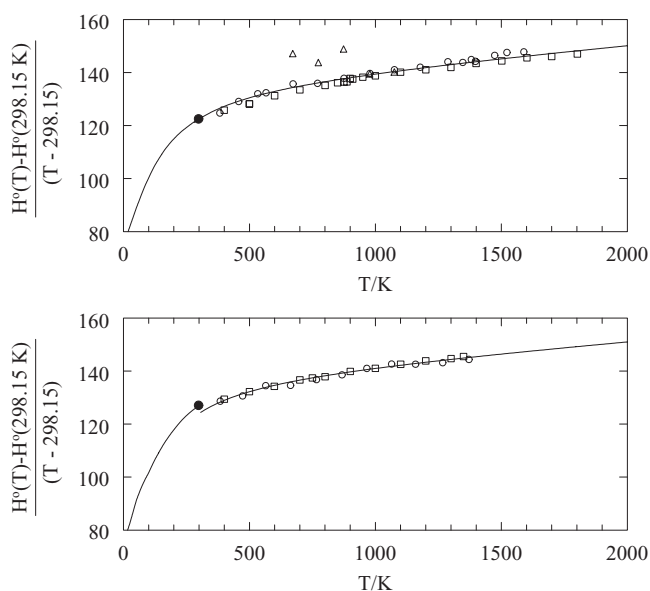


Fig. 8. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{mol}^{-1}$ ) of B- $\text{Eu}_2\text{O}_3$  (top) and C- $\text{Eu}_2\text{O}_3$  (bottom);  $\circ$ , Gvelesiani *et al.*<sup>115</sup>;  $\square$ , Pankratz and King<sup>128</sup>;  $\triangle$ ,<sup>130</sup>  $\bullet$ , value derived from the low-temperature measurements by Lyutsareva *et al.*<sup>126</sup>; the curves show the recommended equations.

was observed by Pankratz and King<sup>128</sup> which was not present in the results of Gvelesiani *et al.*,<sup>115</sup> nor found in high temperature dilatometry studies.<sup>130,131</sup> The recommended value is derived from the polynomial fit of the data by Pankratz and King<sup>128</sup> and Gvelesiani *et al.*,<sup>115</sup> constrained to  $C_p^\circ(298.15 \text{ K}) = 122.33 \text{ J K}^{-1} \text{mol}^{-1}$ ,<sup>126</sup> yielding for the heat capacity:

$$C_p^\circ/(\text{J K}^{-1} \text{mol}^{-1}) = 133.3906 + 16.6443 \times 10^{-3}(T/\text{K}) - 1.42435 \times 10^6(T/\text{K})^{-2}.$$

The enthalpy of the C  $\rightarrow$  B transition can be approximated from the differences of the enthalpy equations at the transition temperature ( $-1.2 \text{ kJ mol}^{-1}$  at 1300 K), plus the difference in the enthalpies of formation at 298.15 K ( $+12.1 \text{ kJ mol}^{-1}$ ), yielding  $\Delta_{\text{trs}}S^\circ = 8.1 \text{ J K}^{-1} \text{mol}^{-1}$ , in

fair agreement with Hoekstra's estimate,  $\Delta_{\text{trs}}S^\circ = 6.3 \text{ J K}^{-1} \text{mol}^{-1}$  from high pressure studies.<sup>116</sup> We select the latter value, giving

$$\Delta_{\text{trs}}H^\circ(\text{C} \rightarrow \text{B}) = (9 \pm 3) \text{ kJ mol}^{-1}$$

Heat-capacity or enthalpy measurements have not been reported for the H, X, and liquid phases of  $\text{Eu}_2\text{O}_3$ . Foex and Traverse<sup>10</sup> found that the thermal effects of the B  $\rightarrow$  A transformation are average compared to the other transformations. The properties of these modifications have been estimated in a similar way as for  $\text{Ce}_2\text{O}_3$ . We thus obtain for the transition enthalpies

$$\begin{aligned} \Delta_{\text{trs}}H^\circ(\text{B} \rightarrow \text{A}) &= (7 \pm 2) \text{ kJ mol}^{-1}, \\ \Delta_{\text{trs}}H^\circ(\text{A} \rightarrow \text{H}) &= (33 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{\text{trs}}H^\circ(\text{H} \rightarrow \text{X}) &= (21 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{\text{fus}}H^\circ &= (89 \pm 10) \text{ kJ mol}^{-1}. \end{aligned}$$

For the heat capacity of the high temperature modifications of  $\text{Eu}_2\text{O}_3$  we estimate

$$\begin{aligned} C_p^\circ(\text{A}, T) &= 141 \text{ J K}^{-1} \text{mol}^{-1} \\ C_p^\circ(\text{H}, T) &= C_p^\circ(\text{X}, T) = 144 \text{ J K}^{-1} \text{mol}^{-1} \\ C_p^\circ(\text{liq}, T) &= 156 \text{ J K}^{-1} \text{mol}^{-1} \end{aligned}$$

### 3.10.3. Enthalpy of formation

The enthalpy of formation of monoclinic as well as cubic  $\text{Eu}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\begin{aligned} \Delta_f H^\circ(\text{Eu}_2\text{O}_3, \text{monoclinic}, 298.15 \text{ K}) &= -(1650.4 \pm 4.0) \text{ kJ mol}^{-1}, \\ \Delta_f H^\circ(\text{Eu}_2\text{O}_3, \text{cubic}, 298.15 \text{ K}) &= -(1662.5 \pm 6.0) \text{ kJ mol}^{-1}. \end{aligned}$$

The value for monoclinic europium sesquioxide is based on an analysis of combustion as well as solution calorimetry studies, as shown in Table 15. The selected value is the mean of the results by Holley and coworkers.<sup>132,133</sup> Their results

TABLE 15. The enthalpy of formation of monoclinic  $\text{Eu}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Eu}(\text{cr})$  and  $\text{Eu}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Huber Jr. <i>et al.</i> <sup>132</sup>	C			$-1648.1 \pm 3.8$
Yashvili and Gvelesiani <sup>101</sup>	S (1.0)	$-632.6 \pm 3.8$	$-397.5 \pm 3.8$	$-1725.5 \pm 8.5$
		$[-607 \pm 4]^b$		$-1674.0 \pm 8.9$
Fitzgibbon <i>et al.</i> <sup>133</sup>	C			$-1651.0 \pm 3.8$
	S (4.0)	$-605.2 \pm 2.9$	$-416.8 \pm 1.5$	$-1652.0 \pm 6.0$
	S (6.0)	$[-589.9 \pm 2.9]^c$	$-415.2 \pm 2.7$	$-1624.5 \pm 6.4$
		$[-603 \pm 4]^b$		$-1650.7 \pm 8.0$
Hennig <i>et al.</i> <sup>134</sup>	S (4.0)	$[-583.0 \pm 2.5]^c$	$-338.3 \pm 0.3$	$-1686.2 \pm 5.0$
				$-1730.6 \pm 5.8^d$
	selected value:			$-1650.4 \pm 4.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ ;

<sup>b</sup>Estimated by Cordfunke and Konings<sup>34</sup> from studies for various lanthanide metals at 1.0, 3.0 and 6.0  $\text{mol dm}^{-3}$   $\text{HCl}$  (aq) by Merli *et al.*<sup>39</sup>;

<sup>c</sup>Stuve<sup>135</sup>

<sup>d</sup>Using  $\Delta H_1^\circ$  from Fitzgibbon *et al.*<sup>133</sup>

obtained by combustion calorimetry are in excellent agreement. The value derived from solution calorimetric measurements of Eu(cr) and Eu<sub>2</sub>O<sub>3</sub>(cr) in 4.0 mol dm<sup>-3</sup> HCl(aq) excellently agrees with these. Further support is obtained from the measurements of the enthalpy of solution of Eu<sub>2</sub>O<sub>3</sub>(cr) in 6.0 mol dm<sup>-3</sup> HCl(aq) by Fitzgibbon *et al.*,<sup>133</sup> combined with an estimated value of the enthalpy of solution of Eu(cr) in the same solvent. The value derived from the work by Yashvili and Gvelesiani,<sup>101</sup> who measured the enthalpies of solution of Eu(cr) and Eu<sub>2</sub>O<sub>3</sub>(cr) in 1.0 mol dm<sup>-3</sup> HCl(aq), deviates considerably, even when recalculated with an estimated enthalpy of solution of Eu(cr) in 1.0 HCl(aq). Also the value derived from the measurements by Hennig *et al.*,<sup>134</sup> who measured the enthalpy of solution of Eu<sub>2</sub>O<sub>3</sub> in 4.0 mol dm<sup>-3</sup> HCl(aq), disagrees, already evident from the much lower enthalpy of solution compared to Fitzgibbon *et al.*<sup>133</sup>

The enthalpy of formation of cubic Eu<sub>2</sub>O<sub>3</sub> was derived by Cordfunke and Konings<sup>34</sup> from two different sources. Fitzgibbon *et al.*<sup>133</sup> determined  $\Delta_{trs}H^\circ(\text{monoclinic/cubic}) = -(11.13 \pm 1.17) \text{ kJ mol}^{-1}$  by solution calorimetry in four different solvents, resulting in  $\Delta_f H^\circ(298.15 \text{ K}) = -(1661.1 \pm 6.3) \text{ kJ mol}^{-1}$ . An almost identical value,  $\Delta_f H^\circ(298.15 \text{ K}) = -(1663.3 \pm 5.9) \text{ kJ mol}^{-1}$ , was derived from the enthalpy of solution of cubic Eu<sub>2</sub>O<sub>3</sub> in 4.0 mol dm<sup>-3</sup> HCl(aq) by Stuve<sup>135</sup> and the enthalpy of solution of Eu(cr) by Fitzgibbon *et al.*<sup>133</sup> This latter value was preferred over Stube's value.

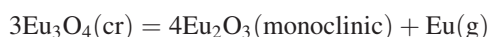
### 3.11. Eu<sub>3</sub>O<sub>4</sub>(cr)

#### 3.11.1. Polymorphism and melting point

Trieuropium tetraoxide has an orthorhombic crystal structure (space group Pnam). Bedford and Catalano<sup>136</sup> suggest that Eu<sub>3</sub>O<sub>4</sub> melts congruently at 2273 K.

#### 3.11.2. Heat capacity and entropy

The low-temperature heat capacity of Eu<sub>3</sub>O<sub>4</sub> has not been measured. The standard entropy can be derived from the second-law reaction entropy derived from vaporisation measurements of the reaction:



for which  $\Delta S^\circ = (118 \pm 3.4) \text{ J K}^{-1} \text{ mol}^{-1}$  at 1810 K was found by Haschke and Eick,<sup>137</sup> yielding  $S^\circ(298.15 \text{ K}) = (314.6 \pm 3.1) \text{ J K}^{-1} \text{ mol}^{-1}$ . This entropy value at 298.15 K is very close the sum of the oxides B-Eu<sub>2</sub>O<sub>3</sub> and EuO,  $(307.9 \pm 2.2) \text{ J K}^{-1} \text{ mol}^{-1}$ . We select

$$S^\circ(298.15 \text{ K}) = (315 \pm 4) \text{ J K}^{-1} \text{ mol}^{-1}$$

The high-temperature enthalpy increment of Eu<sub>3</sub>O<sub>4</sub> has been estimated by Haschke and Eick<sup>137</sup> as

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 182.464 + 26.108 \times 10^{-3}(T/\text{K}).$$

It should be noted that in the range 300–900 K, this equation is substantially higher than the sum of the oxides B-Eu<sub>2</sub>O<sub>3</sub> and EuO.

### 3.11.3. Enthalpy of formation

The enthalpy of formation of Eu<sub>3</sub>O<sub>4</sub> has been derived from the enthalpy of reaction derived by Haschke and Eick<sup>137</sup> of the vaporisation reaction (see Sec. 3.11.2),  $\Delta_r H = (360.7 \pm 5.9) \text{ kJ mol}^{-1}$ , resulting in:

$$\Delta_f H(\text{Eu}_3\text{O}_4, \text{cr}, 298.15 \text{ K}) = (2276 \pm 10) \text{ kJ mol}^{-1}$$

### 3.12. EuO(cr)

#### 3.12.1. Polymorphism and melting point

Europium monoxide has a face-centered cubic structure (space group Fm $\bar{3}$ m).<sup>138</sup> McMasters *et al.*<sup>139</sup> observed no phase transitions up to 1724 K. According to the phase diagram of the Eu-O system proposed by McCarthy and White<sup>140</sup> EuO decomposes peritectically at about 1793 K. Shafer *et al.*<sup>141</sup> suggest that EuO exhibits hypo- as well as hyperstiochiometry composition ranges, and that the liquidus of the oxygen rich compositions increases to  $(2238 \pm 10) \text{ K}$ , in agreement with the values reported by Reed and Fahey,<sup>142</sup>  $(2253 \pm 20) \text{ K}$ , and Bedford and Catalano,<sup>136</sup>  $(2263 \pm 30) \text{ K}$ .

#### 3.12.2. Heat capacity and entropy

The low-temperature heat capacity of EuO has been measured by Teany and Moruzzi<sup>143</sup> from 16 to 300 K. They observed a sharp anomaly with Curie temperature  $T_c = 69.3 \text{ K}$ , also found by Shafer *et al.*<sup>141</sup> by electrical conductivity measurements. The total entropy of the transition was found to be corresponding to the theoretical value  $R \ln(8)$ . The standard entropy of EuO has been derived from these measurements:

$$S^\circ(298.15 \text{ K}) = (83.6 \pm 0.8) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment has been measured by McMasters *et al.*<sup>139</sup> The two data sets are in good agreement and the high-temperature data can be represented by the polynomial equation (419 to 1724 K):

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 46.5453 + 7.360 \times 10^{-3}(T/\text{K})^2$$

which is a refit of the experimental data, omitting several extreme outliers.

### 3.12.3. Enthalpy of formation

The enthalpy of formation of EuO<sub>1.02</sub> was determined by Burnett<sup>144</sup> by solution calorimetry yielding  $\Delta_f H^\circ(298.15 \text{ K}) = -(607.0 \pm 4.1) \text{ kJ mol}^{-1}$ , and Huber and Holley Jr.,<sup>145,146</sup> by oxygen combustion calorimetry yielding  $\Delta_f H^\circ(298.15 \text{ K}) = -(599.6 \pm 2.1) \text{ kJ mol}^{-1}$ , in good agreement. Assuming that EuO<sub>1.02</sub> is a ideal solid solution (0.96EuO + 0.04EuO<sub>1.5</sub>) these values have been recalculated and the mean value has been selected,

$$\Delta_f H^\circ(\text{EuO}, \text{cr}, 298.15 \text{ K}) = -(593.2 \pm 5.0) \text{ kJ mol}^{-1}.$$

3.13. Gd<sub>2</sub>O<sub>3</sub>(cr,l)

## 3.13.1. Polymorphism and melting point

At room temperature, gadolinium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ), though the monoclinic B-type structure can also be maintained to room temperature. The C  $\rightarrow$  B transformation, which is sluggish but reversible, was found at 1298 K by Shafer and Roy,<sup>147</sup> 1523 K by Roth and Schneider<sup>121</sup> and 1473 K by Warshaw and Roy,<sup>89</sup> the latter being selected here because of the careful analysis made in that work. In view of the approximate nature, no attempt has been made to correct these values to ITS-90.

Foex and Traverse<sup>10</sup> determined the temperatures of the B  $\rightarrow$  A, A  $\rightarrow$  H and H  $\rightarrow$  X transformations by thermal analysis, without giving numerical results. From the thermogram we deduce the values  $\sim$ 2403,  $\sim$ 2423, and  $\sim$ 2623 K. They also determined the transformation of H-type Gd<sub>2</sub>O<sub>3</sub> to the X-type Ln<sub>2</sub>O<sub>3</sub> structure at  $\sim$ 2623 K using XRD. These values must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +15 K. Lopato *et al.*<sup>11</sup> reported that the transformations B  $\rightarrow$  A and A  $\rightarrow$  H occur in the temperature range 2443–2473 K and that H  $\rightarrow$  X transformation occurs close to the melting point (2633–2653 K). Barkhatov *et al.*<sup>148</sup> reported the transformation temperature for B  $\rightarrow$  A and A  $\rightarrow$  H as (2443  $\pm$  10) K and (2481  $\pm$  10) K from drop calorimetric measurements and (2436  $\pm$  10) K and (2470  $\pm$  15) K from electrical conductivity measurements, respectively. No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of  $-1$  K needs to be applied to convert to ITS-90. We select  $T_{trs} = (2430 \pm 30)$  K for the B  $\rightarrow$  A transformation,  $T_{trs} = (2470 \pm 30)$  K for the A  $\rightarrow$  H transformation and  $T_{trs} = (2538 \pm 20)$  K for the H  $\rightarrow$  X transformation. The measurements of the melting temperature of Gd<sub>2</sub>O<sub>3</sub> have been summarized in Table 16, which is based on the IUPAC review by Coutures and Rand.<sup>15</sup> The selected melting point is (2693  $\pm$  15) K.

TABLE 16. Temperature of melting of gadolinium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Wisnyi and Pijanowski <sup>111</sup>	2603 $\pm$ 20	2620 $\pm$ 20
Curtis and Johnson <sup>112</sup>	2623 $\pm$ 50	2626 $\pm$ 50
Foex <sup>19</sup>	2668	2683
Mordovin <i>et al.</i> <sup>20</sup>	2597 $\pm$ 30	2599 $\pm$ 30
Noguchi and Mizuno <sup>21</sup>	2603 $\pm$ 20	2606 $\pm$ 20
Spiridonov <i>et al.</i> <sup>149</sup>	2573	2576
Treswjatskii <i>et al.</i> <sup>22</sup>	2653 $\pm$ 20	2652 $\pm$ 20
Coutures <i>et al.</i> <sup>23</sup>	2713 $\pm$ 10	2712 $\pm$ 10
Mizuno <i>et al.</i> <sup>24</sup>	2667	2655
Yoshimura <i>et al.</i> <sup>25</sup>	2666	2672 $\pm$ 15
Mizuno <i>et al.</i> <sup>150</sup>		2686 $\pm$ 25
Shevthenko and Lopato <sup>13</sup>	2683	2682
Salikhov and Kan <sup>95</sup>		2666 $\pm$ 10
Kang <i>et al.</i> <sup>151</sup>		2701 $\pm$ 20
Selected value:		2693 $\pm$ 15

## 3.13.2. Heat capacity and entropy

The low-temperature heat capacity of C-Gd<sub>2</sub>O<sub>3</sub> has been measured by Justice and Westrum Jr.<sup>113</sup> from T = (7 to 346) K and Stewart *et al.*<sup>152</sup> from T = (1.4 to 18) K, the results being in acceptable agreement in the overlapping temperature range. The results show a thermal anomaly, which is related to the removal of the ground state degeneracy. Stewart *et al.*<sup>152</sup> concluded from the relatively large crystal field splitting needed to reproduce the peak and from EPR measurements that the anomaly in the cubic form is not a Schottky-type transition, as suggested by Miller *et al.*,<sup>153</sup> but is related to a magnetic ordering. The entropy of the transition is, however, close to  $2R\ln(2J+1) = 2R\ln(8)$  expected for the contribution of the ground state <sup>8</sup>F<sub>7/2</sub> multiplet. The results give for the entropy of C-Gd<sub>2</sub>O<sub>3</sub> at T = 298.15 K:

$$S^\circ(\text{Gd}_2\text{O}_3, \text{cubic}, 298.15 \text{ K}) = (150.6 \pm 0.2) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The low-temperature heat capacity of B-Gd<sub>2</sub>O<sub>3</sub> was measured by Konings *et al.*<sup>127</sup> from 5 to 400 K and by Rosenblum *et al.*<sup>154</sup> from 1.6 to 12.5 K. The latter authors found a lambda-type transition with a maximum at T = 3.80 K. The transition entropy is almost identical to the theoretical value  $2R\ln(8) = 4.16R$ . The absolute entropy at T = 298.15 K is

$$\begin{aligned} S^\circ(\text{Gd}_2\text{O}_3, \text{monoclinic}, 298.15 \text{ K}) \\ = (157.1 \pm 0.2) \text{ J K}^{-1} \text{ mol}^{-1}. \end{aligned}$$

The high-temperature enthalpy increment of C-Gd<sub>2</sub>O<sub>3</sub> has been determined by Curtis and Johnson,<sup>112</sup> Pankratz and King,<sup>128</sup> and Tsagareishvili *et al.*<sup>155</sup> The results are in good agreement, as shown in Fig. 9. Our recommended heat capacity equation for C-Gd<sub>2</sub>O<sub>3</sub> is based on the combined results of

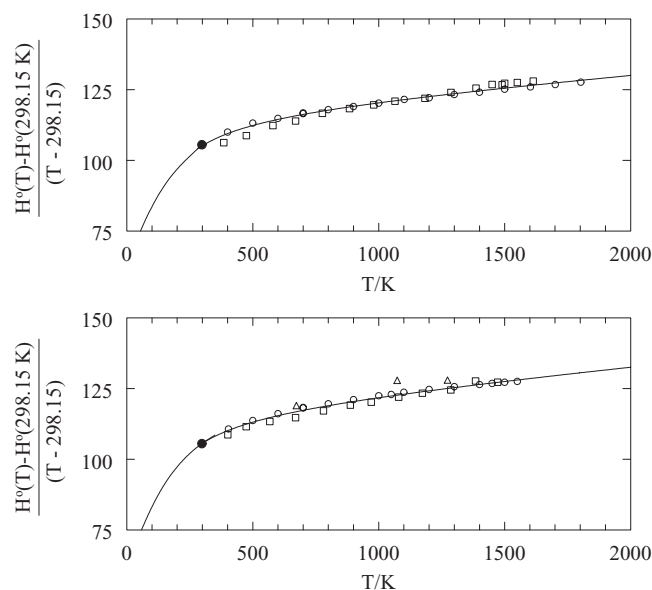


FIG. 9. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of B-Gd<sub>2</sub>O<sub>3</sub> (top) and C-Gd<sub>2</sub>O<sub>3</sub> (bottom);  $\circ$ , Pankratz and King<sup>128</sup>;  $\square$ , Tsagareishvili *et al.*<sup>155</sup>;  $\triangle$ , Curtis and Johnson<sup>112</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice and Westrum Jr.<sup>113</sup> for C-Gd<sub>2</sub>O<sub>3</sub> and Konings *et al.*<sup>127</sup> for B-Gd<sub>2</sub>O<sub>3</sub>; the curves show the recommended equations.



the latter two studies, constrained to  $C_p^\circ = 105.52 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements:<sup>113</sup>

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 114.8086 + 17.2911 \times 10^{-3}(\text{T/K}) - 1.28397 \times 10^6(\text{T/K})^{-2}.$$

The results of Pankratz and King<sup>128</sup> and Tsagareishvili *et al.*<sup>155</sup> for B-Gd<sub>2</sub>O<sub>3</sub> differ significantly at the lowest temperatures. We have fitted the combined results into a polynomial fit, constrained to  $C_p^\circ = 105.1 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements<sup>113</sup> to give for the heat capacity of B-Gd<sub>2</sub>O<sub>3</sub>:

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 114.6104 + 15.2344 \times 10^{-3}(\text{T/K}) - 1.24917 \times 10^6(\text{T/K})^{-2}.$$

The entropy of the C → B transition was estimated from high-pressure studies by Hoekstra<sup>116</sup> as  $6.3 \text{ J K}^{-1} \text{ mol}^{-1}$  using the Clausius-Clapeyron equation. This value is confirmed by the difference in the entropies of the B- and C-modifications of Gd<sub>2</sub>O<sub>3</sub> at 298.15 K (see above),  $6.5 \text{ J K}^{-1} \text{ mol}^{-1}$ . Unlike Sm<sub>2</sub>O<sub>3</sub> and Eu<sub>2</sub>O<sub>3</sub> the transition enthalpy cannot be derived from experimental enthalpy data, as the difference in the enthalpies of formation at 298.15 K is not known. Barkhatov *et al.*<sup>148</sup> reported enthalpy increment measurements for Gd<sub>2</sub>O<sub>3</sub> from which the enthalpies of the B → A and A → H transformations were derived as  $(6.3 \pm 3.3) \text{ kJ mol}^{-1}$  and  $(34.7 \pm 3.3) \text{ kJ mol}^{-1}$ . Heat-capacity or enthalpy measurements have not been reported for the X and liquid phases of Gd<sub>2</sub>O<sub>3</sub>. The properties of these modifications have been estimated in a similar way as for La<sub>2</sub>O<sub>3</sub> and Ce<sub>2</sub>O<sub>3</sub>. We thus obtain for the transition enthalpies

$$\begin{aligned} \Delta_{trs}H^\circ(\text{C} \rightarrow \text{B}) &= (9 \pm 2) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{B} \rightarrow \text{A}) &= (6.3 \pm 3.3) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (34.7 \pm 3.3) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (20 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (92 \pm 10) \text{ kJ mol}^{-1}. \end{aligned}$$

For the heat capacity of the high temperature A, H, and X modifications and liquid Gd<sub>2</sub>O<sub>3</sub> we estimate

$$\begin{aligned} C_p^\circ(\text{A}, \text{T}) &= 142 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{H}, \text{T}) &= C_p^\circ(\text{X}, \text{T}) = 130 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{liq}, \text{T}) &= 140 \text{ J K}^{-1} \text{ mol}^{-1}, \end{aligned}$$

### 3.13.3. Enthalpy of formation

The enthalpy of formation of monoclinic Gd<sub>2</sub>O<sub>3</sub> has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Gd}_2\text{O}_3, \text{ cr}, 298.15 \text{ K}) = -(1819.7 \pm 3.6) \text{ kJ mol}^{-1}.$$

This value is solely based on the combustion calorimetric study by Huber, Jr. and Holley, Jr.,<sup>156</sup> who used 97.05 mass %

TABLE 17. The enthalpy of formation of Gd<sub>2</sub>O<sub>3</sub>(cr) at 298.15 K;  $\Delta H_f^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of Gd(cr) and Gd<sub>2</sub>O<sub>3</sub>(cr) in HCl(aq), respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_f^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Huber Jr. and Holley Jr. <sup>156</sup>	C			$-1819.7 \pm 3.6$
Spedding <i>et al.</i> <sup>99</sup>	C			$-1786.2$
Yashvili and Gvelesiani <sup>101</sup>	S (6.0)	$-694.5 \pm 1.7$	$-422.6 \pm 1.3$	$-1826.3 \pm 3.7$
		$[-694.9 \pm 1.0]^b$		$-1829.5 \pm 2.6$
				$-1828.2 \pm 3.6^c$
Selected value:				$-1819.7 \pm 3.6$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ .

<sup>b</sup>Merli *et al.*<sup>39</sup>

<sup>c</sup>Cycle based on GdCl<sub>3</sub>.<sup>39</sup>

pure gadolinium metal (Table 17). The combustion value by Spedding *et al.*<sup>99</sup> is much less accurate due to incomplete combustion and lack of analytical characterization of the sample of Gd(cr). The values derived from the solution calorimetric study by Yashvili and Gvelesiani,<sup>101</sup> who determined the enthalpies of solution of Gd(cr) and Gd<sub>2</sub>O<sub>3</sub>(cr) in  $6.0 \text{ mol dm}^{-3} \text{ HCl(aq)}$ , are in reasonable agreement, but as discussed by Cordfunke and Konings,<sup>34</sup> they are considered significantly less accurate, especially in view of the difficulties of the slow dissolution of Gd<sub>2</sub>O<sub>3</sub>(cr) in HCl(aq).

## 3.14. TbO<sub>2</sub>(cr)

### 3.14.1. Structure

TbO<sub>2</sub> has a face-centered cubic structure (space group  $Fm\bar{3}m$ ) at room temperature. The upper stability of this phase has not been studied in detail. The phase diagram suggested by Lowe and Eyring<sup>157</sup> indicates that the TbO<sub>2-x</sub> is stable up to about 1400 K.

### 3.14.2. Heat capacity and entropy

No measurements of the heat capacity of TbO<sub>2</sub> have been made. The standard entropy has been estimated as

$$S^\circ(298.15 \text{ K}) = (86.9 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

from the systematics in the lanthanide and actinide oxides. In a similar manner the high temperature heat capacity was estimated as

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 73.259 + 13.2023 \times 10^{-3}(\text{T/K}) - 1.0424 \times 10^6(\text{T/K})^{-2}.$$

### 3.14.3. Enthalpy of formation

The enthalpy of formation of TbO<sub>2</sub> was determined by Stubblefield *et al.*<sup>84</sup> and Fitzgibbon and Holley Jr.<sup>158</sup> using solution calorimetry. Both research groups reported results for samples with varying O/Tb ratios. After recalculation and extrapolation to O/Tb = 2.0, as shown in Fig. 5, we obtain for the selected enthalpy of formation:

$$\Delta_f H^\circ(\text{TbO}_2, \text{ cr}, 298.15 \text{ K}) = -(972.2 \pm 5.0) \text{ kJ mol}^{-1}.$$

TABLE 18. The enthalpy of formation of phases in the TbO<sub>2</sub>-TbO<sub>1.5</sub> system

Composition	$\Delta_f H^\circ(298.15 \text{ K})$	Authors
	$\text{kJ mol}^{-1}$	
TbO <sub>1.510</sub>	$-(933.4 \pm 3.6)$	Fitzgibbon and Holley, Jr. <sup>158</sup>
TbO <sub>1.709</sub>	$-(952.3 \pm 3.4)$	Fitzgibbon and Holley, Jr. <sup>158</sup>
TbO <sub>1.710</sub>	$-(953.1 \pm 3.9)$	Stubblefield <i>et al.</i> <sup>84</sup>
TbO <sub>1.800</sub>	$-(966.1 \pm 3.3)$	Stubblefield <i>et al.</i> <sup>84</sup>
TbO <sub>1.817</sub>	$-(961.9 \pm 3.5)$	Fitzgibbon and Holley, Jr. <sup>158</sup>
TbO <sub>1.975</sub>	$-(970.5 \pm 2.8)$	Fitzgibbon and Holley, Jr. <sup>158</sup>

The enthalpies of formation (recalculated) of the compositions with various O/Tb ratios used for this calculation are given in Table 18.

### 3.15. Tb<sub>6</sub>O<sub>11</sub>(cr), Tb<sub>11</sub>O<sub>20</sub>(cr), Tb<sub>4</sub>O<sub>7</sub>(cr), Tb<sub>7</sub>O<sub>12</sub>(cr)

#### 3.15.1. Structure

In addition to the TbO<sub>2-x</sub> phase, a number of defined stoichiometric Tb<sub>n</sub>O<sub>m</sub> phases were reported to exist: TbO<sub>1.833</sub> (Tb<sub>6</sub>O<sub>11</sub>) as well as TbO<sub>1.818</sub> (Tb<sub>11</sub>O<sub>20</sub>) have a triclinic crystal structure (P $\bar{1}$ ), TbO<sub>1.75</sub> (Tb<sub>4</sub>O<sub>7</sub>) has a cubic crystal structure (Fm $\bar{3}$ m), and TbO<sub>1.714</sub> (Tb<sub>7</sub>O<sub>12</sub>) has a trigonal crystal structure (R $\bar{3}$ ).

#### 3.15.2. Heat capacity and entropy

Heat capacity data for the stoichiometric Tb<sub>n</sub>O<sub>m</sub> phases are limited to the measurement for Tb<sub>4</sub>O<sub>7</sub> by Hill<sup>159</sup> from 0.5 to 22 K, revealing an antiferromagnetic transition at and 7.85 K. For that reason we recommend that the standard entropies and the high temperature heat capacity are interpolated between TbO<sub>2</sub> and Tb<sub>2</sub>O<sub>3</sub>.

#### 3.15.3. Enthalpy of formation

The enthalpies of formation of a number of phases with varying O/Tb ratios have been measured by Stubblefield *et al.*<sup>84</sup> and Fitzgibbon and Holley Jr.<sup>158</sup> and the recalculated values are listed in Table 18. As shown in Fig. 5, the experimental enthalpies of formation of the Tb<sub>n</sub>O<sub>m</sub> phases are in good agreement and fit to a regular trend, slightly more non-linear compared to the Pr-O system. We have interpolated the enthalpies of formation of the Tb<sub>n</sub>O<sub>m</sub> phases from this trend as

$$\begin{aligned} \Delta_f H^\circ(\text{TbO}_{1.833}, \text{cr}, 298.15 \text{ K}) &= -(945.8 \pm 5.0) \text{ kJ mol}^{-1}, \\ \Delta_f H^\circ(\text{TbO}_{1.818}, \text{cr}, 298.15 \text{ K}) &= -(957.8 \pm 5.0) \text{ kJ mol}^{-1}, \\ \Delta_f H^\circ(\text{TbO}_{1.750}, \text{cr}, 298.15 \text{ K}) &= -(962.9 \pm 5.0) \text{ kJ mol}^{-1}, \\ \Delta_f H^\circ(\text{TbO}_{1.714}, \text{cr}, 298.15 \text{ K}) &= -(963.8 \pm 5.0) \text{ kJ mol}^{-1}. \end{aligned}$$

### 3.16. Tb<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.16.1. Polymorphism and melting point

At room temperature, terbium sesquioxide has the rare earth cubic C-type structure (space group *Ia* $\bar{3}$ ). Warshaw and Roy<sup>89</sup> found the C  $\rightarrow$  B transformation at 2148 K. Foex and Traverse<sup>10</sup> determined this transformation temperature as 1823 K.

 TABLE 19. Temperature of melting of terbium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{\text{fus}} \text{ K}$	
	Reported	ITS-90
Foex <sup>19</sup>	2663	2678
Mordovin <i>et al.</i> <sup>20</sup>	$2566 \pm 30$	$2569 \pm 30$
Noguchi and Mizuno <sup>21</sup>	$2577 \pm 20$	$2580 \pm 20$
Treswjatskii <i>et al.</i> <sup>22</sup>	$2643 \pm 20$	$2642 \pm 20$
Coutures <i>et al.</i> <sup>23</sup>	$2683 \pm 10$	$2682 \pm 10$
Shevthenko and Lopato <sup>13</sup>	2673	2672
Selected value:		$2682 \pm 15$

The B  $\rightarrow$  H transformation was also determined by Foex and Traverse<sup>10</sup> using thermal analysis at 2473 K. These values must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +15 K. Lopato *et al.*<sup>11</sup> reported that the B  $\rightarrow$  H transformation occurs at 2448 K and the H  $\rightarrow$  X transformation close to the melting point (2613–2643 K). No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of  $-1$  K needs to be applied to convert to ITS-90. We select  $T_{\text{trs}} = (1823 \pm 30)$  K for the C  $\rightarrow$  B transformation and  $T_{\text{trs}} = (2488 \pm 30)$  K for the B  $\rightarrow$  H transformation, solely based on the work of Foex and Traverse.<sup>10</sup>

The measurements of the melting temperature of Tb<sub>2</sub>O<sub>3</sub> have been summarized in Table 19, which is based on the IUPAC review by Coutures and Rand.<sup>15</sup> The selected melting point is  $(2682 \pm 15)$  K.

#### 3.16.2. Heat capacity and entropy

The low-temperature heat capacity of Tb<sub>2</sub>O<sub>3</sub> has been measured by Hill<sup>159</sup> from 0.5 to 22 K, revealing an antiferromagnetic transition at 2.42 K. These results are too limited to derive the standard entropy and the selected value for this quantity is the value estimated by Konings<sup>109</sup> from the trend in the lattice component in the lanthanide sesquioxides and the calculated excess contribution calculated from the crystal field energies:

$$S^\circ(298.15 \text{ K}) = (159.2 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of C-Tb<sub>2</sub>O<sub>3</sub> has been determined by Pankratz *et al.*<sup>114</sup> from 414 to 1599 K. The results have been fitted to a polynomial equation constrained to  $C_p^\circ(298.15 \text{ K}) = 116.0 \text{ J K}^{-1} \text{ mol}^{-1}$ , estimated in a similar way as the entropy value. We thus derive for the heat capacity,

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 120.6682 + 22.17194 \times 10^{-3}(\text{T/K}) \\ &\quad - 1.00261 \times 10^6(\text{T/K})^{-2}. \end{aligned}$$

For the estimation of the enthalpy change of the C  $\rightarrow$  B phase transformation we accept the Hoekstra's estimate for the entropy of transition  $(6.3 \text{ J K}^{-1} \text{ mol}^{-1})$ ,<sup>116</sup> to give

$$\Delta_{\text{trs}} H^\circ(\text{C} \rightarrow \text{B}) = (12 \pm 4) \text{ kJ mol}^{-1}.$$

For the heat capacity of  $\text{B-Tb}_2\text{O}_3$  we estimate

$$C_p^\circ(\text{B}, T) = 152 \text{ J K}^{-1} \text{ mol}^{-1}.$$

Foex and Traverse<sup>10</sup> found that the  $\text{B} \rightarrow \text{H}$  transformation corresponds to a strong peak in the DTA analysis. Wu and Pelton<sup>33</sup> argued that the  $\text{B} \rightarrow \text{H}$  transformation has the same entropy change as the  $\text{C} \rightarrow \text{H}$  (see under  $\text{Er}_2\text{O}_3$ ) for which they suggested  $\Delta_{trs}S^\circ(\text{B} \rightarrow \text{H}) = 7.2 \text{ J K}^{-1} \text{ mol}^{-1}$  from analysis of  $\text{Ln}_2\text{O}_3\text{-Al}_2\text{O}_3$  phase diagrams. For the entropy of fusion in these systems they derived  $\Delta_{fus}S^\circ(\text{H} \rightarrow \text{liquid}) = 17.9 \text{ J K}^{-1} \text{ mol}^{-1}$  in a similar manner. These numbers would suggest an appreciably lower sum for the entropies of the transformation from  $\text{C}$  to liquid compared to  $\text{Gd}_2\text{O}_3$ , whereas an almost constant value could be expected, based on the known data for the lanthanide trifluorides.<sup>160</sup> The selected values are consistent with the latter assumption:

$$\begin{aligned}\Delta_{trs}H^\circ(\text{B} \rightarrow \text{H}) &= (55 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (83 \pm 8) \text{ kJ mol}^{-1}.\end{aligned}$$

For the heat capacity of the high temperature modification  $\text{H}$  and the liquid phase of  $\text{Tb}_2\text{O}_3$  we estimate

$$\begin{aligned}C_p^\circ(\text{H}, T) &= 170 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{liq}, T) &= 182 \text{ J K}^{-1} \text{ mol}^{-1}.\end{aligned}$$

### 3.16.3. Enthalpy of formation

The enthalpy of formation of monoclinic cubic  $\text{Tb}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Tb}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1865.2 \pm 6.0) \text{ kJ mol}^{-1}.$$

This value is based on an analysis of two solution calorimetry studies, as shown in Table 20. Stubblefield *et al.*<sup>84</sup> measured the enthalpy of solution of  $\text{Tb}_2\text{O}_3(\text{cr})$  in  $6.0 \text{ mol dm}^{-3} \text{ HNO}_3(\text{aq})$ . However, the thermochemical reaction cycle was considered to be not highly reliable, since not all the auxiliary thermodynamic data important for that calculation are known with sufficient accuracy. Fitzgibbon and Holley, Jr.<sup>158</sup> measured the enthalpy of formation of  $\text{TbO}_{1.510}(\text{cr})$  using a thermochemical cycle which involves the solution of terbium metal and the oxide in  $1.0 \text{ mol dm}^{-3} \text{ HCl}(\text{aq})$  and

$6.0 \text{ mol dm}^{-3} \text{ HNO}_3(\text{aq})$ , respectively. Different solvents were used to avoid reduction of  $\text{HNO}_3(\text{aq})$  by terbium metal and oxidation of some of the  $\text{HCl}$  by the oxide. In order to combine the results in the two solvents, the enthalpy of solution of terbium carbonate was measured in both solvents. The enthalpy of formation of  $\text{TbO}_{1.50}$  was obtained by extrapolation (Fig. 5), being in excellent agreement with the value by Stubblefield *et al.*<sup>84</sup>

## 3.17. $\text{Dy}_2\text{O}_3(\text{cr,l})$

### 3.17.1. Polymorphism and melting point

At room temperature, dysprosium sesquioxide has the rare earth cubic  $\text{C}$ -type structure (space group  $Ia\bar{3}$ ). Warshaw and Roy<sup>89</sup> found the  $\text{C} \rightarrow \text{B}$  transformation at 2423 K. Foex and Traverse<sup>10</sup> and Lopato *et al.*<sup>11</sup> determined this transformation temperature as 2223 K. In view of the approximate nature, no attempt has been made to correct these values to ITS-90. The  $\text{B} \rightarrow \text{H}$  transformation was determined by Foex and Traverse<sup>10</sup> using thermal analysis at  $T = 2473 \text{ K}$ . This values must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +15 K. Lopato *et al.*<sup>11</sup> reported that the  $\text{B} \rightarrow \text{H}$  transformation occurs at 2463 K. No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of  $-1 \text{ K}$  needs to be applied to convert to ITS-90. We select  $T_{trs} = (2223 \pm 30) \text{ K}$  for the  $\text{C} \rightarrow \text{B}$  transformation and  $T_{trs} = (2488 \pm 30) \text{ K}$  for the  $\text{B} \rightarrow \text{H}$  transformation, solely based on the work of Foex and Traverse.<sup>10</sup>

The measurements of the melting temperature of  $\text{Dy}_2\text{O}_3$  have been summarized in Table 21, which is based on the IUPAC review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2680 \pm 15) \text{ K}$ .

### 3.17.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Dy}_2\text{O}_3$  has been measured by Justice and Westrum, Jr.<sup>161</sup> from 10 to 350 K. At the lower temperature range, the tail of a thermal anomaly was observed, which is related to the removal of the ground state degeneracy. Extrapolation of these results to  $T = 0 \text{ K}$ , and accounting for the  $2R \ln(2)$  contribution of the lowest level of the  ${}^6\text{H}_{15/2}$  ground state multiplet, gives for the entropy at

TABLE 20. The enthalpy of formation of  $\text{Tb}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Tb}(\text{cr})$  and  $\text{Tb}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Stubblefield <i>et al.</i> <sup>84</sup>	S (1.0) <sup>b</sup>		$-395.0 \pm 2.5$	$-1864.5 \pm 8.4$
Fitzgibbon and Holley, Jr. <sup>158</sup>	S (1.0) <sup>b</sup>		$-392.5 \pm 5.0$	$-1865.2 \pm 6.0$
Selected value:				$-1865.2 \pm 6.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ .

<sup>b</sup>Solvent was  $\text{HNO}_3(\text{aq})$ .

TABLE 21. Temperature of melting of dysprosium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Wisnyi and Pijanowski <sup>111</sup>	2613 ± 20	2629 ± 20
Foex <sup>19</sup>	2663	2678
Mordovin <i>et al.</i> <sup>20</sup>	2566 ± 30	2569 ± 30
Noguchi and Mizuno <sup>21</sup>	2501 ± 20	2504 ± 20
Treswjatskii <i>et al.</i> <sup>22</sup>	2633 ± 20	2632 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2683 ± 10	2682 ± 10
Mizuno <i>et al.</i> <sup>24</sup>	2628 ± 20	2615 ± 20
Shevthenko and Lopato <sup>13</sup>	2673	2672
Selected value:		2680 ± 15

$T = 298.15 \text{ K}$ :

$$S^\circ(298.15 \text{ K}) = (149.8 \pm 0.15) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of  $\text{Dy}_2\text{O}_3$  has been measured by Pankratz and Kelly<sup>65</sup> from 417 to 1801 K. The results are in good agreement with the low temperature heat capacity data. Pankratz and Kelly<sup>65</sup> observed a minor thermal anomaly between 1823 and 1863 K, whose origin is not clear. Since the enthalpy effect is very small ( $<1 \text{ kJ mol}^{-1}$ ) and has not yet been confirmed by other techniques, and similar unconfirmed transitions have been reported by this group of researchers for other rare-earth sesquioxides, we have neglected it. The data have been fitted to a polynomial equation, constrained to  $C_p^\circ = 116.27 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements,<sup>161</sup> yielding for the heat capacity:

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 121.2302 + 15.27609 \times 10^{-3}(T/K) - 0.84580 \times 10^6(T/K)^{-2}.$$

For the heat capacity of B- $\text{Dy}_2\text{O}_3$  we estimate

$$C_p^\circ(\text{B}, T) = 155 \text{ J K}^{-1} \text{ mol}^{-1}.$$

For transition and fusion entropies we assume the same values as for the iso-structural changes in  $\text{Tb}_2\text{O}_3$ , yielding

$$\Delta_{trs}H^\circ(\text{C} \rightarrow \text{B}) = (14 \pm 5) \text{ kJ mol}^{-1},$$

$$\Delta_{trs}H^\circ(\text{B} \rightarrow \text{H}) = (55 \pm 8) \text{ kJ mol}^{-1},$$

$$\Delta_{fus}H^\circ = (83 \pm 8) \text{ kJ mol}^{-1}.$$

For the heat capacity of the high temperature H modification and liquid phase of  $\text{Dy}_2\text{O}_3$  we estimate

$$C_p^\circ(\text{H}, T) = 173 \text{ J K}^{-1} \text{ mol}^{-1}$$

$$C_p^\circ(\text{liq}, T) = 188 \text{ J K}^{-1} \text{ mol}^{-1}.$$

### 3.17.3. Enthalpy of formation

The enthalpy of formation of cubic  $\text{Dy}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Dy}_2\text{O}_3, \text{cr}, 298.15\text{K}) = -(1863.4 \pm 5.0) \text{ kJ mol}^{-1}.$$

The selected value is based on the study by Huber, Jr. *et al.*<sup>162</sup> who used both solution calorimetry and oxygen bomb combustion calorimetry (Table 22). The results from the earlier publication by the same group<sup>163</sup> based on the combustion calorimetric measurements, seems to be less accurate, since the sample of dysprosium metal was less pure than that used in the later investigation.

## 3.18. $\text{Ho}_2\text{O}_3(\text{cr,l})$

### 3.18.1. Polymorphism and melting point

At room temperature, holmium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ). Foex and Traverse<sup>10</sup> found the C  $\rightarrow$  B transformation at 2523 K and Lopato *et al.*<sup>11</sup> between 2463 K and 2493 K. The B  $\rightarrow$  H transformation was only determined by Foex and Traverse<sup>10</sup> using thermal analysis near 2573 K. The values of Foex and Traverse<sup>10</sup> values must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +15 K. We select  $T_{trs} = (2538 \pm 30) \text{ K}$  for the C  $\rightarrow$  B transformation and  $T_{trs} = (2588 \pm 30) \text{ K}$  for the B  $\rightarrow$  H transformation.

The measurements of the melting temperature of  $\text{Ho}_2\text{O}_3$  have been summarized in Table 23, which is based on the IUPAC review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2686 \pm 15) \text{ K}$ .

### 3.18.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Ho}_2\text{O}_3$  has been measured by Justice and Westrum Jr.<sup>161</sup> from 10 to 350 K.

TABLE 22. The enthalpy of formation of  $\text{Dy}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Dy}(\text{cr})$  and  $\text{Dy}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Huber, Jr. <i>et al.</i> <sup>163</sup>	C			$-1865.2 \pm 3.8$
Huber, Jr. <i>et al.</i> <sup>162</sup>	C			$-1862.9 \pm 4.2$
	S (4.0) <sup>b</sup>	$-695.3 \pm 2.9$	$-385.1 \pm 3.4$	$-1863.9 \pm 6.7$
Selected value:				$-1863.4 \pm 5.0$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ .

<sup>b</sup>solvent was  $\text{HNO}_3(\text{aq})$ .



TABLE 23. Temperature of melting of holmium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Foex <sup>19</sup>	2668	2683
Mordovin <i>et al.</i> <sup>20</sup>	2626 ± 30	2629 ± 30
Noguchi and Mizuno <sup>21</sup>	2603 ± 20	2606 ± 20
Treswjatskii <i>et al.</i> <sup>22</sup>	2643 ± 20	2642 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2693 ± 10	2692 ± 10
Mizuno <i>et al.</i> <sup>24</sup>	2638 ± 20	2625 ± 20
Shevthenko and Lopato <sup>13</sup>	2673	2672
Salikhov and Kan <sup>95</sup>		2664 ± 10
Selected value:		2686 ± 15

Extrapolation of these results to  $T = 0$  K, and accounting for the contribution of the lowest level of the  $^5I_{15/2}$  ground state multiplet, gives for the entropy at  $T = 298.15$  K:

$$S^\circ(298.15 \text{ K}) = (156.38 \pm 0.15) \text{ J K}^{-1} \text{ mol}^{-1}$$

The high-temperature enthalpy increment of  $\text{Ho}_2\text{O}_3$  has been measured by Pankratz *et al.*<sup>90</sup> from 400 to 1799 K and by Tsagareishvili and Gvelesiani<sup>164</sup> from 397 to 1621 K. The results of Pankratz *et al.*<sup>90</sup> tend to be somewhat lower at the lowest temperatures and moreover, do not agree particularly well with the low-temperature heat capacity data (Fig. 10). Our recommended heat-capacity equation is based on a polynomial fit of the results of both studies. The equation is constrained to  $C_p^\circ = 114.98 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements,<sup>113</sup> thus agreeing better with the results of Tsagareishvili and Gvelesiani<sup>164</sup> in the low temperature range. We thus obtain

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 121.9340 + 10.11623 \times 10^{-3}(T/\text{K}) - 0.886280 \times 10^6(T/\text{K})^{-2}.$$

For transition and fusion entropies we assume the same values as for the isostructural changes in  $\text{Tb}_2\text{O}_3$ , yielding

$$\begin{aligned} \Delta_{trs}H^\circ(C \rightarrow B) &= (16 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(B \rightarrow H) &= (57 \pm 8) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (83 \pm 8) \text{ kJ mol}^{-1}. \end{aligned}$$

TABLE 24. The enthalpy of formation of  $\text{Ho}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Ho}(\text{cr})$  and  $\text{Ho}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Huber Jr. <i>et al.</i> <sup>165</sup>	C			-1881.0 ± 5.0
Morss <i>et al.</i> <sup>166</sup>	S (4.0)		-379.1 ± 5.2 <sup>b</sup>	-1887.3 ± 9.5 <sup>c</sup>
		[-710.5 ± 7.1] <sup>c</sup>		-1900.3 ± 15.1
				-1885.7 ± 7.3 <sup>d</sup>
Selected value:				-1883.3 ± 8.2

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ ;

<sup>b</sup>uncertainty recalculated.

<sup>c</sup>using  $\Delta H_1^\circ = -704 \pm 4$  as suggested by Morss *et al.*<sup>166</sup>

<sup>d</sup>cycle based on  $\text{HoCl}_3$  as explained in the text.

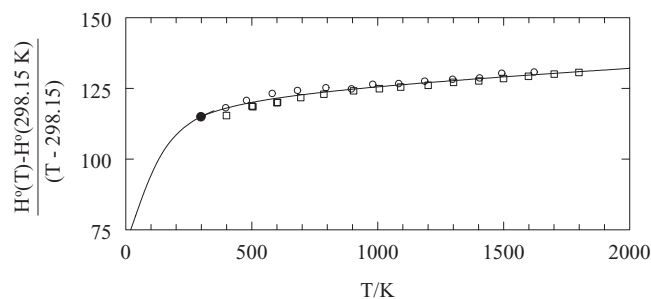


FIG. 10. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{Ho}_2\text{O}_3$ ;  $\circ$ , Tsagareishvili and Gvelesiani<sup>164</sup>;  $\square$ , Pankratz *et al.*<sup>90</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice and Westrum Jr.<sup>113</sup>; the curve shows the recommended equation.

For the heat capacity of the high temperature B and H modifications and of liquid  $\text{Ho}_2\text{O}_3$  we estimate

$$\begin{aligned} C_p^\circ(\text{Ho}_2\text{O}_3, \text{B}, T) &= 135 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{Ho}_2\text{O}_3, \text{H}, T) &= 149 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{Ho}_2\text{O}_3, \text{liq}, T) &= 162 \text{ J K}^{-1} \text{ mol}^{-1}. \end{aligned}$$

### 3.18.3. Enthalpy of formation

The enthalpy of formation of cubic  $\text{Ho}_2\text{O}_3$  has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Ho}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1883.3 \pm 8.2) \text{ kJ mol}^{-1}.$$

This value is based on the oxygen-bomb combustion calorimetric measurements by Huber Jr. *et al.*<sup>165</sup> and the solution calorimetric measurements of  $\text{Ho}_2\text{O}_3(\text{cr})$  in  $4.0 \text{ mol dm}^{-3} \text{ HCl}(\text{aq})$  by Morss *et al.*<sup>166</sup> For the analysis of the latter study, Morss *et al.*<sup>166</sup> used an estimated enthalpy of solution of  $\text{Ho}(\text{cr})$  in the same medium, as the measurement of the latter quantity by Stuve<sup>167</sup> was considered not reliable. Cordfunke and Konings<sup>34</sup> combined the enthalpy of solution of  $\text{Ho}_2\text{O}_3(\text{cr})$  in  $4.0 \text{ mol dm}^{-3} \text{ HCl}(\text{aq})$  with the enthalpy of solution of  $\text{HoCl}_3(\text{cr})$  in the same medium by Stuve,<sup>167</sup> obtaining a value in good agreement with the combustion value (see Table 24).

TABLE 25. Temperature of melting of erbium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Foex <sup>19</sup>	2673	2688
Mordovin <i>et al.</i> <sup>20</sup>	2661 ± 30	2664 ± 30
Noguchi and Mizuno <sup>21</sup>	2618 ± 20	2621 ± 20
Treswjatskii <i>et al.</i> <sup>22</sup>	2663 ± 20	2662 ± 20
Coutures <i>et al.</i> <sup>23</sup>	2693 ± 10	2692 ± 10
Mizuno <i>et al.</i> <sup>24</sup>	2648 ± 20	2636 ± 20
Shevthenko and Lopato <sup>13</sup>	2693	2692
Salikhov and Kan <sup>95</sup>		2686 ± 10
Selected value:		2690 ± 15

### 3.19. Er<sub>2</sub>O<sub>3</sub>(cr,l)

#### 3.19.1. Polymorphism and melting point

At room temperature, erbium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ). Foex and Traverse<sup>10</sup> found the C → H transformation near  $T = 2523$  K using thermal analysis. This value must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of +15 K. Lopato *et al.*<sup>11</sup> found the transformation at 2593 K. No information on the calibration of these measurements has been found, but assuming the data refer to IPTS-68, a correction of −1 K needs to be applied to convert to ITS-90. We select  $T_{trs} = (2538 \pm 30)$  K, solely based on the results of Foex and Traverse.<sup>10</sup>

The measurements of the melting temperature of Er<sub>2</sub>O<sub>3</sub> have been summarized in Table 25, which is based on the IUPAC review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2690 \pm 15)$  K.

#### 3.19.2. Heat capacity and entropy

The low-temperature heat capacity of Er<sub>2</sub>O<sub>3</sub> has been measured by Justice and Westrum Jr.<sup>161</sup> from 10 to 350 K. Extrapolation of these results to  $T = 0$  K, and accounting for the  $2R\ln(2)$  contribution of the lowest level of the <sup>5</sup>I<sub>15/2</sub> ground state multiplet. Tang *et al.*<sup>168</sup> found by calorimetric measurements in the temperature range 1.5 to 15 K that Er<sub>2</sub>O<sub>3</sub> orders antiferromagnetically with Néel temperature of 3.3 K. The entropy associated with the measured peak is close to 3/4 of  $2R\ln(2)$ . Since there are two nonequivalent Er<sup>3+</sup> sites in Er<sub>2</sub>O<sub>3</sub>, occurring in the ratio 3:1, the observed anomaly corresponds to the ordering of one of the sites, the ordering of the other occurring at lower temperature. The total entropy at  $T = 298.15$  K derived from these results is

$$S^\circ(298.15 \text{ K}) = (153.13 \pm 0.15) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of Er<sub>2</sub>O<sub>3</sub> has been measured by Pankratz *et al.*<sup>90</sup> from 399 to 1797 K and by Tsagareishvili and Gvelesiani<sup>169</sup> from 387 to 1625 K. The results of Pankratz *et al.*<sup>90</sup> tend to be somewhat lower at the lowest temperatures (Fig. 11). Our recommended heat-capacity equation is based on a polynomial fit of the results of both studies. The equation is constrained to  $C_p^\circ = 108.49 \text{ J K}^{-1}$

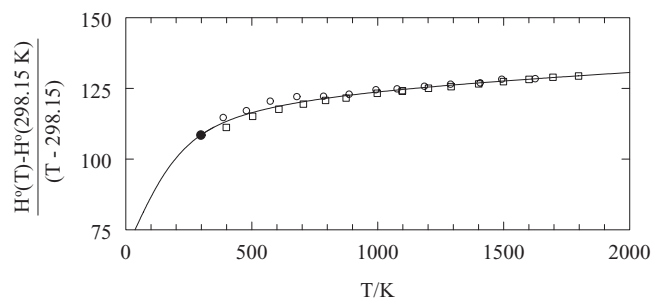


FIG. 11. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of Er<sub>2</sub>O<sub>3</sub>; ○, Tsagareishvili and Gvelesiani<sup>169</sup>; □, Pankratz *et al.*<sup>90</sup>; ●, value derived from the low-temperature measurements by Justice and Westrum Jr.<sup>161</sup>; the curve shows the recommended equation.

$\text{mol}^{-1}$  from the low-temperature measurements.<sup>161</sup> We thus obtain

$$C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = 123.2921 + 8.62245 \times 10^{-3} (T/\text{K}) - 1.54433 \times 10^{-6} (T/\text{K})^2.$$

Wu and Pelton<sup>33</sup> argued that the C → H transformation has a significant entropy effect, as changes in the liquidus slopes can be observed in the Er<sub>2</sub>O<sub>3</sub>–Al<sub>2</sub>O<sub>3</sub> and Ho<sub>2</sub>O<sub>3</sub>–Al<sub>2</sub>O<sub>3</sub> phase diagrams. From the limiting slope at the Ln<sub>2</sub>O<sub>3</sub> side and the change in slope at the transformation temperatures they derived  $\Delta_{trs}S^\circ(\text{C} \rightarrow \text{H}) = 7.2 \text{ J K}^{-1} \text{ mol}^{-1}$ , and  $\Delta_{trs}S^\circ(\text{H} \rightarrow \text{liquid}) = 17.9 \text{ J K}^{-1} \text{ mol}^{-1}$ . These numbers would suggest an appreciable lower sum for the entropies of the transformation from the C form to the liquid phase compared to the other cubic lanthanides, whereas an almost constant value could be expected, based on the known data for the lanthanide trifluorides.<sup>160</sup> Our selected values are based on the assumption that the entropy changes for the C → H transformation and fusion are identical to those of Y<sub>2</sub>O<sub>3</sub> as measured by Shpil'rain *et al.*<sup>170</sup> using drop calorimetry:

$$\begin{aligned} \Delta_{trs}H^\circ(\text{C} \rightarrow \text{H}) &= (25 \pm 5) \text{ kJ mol}^{-1} \\ \Delta_{fus}H^\circ &= (83 \pm 5) \text{ kJ mol}^{-1} \end{aligned}$$

For the heat capacity of the high temperature modification H and liquid Er<sub>2</sub>O<sub>3</sub> we estimate

$$\begin{aligned} C_p^\circ(\text{Er}_2\text{O}_3, \text{H}, T) &= 162 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{Er}_2\text{O}_3, \text{liq}, T) &= 176 \text{ J K}^{-1} \text{ mol}^{-1}. \end{aligned}$$

#### 3.19.3. Enthalpy of formation

The enthalpy of formation of cubic Er<sub>2</sub>O<sub>3</sub> has been assessed by Cordfunke and Konings<sup>34</sup> recently, and we accept the selected values from that work, since no new information has been published since

$$\Delta_f H^\circ(\text{Er}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1900.1 \pm 6.5) \text{ kJ mol}^{-1}.$$

This value is based on an analysis of the calorimetric studies listed in Table 26. Huber Jr. *et al.*<sup>171</sup> and Spedding *et al.*<sup>99</sup> determined the enthalpy of formation of Er<sub>2</sub>O<sub>3</sub>(cr) by oxygen bomb calorimetry. The derived values differ considerable, but

TABLE 26. The enthalpy of formation of  $\text{Er}_2\text{O}_3(\text{cr})$  at 298.15 K;  $\Delta H_1^\circ$  and  $\Delta H_2^\circ$  are the enthalpies of solution of  $\text{Er}(\text{cr})$  and  $\text{Er}_2\text{O}_3(\text{cr})$  in  $\text{HCl}(\text{aq})$ , respectively (after Cordfunke and Konings<sup>34</sup>)

Authors	Method <sup>a</sup>	$\Delta H_1^\circ/\text{kJ mol}^{-1}$	$\Delta H_2^\circ/\text{kJ mol}^{-1}$	$\Delta_f H^\circ/\text{kJ mol}^{-1}$
Huber Jr. <i>et al.</i> <sup>171</sup>	C			$-1897.8 \pm 3.8$
Spedding <i>et al.</i> <sup>99</sup>	C			$-1762.8$
Montgomery and Stuve <sup>172</sup>	S (1.40)	$[-705.6 \pm 1.4]^b$	$-370.6 \pm 3.7$	$-1898.2 \pm 4.6$
Morss <i>et al.</i> <sup>166</sup>	S (1.40)		$-364.6 \pm 1.9^c$	$-1904.2 \pm 3.4$
Selected value:				$-1900.1 \pm 6.5$

<sup>a</sup>C: combustion calorimetry; S: solution calorimetry; values in parentheses give the concentration of the solvent in  $\text{mol dm}^{-3}$ .

<sup>b</sup>Fuger and Morss.<sup>173</sup>

<sup>c</sup>Uncertainty recalculated.

the work of the former authors is considered to be much more reliable due to the fact that the erbium metal used was well characterised. This is supported by the good agreement with the enthalpy of formation values derived from solution calorimetry  $\sim 1.4 \text{ mol dm}^{-3} \text{ HCl}(\text{aq})$  by Montgomery and Stuve<sup>172</sup> and Morss *et al.*<sup>166</sup> These values were combined with the enthalpy of solution of  $\text{Er}(\text{cr})$  by Fuger *et al.*<sup>173</sup>

### 3.20. $\text{Tm}_2\text{O}_3(\text{cr,l})$

#### 3.20.1. Polymorphism and melting point

At room temperature, thulium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ). Foex and Traverse<sup>10</sup> found the  $B \rightarrow H$  transformation near  $T = 2573 \text{ K}$  using thermal analysis. This value must be converted to ITS-90 using the procedure outlined by Coutures and Rand,<sup>15</sup> leading to a correction of  $+15 \text{ K}$ . We thus select  $T_{\text{trs}} = (2588 \pm 30) \text{ K}$ .

The melting temperature of  $\text{Tm}_2\text{O}_3$  has been measured by Treswjatskii *et al.*<sup>22</sup> to be  $(2653 \pm 20) \text{ K}$ . However, the melting temperature determinations of the lanthanide sesquioxides by these authors are systematically low by 30–50 K compared to our selected values. Shevthenko and Lopato<sup>13</sup> measured the melting point to be  $T = 2683 \text{ K}$ , which would be  $T = 2682 \text{ K}$  on ITS-90. Coutures *et al.*<sup>23</sup> estimated the melting temperature as  $(2698 \pm 20) \text{ K}$  from the trend in the lanthanide sesquioxides. We select  $T_{\text{fus}} = (2682 \pm 30) \text{ K}$ .

#### 3.20.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Tm}_2\text{O}_3$  has been measured by Justice *et al.*<sup>174</sup> from 6 to 350 K. They obtained  $S^\circ(298.15 \text{ K}) - S^\circ(10 \text{ K}) = 138.70 \text{ J K}^{-1} \text{ mol}^{-1}$ , and extrapolated the results to  $T = 0 \text{ K}$  using lattice and electronic contributions. This gives

$$S^\circ(298.15 \text{ K}) = (139.7 \pm 0.4) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of  $\text{Tm}_2\text{O}_3$  has been measured by Pankratz *et al.*<sup>90</sup> from 397 to 1801 K and Tsagareishvili and Gvelesiani<sup>129</sup> from 397 to 1607 K. These data are in fair agreement, but both are somewhat low at the lowest temperatures compared to the low-temperature data

(Fig. 12). At high temperatures the results tend to diverge, the results of Tsagareishvili and Gvelesiani<sup>129</sup> being significantly higher. Moreover, the results of Pankratz *et al.*<sup>90</sup> indicate a minor thermal anomaly near 1680 K, whose origin is not clear. Since the enthalpy effect is very small ( $\sim 1.3 \text{ kJ mol}^{-1}$ ), and similar unconfirmed transitions have been reported by this group for other rare-earth sesquioxides, we have neglected it. Our recommended heat-capacity equation is based on a polynomial fit of the results of both studies, but since the results of Pankratz *et al.*<sup>90</sup> cover a wider temperature range, they dominate the high temperature part of the curve. The equation is constrained to  $C_p^\circ = 116.73 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements.<sup>161</sup> We thus obtain

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 128.4322 + 5.23209 \times 10^{-3}(T/\text{K}) - 1.17891 \times 10^6(T/\text{K})^{-2}.$$

The entropies of the  $C \rightarrow H$  transformation and melting are assumed to be the same as for  $\text{Er}_2\text{O}_3$ . This yields

$$\begin{aligned} \Delta_{\text{trs}} H^\circ(C \rightarrow H) &= (26 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{\text{fus}} H^\circ &= (83 \pm 8) \text{ kJ mol}^{-1}. \end{aligned}$$

For the heat capacity of the high temperature modification H and liquid  $\text{Tm}_2\text{O}_3$  we estimate

$$\begin{aligned} C_p^\circ(\text{Tm}_2\text{O}_3, \text{H}, T) &= 155 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{Tm}_2\text{O}_3, \text{liq}, T) &= 168 \text{ J K}^{-1} \text{ mol}^{-1}. \end{aligned}$$

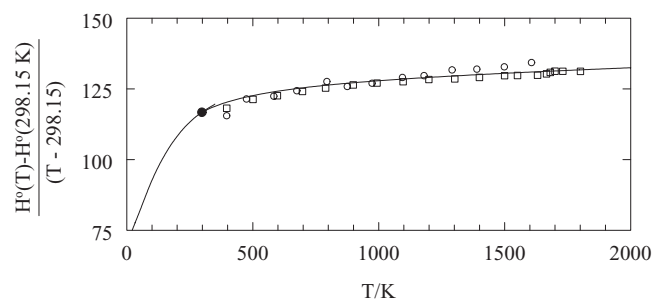


FIG. 12. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{Tm}_2\text{O}_3$ ;  $\circ$ , Tsagareishvili and Gvelesiani<sup>129</sup>;  $\square$ , Pankratz *et al.*<sup>90</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice *et al.*<sup>174</sup>; the curve shows the recommended equation.

### 3.20.3. Enthalpy of formation

There is only one determination of the standard molar enthalpy of formation of cubic thulium sesquioxide, which was performed by Huber Jr. *et al.*<sup>175</sup> using oxygen-bomb combustion calorimetry, and this value was selected by Cordfunke and Konings<sup>34</sup>:

$$\Delta_f H^\circ(\text{Tm}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1889.3 \pm 5.7) \text{ kJ mol}^{-1}.$$

It is the weighted mean of the results for two samples of Tm(cr) containing significantly different amounts of impurities. Also the extent of combustion varied significantly (88.67% to 99.67% of completion). After correction, Huber Jr. *et al.*<sup>175</sup> obtained the values  $\Delta_f H^\circ(298.15 \text{ K}) = -(1894.8 \pm 8.3) \text{ kJ mol}^{-1}$  and  $\Delta_f H^\circ(298.15 \text{ K}) = -(1884.3 \pm 7.9) \text{ kJ mol}^{-1}$ , respectively.

## 3.21. Yb<sub>2</sub>O<sub>3</sub>(cr,l)

### 3.21.1. Polymorphism and melting point

At room temperature, ytterbium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ). Foex and Traverse<sup>10</sup> found the B  $\rightarrow$  H transformation very close to the melting point. Lopato *et al.*<sup>11</sup> reported this transformation 20 K below the melting point. We thus select  $T_{\text{trs}} = (2687 \pm 20) \text{ K}$ .

The measurements of the melting temperature of Yb<sub>2</sub>O<sub>3</sub> have been summarized in Table 27, which is based on the IUPAC review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2707 \pm 15) \text{ K}$ .

### 3.21.2. Heat capacity and entropy

The low-temperature heat capacity of Yb<sub>2</sub>O<sub>3</sub> has been measured by Justice and Westrum Jr.<sup>113</sup> from 10 to 350 K. They obtained  $S^\circ(298.15 \text{ K}) - S^\circ(10 \text{ K}) = 121.42 \text{ J K}^{-1} \text{ mol}^{-1}$ . They extrapolated the results to  $T = 0 \text{ K}$ , and added  $2R \ln(2)$  for the contribution of the ground state doublet of the split  $^2F_{7/2}$  multiplet. Li *et al.*<sup>176</sup> confirmed this by calorimetric measurements in the temperature range 1.5–15 K. These measurements reveal that Yb<sub>2</sub>O<sub>3</sub> orders antiferromagnetically with Néel temperature of 2.3 K, with an entropy close to

TABLE 27. Temperature of melting of ytterbium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{\text{fus}}/\text{K}$	
	Reported	ITS-90
Foex <sup>19</sup>	2693	2708
Mordovin <i>et al.</i> <sup>20</sup>	2646 $\pm$ 30	2649 $\pm$ 30
Noguchi and Mizuno <sup>21</sup>	2628 $\pm$ 20	2631 $\pm$ 20
Treswjatskii <i>et al.</i> <sup>22</sup>	2673 $\pm$ 20	2672 $\pm$ 20
Coutures <i>et al.</i> <sup>23</sup>	2708 $\pm$ 10	2707 $\pm$ 10
Mizuno <i>et al.</i> <sup>24</sup>	2660 $\pm$ 20	2648 $\pm$ 20
Shevthenko and Lopato <sup>13</sup>	2723	2722
Selected value:		2707 $\pm$ 15

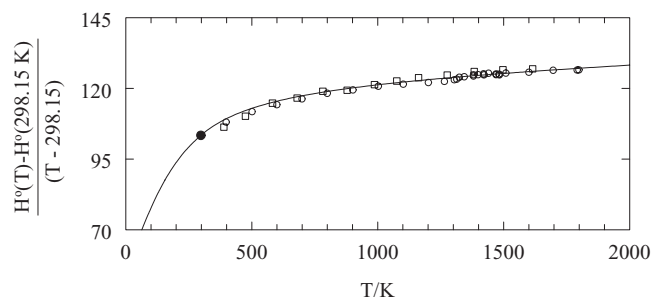


FIG. 13. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of Yb<sub>2</sub>O<sub>3</sub>;  $\circ$ , Tsagarishvili and Gvelesiani<sup>129</sup>;  $\square$ , Pankratz *et al.*<sup>90</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice *et al.*<sup>174</sup>; the curve shows the recommended equation.

$2R \ln(2)$ . Thus, we obtain for the entropy at  $T = 298.15 \text{ K}$ :

$$S^\circ(298.15 \text{ K}) = (133.1 \pm 0.3) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature enthalpy increment of Yb<sub>2</sub>O<sub>3</sub> has been measured by Pankratz *et al.*<sup>90</sup> from 400 to 1798 K and Tsagarishvili and Gvelesiani<sup>129</sup> from 384 to 1588 K. These data are in good agreement, and fit the low-temperature heat capacity data very well (Fig. 13). Our recommended heat-capacity equation is based on a polynomial fit of the results of both studies, which is constrained to  $C_p^\circ = 115.35 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements.<sup>113</sup> We thus obtain

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 130.6438 + 3.34628 \times 10^{-3}(T/\text{K}) - 1.44820 \times 10^6(T/\text{K})^{-2}.$$

The entropies of the C  $\rightarrow$  H transformation and melting are assumed to be the same as for Er<sub>2</sub>O<sub>3</sub>. This yields

$$\begin{aligned} \Delta_{\text{trs}} H^\circ(\text{C} \rightarrow \text{H}) &= (27 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{\text{fus}} H^\circ &= (84 \pm 8) \text{ kJ mol}^{-1}. \end{aligned}$$

For the heat capacity of the high temperature modification H and liquid Yb<sub>2</sub>O<sub>3</sub> we estimate

$$\begin{aligned} C_p^\circ(\text{Yb}_2\text{O}_3, \text{H}, T) &= 134 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{Yb}_2\text{O}_3, \text{liq}, T) &= 146 \text{ J K}^{-1} \text{ mol}^{-1}. \end{aligned}$$

### 3.21.3. Enthalpy of formation

The only experimental value has been reported by Huber Jr. *et al.*,<sup>163</sup> based on the results of oxygen-bomb combustion calorimetric measurements of 97.2 mass % pure Yb(cr) sample. This value, which includes a careful correction for impurities, was adopted by Cordfunke and Konings<sup>34</sup> with an increased uncertainty, and is also selected here

$$\Delta_f H^\circ(\text{Yb}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1814.5 \pm 6.0) \text{ kJ mol}^{-1}.$$

## 3.22. Lu<sub>2</sub>O<sub>3</sub>(cr,l)

### 3.22.1. Melting point

At room temperature, lutetium sesquioxide has the rare earth cubic C-type structure (space group  $Ia\bar{3}$ ), which is stable



TABLE 28. Temperature of melting of lutetium sesquioxide (after Coutures and Rand<sup>15</sup>)

Authors	$T_{fus}/K$	
	Reported	ITS-90
Mordovin <i>et al.</i> <sup>20</sup>	2741 ± 30	2744 ± 30
Noguchi and Mizuno <sup>21</sup>	2700 ± 10	2704 ± 10
Coutures <i>et al.</i> <sup>23</sup>	2763 ± 10	2762 ± 10
Shirvinskaya and Popova <sup>177</sup>	2643	2642
Mizuno <i>et al.</i> <sup>24</sup>	2733 ± 20	2722 ± 20
Shevthenko and Lopato <sup>13</sup>	2783	2782
Selected value:		2762 ± 15

up to the melting point. The measurements of the melting temperature of  $\text{Lu}_2\text{O}_3$  have been summarized in Table 28, which is based on the review by Coutures and Rand<sup>15</sup>; the results being corrected to ITS-90. The selected melting point is  $(2762 \pm 15)$  K.

### 3.22.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Lu}_2\text{O}_3$  has been measured by Justice *et al.*<sup>174</sup> from 6 to 350 K. The standard entropy derived from these data is

$$S^\circ(298.15 \text{ K}) = (109.96 \pm 0.13) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature heat capacity is derived from the enthalpy drop calorimetric measurements by Pankratz and Kelly<sup>65</sup> and Yashvili *et al.*,<sup>30</sup> which are in excellent agreement (Fig. 14). The experimental data have been fitted to a polynomial equation, with boundary condition  $C_p^\circ(298.15) = 101.75 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements.<sup>174</sup>

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 122.4593 + 7.29001 \times 10^{-3}(T/\text{K}) - 2.03414 \times 10^6(T/\text{K})^{-2}.$$

This equation is extrapolated to the melting point. Neither the heat capacity of the liquid phase nor the enthalpy of fusion are known. Our estimates for these quantities are approximate values obtained from the trend in the lanthanide sesquioxide series and comparison to the lanthanide trifluorides:

$$C_p^\circ(\text{Lu}_2\text{O}_3, \text{liq}, T) = 152 \text{ J K}^{-1} \text{ mol}^{-1}.$$

$$\Delta_{fus}H^\circ(\text{Lu}_2\text{O}_3) = (113 \pm 10) \text{ kJ mol}^{-1}.$$

### 3.22.3. Enthalpy of formation

The standard enthalpy of formation of cubic  $\text{Lu}_2\text{O}_3(\text{cr})$  is based on the oxygen-bomb combustion calorimetric study by Huber Jr. *et al.*<sup>178</sup> These authors used two well-analysed samples of lutetium metal, and obtained for the enthalpy of formation the values  $\Delta_f H^\circ(298.15 \text{ K}) = -(1891.8 \pm 14.2) \text{ kJ mol}^{-1}$  and  $\Delta_f H^\circ(298.15 \text{ K}) = -(1870.9 \pm 9.1) \text{ kJ mol}^{-1}$ ,

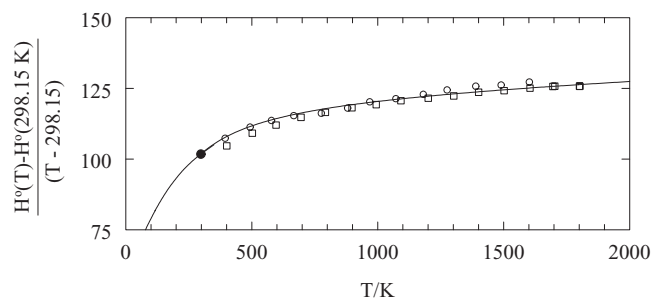


FIG. 14. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{Lu}_2\text{O}_3$ ;  $\circ$ , Yashvili *et al.*<sup>30</sup>;  $\square$ , Pankratz and Kelly<sup>65</sup>;  $\bullet$ , value derived from the low-temperature measurements by Justice *et al.*<sup>174</sup>; the curve shows the recommended equation.

after correction for impurities. As suggested by Cordfunke and Konings<sup>34</sup> the weighted mean of the two results is selected

$$\Delta_f H^\circ(\text{Lu}_2\text{O}_3, \text{cr}, 298.15 \text{ K}) = -(1877.0 \pm 7.7) \text{ kJ mol}^{-1}.$$

## 4. The Gaseous Lanthanide Oxides

### 4.1. LaO(g)

#### 4.1.1. Heat capacity and entropy

Thermal functions of  $\text{LaO}(\text{g})$  in the standard state have been calculated using the molecular constants given in Table 29.

The molecule of  $\text{LaO}$  is well investigated. Seven electronic states  $X^2\Sigma^+$ ,  $A^2\Delta$ ,  $A^2\Pi$ ,  $B^2\Sigma^+$ ,  $C^2\Pi$ ,  $D^2\Sigma^+$ , and  $F^2\Sigma^+$  were observed and analyzed in emission and absorption spectra. The comprehensive reviews of spectral data published before 1975 were given by Huber and Herzberg<sup>179</sup> and up to 1980 by Gurvich *et al.*<sup>180</sup> Later numerous works were done to improve the molecular constants of  $\text{LaO}$  by Bernard and Sibai,<sup>181</sup> Suarez,<sup>182</sup> Carette,<sup>183</sup> Bernard and Vergès,<sup>184</sup> and Steimle and Virgo<sup>185</sup> using traditional methods of investigation; by Bernard *et al.*<sup>186</sup> and Childs *et al.*<sup>187</sup> applying the laser induced fluorescence method; by Suenram *et al.*<sup>188</sup> and Törring *et al.*,<sup>189</sup> who studied the microwave rotational spectrum of  $\text{LaO}$ . The fundamental frequencies for  $\text{La}^{16}\text{O}$  and  $\text{La}^{18}\text{O}$  were observed in Ar-matrix by Andrews *et al.*<sup>190</sup> (796.7 and  $756.1 \text{ cm}^{-1}$  for, respectively), and by Zhang *et al.*<sup>191</sup> (796.7 and  $756.8 \text{ cm}^{-1}$ , respectively) Taking into account the matrix shift the fundamental frequencies obtained in matrices are in agreement with the gas phase data.

*Ab initio* results for the ground state of  $\text{LaO}$  by Hong *et al.*,<sup>192</sup> Dolg and Stoll,<sup>193</sup> Dolg *et al.*,<sup>194</sup> and Cao *et al.*<sup>195</sup> show remarkable agreement with the experimental data. *Ab initio* calculations for excited states predicted also the existence of the unobserved doublet states  $^2\Phi$ ,  $^2\Delta$ ,  $^2\Pi$ ,  $^2\Sigma$  by Marquez *et al.*<sup>196</sup> and Kotzian *et al.*<sup>197</sup> and quartet states  $1^4\Pi$ ,  $1^4\Phi$ ,  $1^4\Delta$  and  $2^4\Pi$  by Schampsand *et al.*<sup>198</sup> The Ligand field calculations by Kaledin *et al.*<sup>199</sup> and Schampsand *et al.*<sup>198</sup> resulted in the configuration assignment of the experimentally known states. The existence of the quartet state at the energy  $17200 \pm 800 \text{ cm}^{-1}$  was confirmed by Klingeler *et al.*<sup>200</sup> when studying the photoelectron spectra of  $\text{LaO}^-$ . All the theoretical

TABLE 29. Molecular constants of LaO(g)

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		cm <sup>-1</sup>						pm	
0 <sup>a</sup>	X <sup>2</sup> Σ <sup>+</sup>	0	817.026	2.1292 <sup>b</sup>	0.3525201	1.424 <sup>c</sup>	2.63	182.5	2
1 <sup>a</sup>	A' <sup>2</sup> Δ <sub>3/2</sub>	7493.4							2
2 <sup>a</sup>	A' <sup>2</sup> Δ <sub>5/2</sub>	8191.2							2
3 <sup>a</sup>	A <sup>2</sup> Π	13094.53							4
4 <sup>a</sup>	B <sup>2</sup> Σ	17878.73							2
5 <sup>a</sup>	C <sup>2</sup> Π	22740.38							4
6 <sup>a</sup>	D <sup>2</sup> Σ	26958.96							2
7 <sup>a</sup>	F <sup>2</sup> Σ	28 015							4
8 <sup>d</sup>		17 200							8
9 <sup>d</sup>		18 500							8
10 <sup>d</sup>		21 200							8
11 <sup>d</sup>		24 000							20
12 <sup>d</sup>		25 900							8
13 <sup>d</sup>		30 200							12
14 <sup>d</sup>		35 000							30
15 <sup>d</sup>		40 000							36

<sup>a</sup>Experimental state.<sup>b</sup> $\omega_e y_e = -3.15 \times 10^{-3} \text{ cm}^{-1}$ .<sup>c</sup> $\alpha_2 10^6 = -2.97 \text{ cm}^{-1}$ .<sup>d</sup>Estimated state.

data with some corrections were used in the present work to estimate unobserved electronic states (see Table 29).

The accepted constants for the <sup>2</sup>Σ<sup>+</sup> state (see Table 29) were obtained from the analysis of the high-resolution laser excited fluorescence B<sup>2</sup>Σ<sup>+</sup> - X<sup>2</sup>Σ<sup>+</sup> transition ( $v' \leq 4, v'' \leq 5$ ) by Bernard *et al.*<sup>186</sup> In their treatment the authors fixed rotational constants at the very precise values derived by Törring *et al.*<sup>189</sup> from the microwave spectrum for the levels  $v = 0-2$ . The selected vibrational constants quite well describe the extensive but less accurate data given by Schoonveld and Sundaram<sup>201</sup> for  $v'' \leq 11$ .

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (239.594 \pm 0.03) \text{ J K}^{-1} \text{ mol}^{-1}.$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 28.0550 + 21.9688 \times 10^{-3} (\text{T/K}) \\ &\quad - 19.25691 \times 10^{-6} (\text{T/K})^2 + 6.083418 \\ &\quad \times 10^{-9} (\text{T/K})^3 - 1.145313 \\ &\quad \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1200 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 41.7593 - 6.82858 \times 10^{-3} (\text{T/K}) \\ &\quad + 3.529957 \times 10^{-6} (\text{T/K})^2 \\ &\quad - 0.281197 \times 10^{-9} (\text{T/K})^3 \\ &\quad - 1.500459 \times 10^6 (\text{T/K})^{-2} \end{aligned}$$

for the 1200–4000 K range.

#### 4.1.2. Enthalpy of formation

Results of determination of LaO(g) enthalpy of formation are presented in Table 30. The experimental data considered in the calculations consist mainly of Knudsen effusion, mass

spectrometric, and combined measurements. In an early investigation of lanthanum sesquioxide vaporization behavior<sup>202</sup> using Knudsen effusion method, the principal mode of vaporization from tungsten cell was found to be



with the steady-state condition corresponding to slightly substoichiometric composition La<sub>2</sub>O<sub>2.96</sub>, and thermochemical assessment of the mass-loss data was made according to this equation. In a later detailed study by Ackermann and Rauh<sup>203</sup> it was found, however, that evaporation from tungsten results in LaO partial pressures higher than in the case of evaporation from rhenium cell, due to significant decrease of oxygen pressure for substoichiometric lanthanum sesquioxide. In the case of a rhenium cell, a substantially lower reducing effect of the cell material was found. From the data obtained by Ackermann and Rauh<sup>203</sup> it follows that Knudsen effusion data assessed under assumption of congruent lanthanum sesquioxide vaporization will lead to overestimated LnO stability. The only exception is the set of experimental data for lanthanum sesquioxide vaporization<sup>203</sup> from the rhenium effusion cell, corrected for slight deviation of the sample composition from exact stoichiometry.

The first mass spectrometric determination of the enthalpy of formation of LaO(g) has been carried out by Chupka *et al.*<sup>204</sup> who investigated the La<sub>2</sub>O<sub>3</sub>(cr) + La(l) system. Later, Ackermann and Rauh<sup>203</sup> investigated the same system in more detail using a combination of Knudsen effusion and mass spectrometry. It was shown that the vapor pressure of La in the system is slightly lower than over pure lanthanum but tends to that at decreasing temperatures. In general, the measurements performed in both investigations can be used in selecting of recommended value for the enthalpy of formation of LaO(g), with more weight to more precise and reliable data of Ackermann and Rauh.<sup>203</sup>

TABLE 30. The enthalpy of formation of LaO(g), in kJ mol<sup>-1</sup>

Authors	T/K	Method <sup>a</sup>	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Chupka <i>et al.</i> <sup>204</sup>	M	1775–1865	$\frac{1}{3}\text{La(g)} + \frac{1}{3}\text{La}_2\text{O}_3(\text{cr}) = \text{LaO(g)}$	$348.8 \pm 19$	$-105.1 \pm 20$
Goldstein <i>et al.</i> <sup>202</sup>	K	2234–2441	$\text{La}_2\text{O}_3(\text{cr}) = 2\text{LaO(g)} + \text{O(g)}$	$1774.2 \pm 8.3$	$-133.3 \pm 10$
Smoes <i>et al.</i> <sup>205</sup>	M	1810–2220	$\text{ScO(g)} + \text{La(g)} = \text{Sc(g)} + \text{LaO(g)}$	$-119.3 \pm 6.8$	$-122.7 \pm 8$
	M	1890–2270	$\text{YO(g)} + \text{La(g)} = \text{Y(g)} + \text{LaO(g)}$	$-81.0 \pm 7.2$	$-120.6 \pm 8$
Ames <i>et al.</i> <sup>206</sup>	M	1890–2220	$\text{Y(g)} + \text{LaO(g)} = \text{YO(g)} + \text{La(g)}$	$77.0 \pm 7.1$	$-116.6 \pm 8$
	M	1870–2080	$\text{Sc(g)} + \text{LaO(g)} = \text{ScO(g)} + \text{La(g)}$	$111.8 \pm 6.7$	$-115.2 \pm 8$
Coppens <i>et al.</i> <sup>207</sup>	M	1877–2088	$\text{SiO(g)} + \text{La(g)} = \text{Si(g)} + \text{LaO(g)}$	$-2.0 \pm 7.8$	$-120.8 \pm 9$
Coppens <i>et al.</i> <sup>208</sup>	M	2146–2270	$\text{B(g)} + \text{LaO(g)} = \text{BO(g)} + \text{La(g)}$	$-11.5 \pm 13$	$-113.7 \pm 15$
Ackermann and Rauh <sup>203</sup>	K (W cell)	2258–2427	$\text{La}_2\text{O}_3(\text{cr}) = 2\text{LaO(g)} + \text{O(g)}$	$1785.6 \pm 6.8$	$-127.6 \pm 8$
	M/K (Re cell)	1800–2300	$\text{La}_2\text{O}_3(\text{cr}) = 2\text{LaO(g)} + \text{O(g)}$	$1801.9 \pm 12$	$-119.4 \pm 8$
	M/K	1516–1904	$\text{La}_2\text{O}_3(\text{cr}) + \text{La(l)} = 3\text{LaO(g)}$	$1435.7 \pm 5.8$	$-118.6 \pm 4$
	M	1783–2184	$\text{La(g)} + \text{YO(g)} = \text{LaO(g)} + \text{Y(g)}$	$-82.4 \pm 6.7$	$-122.0 \pm 8$
Parr <sup>210</sup>	M/P		$D_0(\text{LaO}) = 793.7 \pm 3.0$		$-118.7 \pm 4$
Gole and Chalek <sup>209</sup>	C		$D_0(\text{LaO}) \geq 790.1 \pm 4.2$		$\leq -115.1 \pm 5$
Selected value:			$D_0(\text{LaO}) = 796.1 \pm 8$		$-119.0 \pm 8$

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; P = photoionization; C = beam-gas chemiluminescence.

Several mass spectrometric works deal with the LaO(g) + M(g) oxygen exchange reactions.<sup>203,205–208</sup> Only reactions without other lanthanides were taken into consideration. All MO molecules involved are characterized by reliable values of enthalpy of formation, taken from the reference book “*Thermodynamic Properties of Individual Substances*.”<sup>7,180</sup>

The selected value

$$\Delta_f H^\circ(\text{LaO, g, 298.15 K}) = -(119 \pm 8) \text{ kJ mol}^{-1}$$

is taken as rounded average of the third-law values calculated from all mentioned works except data of Goldstein *et al.*<sup>202</sup> and Ackermann and Rauh<sup>203</sup> on evaporation of lanthanum sesquioxide from tungsten cells. The data on chemiluminescence<sup>209</sup> and photoionization of LaO molecules<sup>210</sup> confirm the selected value. The selected enthalpy of formation corresponds to a dissociation energy of LaO molecule  $D_0(\text{LaO}) = (796 \pm 8) \text{ kJ mol}^{-1}$ .

## 4.2. CeO<sub>2</sub>(g)

### 4.2.1. Heat capacity and entropy

The infrared spectrum of the CeO<sub>2</sub> molecule isolated in inert gas matrices has been determined by DeKock and Weltner Jr.,<sup>211</sup> Gabelnick *et al.*<sup>212</sup> and Willson and Andrews.<sup>213</sup> The results indicate that the molecule has a bent structure (C<sub>2v</sub>), an approximate bond distance of 188 pm was derived.<sup>212</sup> This geometry was confirmed by quantum chemical calculations by Heinemaan *et al.*,<sup>214</sup> Hülsen *et al.*,<sup>215</sup> and Todorova *et al.*<sup>216</sup> We have selected the bond distance (182 pm) and bond angle (126°) from the CASPT2 computations by Todorova *et al.*<sup>216</sup> The experimental values for the stretching frequencies,  $\nu_1 = 757 \text{ cm}^{-1}$  and  $\nu_3 = 737 \text{ cm}^{-1}$  have been adopted here. The experimental bending frequency,  $\nu_2 = 262 \text{ cm}^{-1}$  that is selected for the calculations of the thermal functions, is significantly higher than the value derived from the CASPT2 calculations,  $\nu_2 = 102 \text{ cm}^{-1}$ .<sup>216</sup> DeKock and Weltner, Jr.<sup>211</sup> studied the electronic spectrum of CeO<sub>2</sub> and found no electronic levels below  $15\,000 \text{ cm}^{-1}$ . In our calculations we have included some estimated higher energy levels (Table 31). The

derived entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (274.417 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 37.7646 + 55.1209 \times 10^{-3}(\text{T/K}) \\ &\quad - 57.1816 \times 10^{-6}(\text{T/K})^2 + 21.2692 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 2.6077 \\ &\quad \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–900 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 55.9864 + 2.90431 \times 10^{-3}(\text{T/K}) \\ &\quad - 1.4361 \times 10^{-6}(\text{T/K})^2 + 0.25254 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 1.11908 \\ &\quad \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 900–4000 K range.

### 4.2.2. Enthalpy of formation

Vapor pressure measurements of CeO<sub>2</sub> have been reported by Shukarev and Semenov,<sup>217</sup> Benezech and Foex,<sup>218</sup> Piacente *et al.*,<sup>219</sup> and Ackermann and Rauh.<sup>220</sup> The results by Shchukarev and Semenov (transpiration method) and Benezech and Foex (mass spectrometry) are very different from those obtained by Ackermann and Rauh<sup>220</sup> and by

TABLE 31. The molecular parameters for CeO<sub>2</sub>(g)

Parameter	Value
Ground electronic state	<sup>1</sup> A <sub>1</sub>
Symmetry group	C <sub>2v</sub>
Symmetry number, $\sigma$	2
$I_A I_B I_C$ (g <sup>3</sup> cm <sup>6</sup> ) <sup>a</sup>	$6.984 \times 10^{-115}$
Vibrational frequencies (cm <sup>-1</sup> )	757.0, 264, 736.8
Electronic states (cm <sup>-1</sup> ) <sup>b</sup>	0(1), 18000(2), 19000(2), 22000(2), 24000(2), 25000(2)

<sup>a</sup>Product of moments of inertia.

<sup>b</sup>Numbers in parentheses represent the statistical weights.

TABLE 32. The enthalpy of formation of CeO<sub>2</sub>(g) at 298.15 K

Authors	Method <sup>a</sup>	T/K	reaction	$\Delta_f H^\circ(298.15 \text{ K})/\text{kJ mol}^{-1}$
Ackermann and Rauh <sup>220</sup>	ME	1846–2318	Ce <sub>2</sub> O <sub>3+x</sub> (cr) = (1-x)CeO(g) + (1+x) CeO <sub>2</sub> (g) <sup>b</sup>	-542.8 ± 8.2
Piacente <i>et al.</i> <sup>219</sup>	K/M	1736–2067	CeO <sub>2</sub> (cr) = CeO <sub>2</sub> (g)	-532.4 ± 2.0
	K/M	1933–2134	Ce(g) + CeO <sub>2</sub> (g) = 2CeO(g)	-573.8 ± 16.3
Staley and Norman <sup>222</sup>	K/M	1845–1975	CeO <sub>2</sub> (g) = CeO(g) + 0.5O <sub>2</sub> (g)	-415.7 ± 8.0 <sup>c</sup>
	K/M	1845–1975	CeO(g) + CaO(g) = CeO <sub>2</sub> (g) + Ca(g)	-511.5 ± 22.5 <sup>c</sup>
Younés <i>et al.</i> <sup>221</sup>	K/M	1720–2040	Ce(g) + CeO <sub>2</sub> (g) = 2CeO(g)	-599.3 ± 9.6 <sup>c</sup>
	K/M	1720–2040	La(g) + CeO <sub>2</sub> (g) = LaO(g) + CeO(g)	-593.8 ± 12.3 <sup>c</sup>
	K/M	1720–2040	CeO(g) + UO <sub>3</sub> (g) = CeO <sub>2</sub> (g) + UO <sub>2</sub> (g)	-489.4 ± 17.9 <sup>c</sup>
	K/M	1720–2040	CeO <sub>2</sub> (g) + UO(g) = CeO(g) + UO <sub>2</sub> (g)	-565.7 ± 17.7 <sup>c</sup>
Selected value:				-538 ± 20

<sup>a</sup>ME = mass effusion; K/M = Knudsen-cell mass spectrometry.

<sup>b</sup>Partial vapor pressures of CeO(g) and CeO<sub>2</sub>(g) measured over the congruently vaporizing composition (cvc).

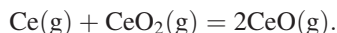
<sup>c</sup>Derived from second-law values given by the authors.

Piacente *et al.*<sup>219</sup> Moreover, the results have been presented in graphical form only.

Ackermann and Rauh<sup>220</sup> measured the vapor pressure of the congruently vaporizing composition and of Ce<sub>2</sub>O<sub>3.03</sub>(cr), corresponding to the reaction:



The enthalpy of formation derived from this study are summarized in Table 32. Piacente *et al.*<sup>219</sup> studied the equilibrium between solid and gaseous CeO<sub>2</sub> by means of Knudsen effusion mass spectrometry, and have been interpreted as congruent vaporisation. In addition, these authors have studied the isomolecular exchange reaction:



This reaction has also been studied by Younés *et al.*,<sup>221</sup> who reported, however, only second-law enthalpies of reaction. Younés *et al.*<sup>221</sup> also reported results for isomolecular reactions with La, UO, and UO<sub>2</sub>, again giving only the second-law enthalpies of reaction (see Table 32). Staley and Norman<sup>222</sup> studied the isomolecular exchange reaction involving CeO(g) and CeO<sub>2</sub>(g), as well as the isomolecular exchange reaction with CaO(g), but also in this work only second-law enthalpies of reaction have been given (Table 32).

The variation in the values obtained is large, particularly among those derived from the second-law analysis, and the selected value for  $\Delta_f H^\circ$  of CeO<sub>2</sub>(g) is the average of the (third-law) values derived from the work of Ackermann and Rauh<sup>220</sup> and Piacente *et al.*,<sup>219</sup> which we considered the most reliable

$$\Delta_f H^\circ(298.15 \text{ K}) = -(538 \pm 20) \text{ kJ mol}^{-1}.$$

### 4.3. CeO(g)

#### 4.3.1. Heat capacity and entropy

The thermal functions of CeO(g) in the standard state have been calculated using the molecular constants presented in Table 33.

The analyses of 34 electronic transitions of CeO was performed by Barrow *et al.*,<sup>223</sup> Linton *et al.*,<sup>224</sup> Linton *et al.*,<sup>225</sup> Linton and Dulick,<sup>226</sup> Linton *et al.*,<sup>227</sup> Kaledin *et al.*<sup>199</sup> and Todorova *et al.*<sup>216</sup> As a result, all 16 lower electronic states belonging to the lowest 4f6s electron configuration of CeO were identified (see Table 33). The configuration assignment for the lower states of these transitions was shown to be doubtless.<sup>197,199,228–232</sup> On the other hand, the configuration assignments for the upper states of these transitions were shown to be contradictory.<sup>197,199,232–234</sup> Certainly these upper states should belong to the 4f6p, 4f5d, and 4f<sup>2</sup> superconfigurations or to the mixed configurations. The estimated statistical weights are presented at fixed energies in Table 33, which are calculated assuming the energy intervals for 4f6p, 4f5d, and 4f<sup>2</sup> states to be 7000–36 000 cm<sup>-1</sup>, 12 000–25 000 cm<sup>-1</sup>, and 17 000–45 000 cm<sup>-1</sup>, respectively. The 6s<sup>2</sup>, 5d6s, 6s6p, 6p<sup>2</sup>, 5d<sup>2</sup>, and 4f6s states are taken into account in the interval 30 000–45 000 cm<sup>-1</sup>.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (246.099 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 22.01944 + 55.5050 \times 10^{-3}(\text{T/K}) \\ &\quad - 43.14095 \times 10^{-6}(\text{T/K})^2 + 10.89494 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 4.23189 \\ &\quad \times 10^4(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1300 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 62.79967 - 17.53953 \times 10^{-3}(\text{T/K}) \\ &\quad + 5.431417 \times 10^{-6}(\text{T/K})^2 - 0.487485 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 4.947446 \\ &\quad \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1300–4000 K range.

#### 4.3.2. Enthalpy of formation

The results of the determination of the enthalpy of formation of CeO(g) are presented in Table 34. Several mass spectrometric measurements of isomolecular oxygen exchange



TABLE 33. Molecular constants of CeO(g)

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$ pm	$p_i$
		cm <sup>-1</sup>							
0 <sup>a</sup>	X(1)2	0	829.5	2.6 <sup>b</sup>	0.3553	1.6 <sup>c</sup>	2.46 <sup>d</sup>	181.198	2
1 <sup>a</sup>	(1)3	82							2
2 <sup>a</sup>	(1)1	812.7							2
3 <sup>a</sup>	(2)2	911.8							2
4 <sup>a</sup>	(1)0	1678.6							1
5 <sup>a</sup>	(2)1	1875.3							2
6 <sup>a</sup>	(2)0	1925.3							1
7 <sup>a</sup>	(1)4	2042.6							2
8 <sup>a</sup>	(2)3	2142.6							2
9 <sup>a</sup>	(2)4	2618.4							2
10 <sup>a</sup>	(3)3	2771.3							2
11 <sup>a</sup>	(3)4	3462.2							2
12 <sup>a</sup>	(3)1	3435.0							2
13 <sup>a</sup>	(3)0	3818.8							1
14 <sup>a</sup>	(4)1	4134.1							2
15 <sup>a</sup>	(4)0	4458.0							1
16 <sup>c</sup>		7000							7
17 <sup>c</sup>		10 000							22
18 <sup>c</sup>		15 000							58
19 <sup>c</sup>		20 000							72
20 <sup>c</sup>		25 000							56
21 <sup>c</sup>		30 000							45
22 <sup>c</sup>		35 000							92
23 <sup>c</sup>		40 000							112

<sup>a</sup>Experimental (4f6s) state.<sup>b</sup>Calculated using  $\Delta G_{1/2} = 824.3 \text{ cm}^{-1}$  from Linton *et al.*<sup>224</sup> and the dissociation energy adopted in this work.<sup>c</sup>Calculated using  $B_0 = 0.35454 \text{ cm}^{-1}$  from Barrow *et al.*<sup>223</sup> and the Pekeris relation.<sup>d</sup> $D_0$ .<sup>e</sup>Estimated state.

reactions have been carried out.<sup>207,220,221,235</sup> Results of all measurements are in remarkable agreement. The selected value for the enthalpy of formation of CeO(g)

$$\Delta_f H^\circ(\text{CeO}, \text{g}, 298.15 \text{ K}) = -(132 \pm 8) \text{ kJ mol}^{-1}$$

is taken as a rounded average of the third-law values calculated from all mentioned works. To the selected enthalpy of formation corresponds the value of dissociation energy of CeO molecule  $D_0(\text{CeO}) = (794.3 \pm 8) \text{ kJ mol}^{-1}$ .

#### 4.4. PrO(g)

##### 4.4.1. Heat capacity and entropy

The thermal functions of PrO(g) in the standard state have been calculated using the molecular constants presented in Table 35.

The analysis of 34 electronic transitions of PrO in emission, absorption, and laser fluorescence spectra was carried out by Shenyavskaya *et al.*,<sup>236</sup> Delaval *et al.*,<sup>237</sup> Beaufils *et al.*,<sup>238</sup> Dulick *et al.*,<sup>239</sup> Dulick *et al.*,<sup>240</sup> Dulick *et al.*,<sup>241</sup> Shenyavskaya and Kaledin,<sup>242</sup> Dulick and Field,<sup>243</sup> and Childs *et al.*<sup>244</sup> As a result, information about 12 low-lying electronic states belonging to the lowest electron configuration  $4f^2 6s$  was obtained (see Table 35). The infrared spectrum of PrO was observed in solid Ar and Kr matrices at 4 K by Weltner and De Kock<sup>245</sup> and Willson *et al.*<sup>246</sup> The values of the fundamental frequency of PrO obtained in matrices are in agreement with the electronic spectra results. The  $4f^2 6s$  electron configuration of PrO contains 91 Hund case “c” molecular states, and each of them is doubly degenerate. Dulick<sup>234</sup> calculated all these states using the crystal field theory and adjusting parameters to reproduce experimental data. The low-lying states of PrO belonging to the  $4f^2 6s$  configuration were calculated also by

TABLE 34. The enthalpy of formation of CeO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Walsh <i>et al.</i> <sup>235b</sup>	M	1700–2040	Ce(g) + LaO(g) = CeO(g) + La(g)	2.26	-131.6
Coppens <i>et al.</i> <sup>207</sup>	M	2146–2270	CeO(g) + La(g) = Ce(g) + LaO(g)		
Ackermann and Rauh <sup>203</sup>	M	1580–1920	Ce(g) + LaO(g) = CeO(g) + La(g)	3.34	-130.5
	M	1760–2160	Ce(g) + YO(g) = CeO(g) + Y(g)	-81.05	-133.5
Younés <i>et al.</i> <sup>221</sup>	M	1490–2030	Ce(g) + LaO(g) = CeO(g) + La(g)	2.05	-131.8
Selected value:			$D_0(\text{CeO}) = 794.8 \pm 8$		-132.0 ± 8

<sup>a</sup>M = mass spectrometry.<sup>b</sup>Recalculated by Ames *et al.*<sup>206</sup>

TABLE 35. Molecular constants of  $^{141}\text{Pr}^{16}\text{O}(\text{g})$ 

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$ pm	$p_i$
		$\text{cm}^{-1}$							
0 <sup>a</sup>	X(1)3.5	0	835.8 <sup>b</sup>	2.22 <sup>b</sup>	0.36175 <sup>b</sup>	1.5 <sup>b</sup>	2.6 <sup>b</sup>	180.115	2
1 <sup>a</sup>	(1)4.5	217							2
2 <sup>a</sup>	(2)3.5	2064							2
3 <sup>a</sup>	(1)5.5	2099							2
4 <sup>a</sup>	(2)4.5	2154							2
5 <sup>a</sup>	(3)4.5	3718							2
6 <sup>a</sup>	(3)3.5	3887							2
7 <sup>a</sup>	(1)6.5	3953							2
8 <sup>a</sup>	(2)5.5	4225							2
9 <sup>a</sup>	(3)5.5	5541							2
10 <sup>a</sup>	(4)5.5	5938							2
11 <sup>c</sup>		1870							2
12 <sup>c</sup>		2880							2
13 <sup>c</sup>		3170							8
14 <sup>c</sup>		4950							10
15 <sup>c</sup>		5247							4
16 <sup>c</sup>		7000							5
17 <sup>c</sup>		10 000							90
18 <sup>c</sup>		15 000							150
19 <sup>c</sup>		20 000							175
20 <sup>c</sup>		25 000							185
21 <sup>c</sup>		30 000							195
22 <sup>c</sup>		35 000							200
23 <sup>c</sup>		40 000							205

<sup>a</sup>Experimental ( $4f^26s$ ) state.<sup>b</sup>From Shenyavskaya *et al.*<sup>236</sup><sup>c</sup>Estimated state.

Dulick *et al.*<sup>247</sup> and Carrete and Hocquet<sup>230</sup> in Ligand field approximation and by Kotzian and Roesch<sup>248</sup> applying the theoretical Intermediate Neglect of Differential Overlap (INDO) MO procedure augmented with a double-group CI technique which includes spin-orbit interactions. Kotzian and Roesch<sup>232</sup> predicted also the lowest states below 40 000  $\text{cm}^{-1}$  belonging to the  $4f^25d$  and  $4f^3$  configurations. The upper states of the transitions observed in the visible apparently are the  $p-s$  transitions, so the  $4f^26p$  states were taken into account in the interval 130 00–63 000  $\text{cm}^{-1}$ .

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (244.367 \pm 0.05) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 22.98903 + 20.3103 \times 10^{-3} (\text{T/K}) \\ &+ 20.2961 \times 10^{-6} (\text{T/K})^2 - 14.6500 \\ &\times 10^{-9} (\text{T/K})^3 + 2.72239 \\ &\times 10^5 (\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1200 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 84.67666 - 28.5857 \times 10^{-3} (\text{T/K}) \\ &+ 8.73036 \times 10^{-6} (\text{T/K})^2 - 0.84566 \\ &\times 10^{-9} (\text{T/K})^3 - 1.44324 \\ &\times 10^7 (\text{T/K})^{-2} \end{aligned}$$

for the 1200–4000 K range.

#### 4.4.2. Enthalpy of formation

The results for the enthalpy of formation of  $\text{PrO}(\text{g})$  are presented in Table 36. Experimental data used in calculations consist of Knudsen effusion and mass spectrometric measurements. Several mass spectrometric measurements of isomolecular oxygen exchange reactions have been carried out.<sup>235,249,250</sup> The selected value

$$\Delta_f H^\circ(\text{PrO}, \text{g}, 298.15 \text{ K}) = -(145.5 \pm 8) \text{ kJ mol}^{-1}$$

is taken as a rounded average of the third-law values calculated from all mentioned works. To the selected enthalpy of formation corresponds the value of dissociation energy of  $\text{PrO}$  molecule  $D_0(\text{PrO}) = (747.1 \pm 8) \text{ kJ mol}^{-1}$ .

#### 4.5. NdO(g)

##### 4.5.1. Heat capacity and entropy

The thermal functions of  $\text{NdO}(\text{g})$  in the standard state have been calculated using the molecular constants presented in Table 37.

The analysis of  $\text{NdO}$  spectra in emission, absorption, and laser fluorescence was carried out by Linton *et al.*,<sup>251</sup> Shenyavskaya *et al.*,<sup>252</sup> Kulikov,<sup>253</sup> Kaledin and Shenyavskaya,<sup>254</sup> and Kaledin.<sup>255</sup> As a result, nine low-lying electronic states with close values of molecular constants were revealed including the ground X4 state, and numerous more or less perturbed excited states with energies 10 000–23 000  $\text{cm}^{-1}$ . The IR spectrum of

TABLE 36. The enthalpy of formation of PrO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Walsh <i>et al.</i> <sup>235b</sup>	M	1750–2070	Pr(g) + LaO(g) = PrO(g) + La(g)	46.45	-148.7
Fries and Cater <sup>249</sup>	M	2053–2191	PrO(g) + Y(g) = Pr(g) + YO(g)	31.4	-145.1
Murad <sup>250</sup>	M		Pr(g) + TiO(g) = PrO(g) + Ti(g)	-71.4	-142.6
Selected value:			$D_0(\text{PrO}) = 747.1 \pm 8$		-145.5 $\pm$ 8

<sup>a</sup>M = mass spectrometry.<sup>b</sup>Recalculated by Ames *et al.*<sup>206</sup>

NdO was observed in solid Ar and Kr matrices at 4 K by Weltner and De Kock<sup>245</sup> and Willson *et al.*<sup>246</sup> The values of the fundamental frequency of NdO obtained in electronic and matrix IR spectra are in agreement.

The theoretical calculations by Dolg and Stoll<sup>193</sup> dealt with the molecular constants of lanthanide monoxides in their ground states. Boudreaux and Baxter<sup>256</sup> calculated and compared electronic structures of NdO and UO.

The low-lying electronic states of NdO were assigned to the  $4f^3 6s$  electron configuration which gives rise to the electronic states with the total statistical weight 728. The states belonging to the subconfiguration  $4f^3(4I)6s$  were calculated using the crystal field theory by Kulikov<sup>253</sup> and Carrete and Hocquet<sup>230</sup> (56 states with  $\Sigma p = 104$ ), and by Dulick *et al.*<sup>247</sup> (up to 1000 cm<sup>-1</sup>,  $\Sigma p = 98$ ). The best agreement with experimental data was shown by Kulikov<sup>253</sup> who used adjusted parameters to reproduce experimental data known at that time (4 low-lying states). Krauss and Stevens<sup>257</sup> confirmed above mentioned

crystal field results by *ab initio* calculations and predicted also the low-lying  $4f^3 5d$  and  $4f^2 s^2$  states (the lower limits are 3390 cm<sup>-1</sup> and 4267 cm<sup>-1</sup>, respectively). Shenyavskaya *et al.*<sup>252</sup> tentatively assigned the [12.171]4 state to the lowest state of the  $4f^2 s^2$  configuration. The strong bands observed in visible spectral region apparently are the  $p-s$  or  $d-s$  transitions. This implies that the  $4f^3 5d$  and  $4f^3 6p$  states should be taken into account when calculating the thermal functions. In Table 37 estimates are presented of the unobserved states belonging to the  $4f^3 6s$ ,  $4f^3 5d$ ,  $4f^3 5p$ , and  $4f^3 6s^2$  configurations. The states of the  $4f^3(4I)6s$  subconfiguration are taken from the work by Kulikov.<sup>253</sup> The lower limits for the  $4f^3(4I) 5d$  and  $4f^3(4I)5p$  states are assumed to be equal to 6000 and 13 500 cm<sup>-1</sup>, respectively. The lower limits for the states belonging to the subconfigurations  $4f^3(4F)nl$  are estimated based on the spectrum NdIV: lower state  $4I(4f^3)$  is separated from the others by  $\sim 12\,000$  cm<sup>-1</sup>, and addition of one electron does not change considerably the interval between centers of

TABLE 37. Molecular constants of <sup>142</sup>Nd<sup>16</sup>O(g)

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$ pm	$p_i$
		cm <sup>-1</sup>							
0 <sup>a</sup>	X(1)4	0	834.083 <sup>b</sup>	2.2855 <sup>b</sup>	0.362346 <sup>b</sup>	1.4291 <sup>b</sup>	2.7296 <sup>b</sup>	179.91	2
1 <sup>a</sup>	(1)5	474							2
2 <sup>a</sup>	(1)3	1156							2
3 <sup>a</sup>	(1)2	1351							2
4 <sup>a</sup>	(2)4	1587							2
5 <sup>a</sup>	(2)3	1793							2
6 <sup>a</sup>	(2)2	1914							2
7 <sup>a</sup>	(1)6	2124							2
8 <sup>a</sup>	(2)5	2153							2
9 <sup>c</sup>		1470							3
10 <sup>c</sup>		1914							3
11 <sup>c</sup>		3180							22
12 <sup>c</sup>		3987							2
13 <sup>c</sup>		5600							28
14 <sup>c</sup>		7200							17
15 <sup>c</sup>		7920							20
16 <sup>c</sup>		10 000							30
17 <sup>c</sup>		15 000							80
18 <sup>c</sup>		20 000							250
19 <sup>c</sup>		25 000							320
20 <sup>c</sup>		30 000							400
21 <sup>c</sup>		35 000							450
22 <sup>c</sup>		40 000							520

<sup>a</sup>Experimental ( $4f^3 6s$ ) state.<sup>b</sup>From Shenyavskaya *et al.*<sup>252</sup><sup>c</sup>Estimated state.

gravity of  $4f^3(4F)nl$  and  $4f^3(4I)nl$  configurations, and these limits are assumed to be lower for all remaining states of corresponding configurations.

Our estimates of the positions of the excited configurations are based on the experimental data by Shenyavskaya *et al.*<sup>252</sup> for  $f^2s^2$  configuration and on comparison with the positions of the analogous states of CeO and PrO (for  $f^3d$ ,  $f^3p$ , and  $f^4$  configurations). The energy ranges for the states included in the configurations were taken equal to those for UO (see Kaledin *et al.*<sup>258</sup>)

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (242.817 \pm 0.05) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & -9.80368 + 188.146 \times 10^{-3}(\text{T/K}) \\ & - 190.285 \times 10^{-6}(\text{T/K})^2 + 62.3079 \\ & \times 10^{-9}(\text{T/K})^3 + 5.70945 \\ & \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1100 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 57.65812 - 9.78465 \times 10^{-3}(\text{T/K}) \\ & + 2.76479 \times 10^{-6}(\text{T/K})^2 - 0.218778 \\ & \times 10^{-9}(\text{T/K})^3 + 4.4284910^5(\text{T/K})^{-2} \end{aligned}$$

for the 1100–4000 K range.

#### 4.5.2. Enthalpy of formation

The results of the determination of the enthalpy of formation of NdO(g) are presented in Table 38. Experimental data have been obtained by Knudsen effusion, transpiration, and mass-spectrometric measurements. Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurement of the weight loss for Nd<sub>2</sub>O<sub>3</sub>(cr). Results of these measurements were treated in this work under assumption of congruent vaporization of Nd<sub>2</sub>O<sub>3</sub>(cr) according to reaction



neglecting the possibility of formation of Nd(g) atoms in the vapor [see text on LaO(g)]. The enthalpy of formation of NdO(g) thus calculated can be more negative than the correct value, the degree of deviation being dependent on the amount of Nd atoms in vapors. Comparison with the results of Tetenbaum<sup>259</sup> and Murad and Hildenbrand<sup>260</sup> confirms this conclusion. Results of transpiration measurements<sup>259</sup> have

significant advantage in comparison to the Knudsen effusion measurements, due to well-defined oxygen pressure, making interpretation of results more certain. The measurements of Murad<sup>250</sup> for two oxygen exchange reactions result in practically coinciding values of the enthalpy of formation of NdO(g). The selected value

$$\Delta_f H^\circ(\text{NdO}, \text{g}, 298.15\text{K}) = -(120 \pm 8) \text{ kJ mol}^{-1}$$

is taken as the rounded average of the third-law values calculated from the works of Tetenbaum<sup>259</sup> and Murad.<sup>250</sup> The selected enthalpy of formation corresponds to  $D_0(\text{NdO}) = (692.8 \pm 8) \text{ kJ mol}^{-1}$ .

## 4.6. PmO(g)

### 4.6.1. Heat capacity and entropy

The thermal functions of PmO(g) in the standard state have been calculated using the molecular constants given in Table 39.

Experimental data on the PmO molecular spectra are absent. The constants presented in Table 39 are estimated. Dolg and Stoll<sup>193</sup> showed that the molecular constants of the lanthanide monoxides in the electronic states of the same electron configuration change regularly. Thus the constants can be estimated by interpolation. Field<sup>233</sup> predicted the ground state X3.5 belonging to the  $4f^4(5I)6s$  subconfiguration. Dulick *et al.*<sup>247</sup> carried out a Ligand field calculation of this subconfiguration and obtained for the ground state  $\Omega = 0.5$ . For the calculation of the thermal functions this difference in  $\Omega$  is of little importance because all the states of PmO belonging to this subconfiguration have statistical weight 2.

In the present work, the molecular constants of PmO are adopted (see Table 39) from the results of calculations by Dulick and Field<sup>243</sup> up to  $10\,000 \text{ cm}^{-1}$  and on the basis of rough estimates of statistical weights for the  $4f^46s$ ,  $4f^46p$ , and  $4f^45d$  states at  $T \geq 10\,000 \text{ cm}^{-1}$ .

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (246.519 \pm 2.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 31.64170 + 38.6653 \times 10^{-3}(\text{T/K}) \\ & - 30.3021 \times 10^{-6}(\text{T/K})^2 + 8.45852 \\ & \times 10^{-9}(\text{T/K})^3 - 2.08747 \\ & \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1100 K range, and

TABLE 38. The enthalpy of formation of NdO(g), in  $\text{kJ mol}^{-1}$

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Goldstein <i>et al.</i> <sup>202</sup>	K	2255–2408	$\text{Nd}_2\text{O}_3 = 2\text{NdO}(\text{g}) + \text{O}(\text{g})$	1790.3	–133.6
Tetenbaum <sup>259</sup>	T	2155–2485	$\text{Nd}_2\text{O}_3 = 2\text{NdO}(\text{g}) + \text{O}(\text{g})$	1576.6	–115.8
Murad <sup>250</sup>	M	1933	$\text{Nd}(\text{g}) + \text{ScO}(\text{g}) = \text{PrO}(\text{g}) + \text{Sc}(\text{g})$	–16.4	–122.1
	M	1999	$\text{Nd}(\text{g}) + \text{TiO}(\text{g}) = \text{PrO}(\text{g}) + \text{Ti}(\text{g})$	–26.5	–122.9
Selected value:			$D_0(\text{NdO}) = 692.8 \pm 8$		–120.0 ± 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; T = transpiration.



TABLE 39. Molecular constants of  $^{145}\text{Pm}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$						$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$		
0 <sup>a</sup>	X0.5	0	832	2.9 <sup>b</sup>	0.358 <sup>c</sup>	1.7 <sup>c</sup>	2.65 <sup>d</sup>	180.7	2
1 <sup>a</sup>		225							2
2 <sup>a</sup>		660							2
3 <sup>a</sup>		820							2
4 <sup>a</sup>		1240							6
5 <sup>a</sup>		2280							14
6 <sup>a</sup>		2890							4
7 <sup>a</sup>		3565							12
8 <sup>a</sup>		4340							12
9 <sup>a</sup>		5065							10
10 <sup>a</sup>		5470							6
11 <sup>a</sup>		6155							12
12 <sup>a</sup>		6950							20
13 <sup>a</sup>		8130							16
14 <sup>a</sup>		9070							6
15 <sup>a</sup>		10 000							35
16 <sup>a</sup>		15 000							60
17 <sup>a</sup>		20 000							180
18 <sup>a</sup>		25 000							400
19 <sup>a</sup>		30 000							800
20 <sup>a</sup>		35 000							1000
21 <sup>a</sup>		40 000							1300

<sup>a</sup>Experimental ( $4f^46s$ ) state.<sup>b</sup>Estimated state.<sup>c</sup>From Shenyavskaya *et al.*<sup>252</sup><sup>d</sup>.....

$$C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = 50.95780 + 1.31663 \times 10^{-3} (\text{T/K}) - 1.57622 \times 10^{-6} (\text{T/K})^2 + 0.282026 \times 10^{-9} (\text{T/K})^3 - 2.75930 \times 10^6 (\text{T/K})^{-2}$$

for the 1100–4000 K range.

#### 4.6.2. Enthalpy of formation

No experimental data are available for the enthalpy of formation of promethium monoxide  $\text{PmO}(\text{g})$ . The quantum-mechanical calculations for the lanthanide monoxides performed till now do not have a thermochemical quality necessary for inclusion in the thermodynamic evaluations. Instead, a model based on the conception of promotion of a free atom into a “valence state” was used for evaluation of the  $\text{PmO}$  molecule stability. For the lanthanide monoxide molecules, the idea was for the first time expressed by Ames *et al.*<sup>206</sup> It was shown that variations in the dissociation energies of the lanthanide monoxide series correspond to the magnitude of the  $4f^n \rightarrow 4f^{n-1}5d$  transitions of the  $\text{Ln}^{2+}$  ions. Murad and Hildenbrand,<sup>260</sup> using more reliable and more broad experimental basis, came to analogous conclusion that the  $4f^n \rightarrow 4f^{n-1}5d$  excitation energy plays a major role in determining the energetics of most of the lanthanide monoxide bonds.

The model was further developed by Gibson,<sup>261</sup> who came to conclusion that the  $\text{Ln}-\text{O}$  and  $\text{Ln}-\text{O}^+$  bonds “are evidently formed using two  $5d$  electrons, rather than one  $5d$  and one  $6s$  electron”. Using estimated excitation energies for the formation of atomic  $\text{Pm}$  in the  $5d6s$  and  $5d^26s$  configurations, 115

and 129  $\text{kJ mol}^{-1}$ , Gibson obtained two values for the dissociation energy of  $\text{PmO}(\text{g})$ : 712  $\text{kJ mol}^{-1}$  and 698  $\text{kJ mol}^{-1}$ . Taking into account the monotonous decrease of  $D_0(\text{LnO})$  values going from  $\text{LaO}$  to  $\text{EuO}$ , those values seem to be too high. A lower value  $D_0(\text{PmO}) \sim 630 \text{ kJ mol}^{-1}$  can be obtained as a mean value between  $D_0(\text{NdO})$  and  $D_0(\text{SmO})$ . In the present assessment a compromise between the two ways of estimation is selected:  $D_0(\text{PmO}) = (650 \pm 50) \text{ kJ mol}^{-1}$ . This value corresponds to

$$\Delta_f H^\circ(\text{PmO, g, 298.15 K}) = -(145 \pm 50) \text{ kJ mol}^{-1}.$$

### 4.7. SmO(g)

#### 4.7.1. Heat capacity and entropy

The thermal functions of  $\text{SmO}(\text{g})$  in the standard state have been calculated using the molecular constants given in Table 40.

Analysis of electronic transitions of  $\text{SmO}$  by Linton *et al.*,<sup>262</sup> Bujin and Linton,<sup>263</sup> Bujin and Linton,<sup>264</sup> Linton *et al.*,<sup>265</sup> Linton *et al.*,<sup>266</sup> and Hannigan<sup>267</sup> revealed 15 low-lying states including the ground  $X0^-$  state. The vibrational structure of the observed transitions was not developed. The  $\Delta G_{1/2}$  values were detected by Linton *et al.*<sup>265</sup> only for four low-lying states, but not for the ground state. The infrared spectrum of  $\text{SmO}$  was observed in solid Ar and Kr matrices by Weltner and De Kock,<sup>245</sup> Willson and Andrews,<sup>213</sup> and Willson *et al.*<sup>246</sup> Taking into account the matrix shift, the obtained values of the fundamental frequency are in agreement with the  $\Delta G_{1/2}$  value for the first excited state of  $\text{SmO}$  in gas phase. That permits us to estimate the vibrational constant  $\omega_e$  with an accuracy of  $2 \text{ cm}^{-1}$ . *Ab initio* calculations by Dolg and Stoll<sup>193</sup> dealt with the ground state of the molecule and resulted in theoretical spectroscopic constants.

The electron configuration assignments for the lower states of the transitions to the ground state configuration  $4f^56s$  are doubtless. The  $4f^5(6H)6s$  states were calculated using Ligand field theory by Carrete and Hocquet<sup>230</sup> and Dulick *et al.*<sup>247</sup> The configuration assignments for the upper states of the transitions were not considered. Table 40 gives the experimental  $4f^5(6H)6s$  states according to Bujin and Linton<sup>263</sup> and the estimated statistical weights of the  $4f^56s$ ,  $4f^56p$ , and  $4f^55d$  superconfigurations at fixed energies up to  $40\,000 \text{ cm}^{-1}$ .

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (246.592 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}.$$

and the coefficients of the equations for the heat capacity are

$$C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = 28.65158 + 45.8171 \times 10^{-3} (\text{T/K}) - 34.4081 \times 10^{-6} (\text{T/K})^2 + 7.99213 \times 10^{-9} (\text{T/K})^3 - 4.99673 \times 10^4 (\text{T/K})^{-2}$$

for the 298.15–1400 K range, and

TABLE 40. Molecular constants of  $^{152}\text{Sm}^{16}\text{O}(\text{g})$ 

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		$\text{cm}^{-1}$					pm		
0 <sup>a</sup>	X(1)0 <sup>-</sup>	0	829.5 <sup>b</sup>	3.58	0.353952 <sup>c</sup>	2.0	2.58 <sup>d</sup>	181.415	1
1 <sup>a</sup>	(1)1	147							2
2 <sup>a</sup>	(1)2	567							2
3 <sup>a</sup>	(2)0 <sup>+</sup>	582							1
4 <sup>a</sup>	(2)1	879							2
5 <sup>a</sup>	(1)3	1280							2
6 <sup>a</sup>	(3)0 <sup>+</sup>	1546							1
7 <sup>a</sup>	(2)2	1604							2
8 <sup>a</sup>	(4)0 <sup>+</sup>	1661							1
9 <sup>a</sup>	(3)1	1661							2
10 <sup>a</sup>	(4)1	2014							2
11 <sup>a</sup>	(3)2	2240							2
12 <sup>a</sup>	(1)4	2287							2
13 <sup>a</sup>	(5)0 <sup>+</sup>	2520							2
14 <sup>a</sup>	(5)1	2867							2
15 <sup>c</sup>		2710							2
16 <sup>c</sup>		3040							4
17 <sup>c</sup>		3450							4
18 <sup>c</sup>		3860							12
19 <sup>c</sup>		4490							8
20 <sup>c</sup>		5000							4
21 <sup>c</sup>		10 000							25
22 <sup>c</sup>		15 000							60
23 <sup>c</sup>		20 000							290
24 <sup>c</sup>		25 000							380
25 <sup>c</sup>		30 000							400
26 <sup>c</sup>		35 000							1200
27 <sup>c</sup>		40 000							2350

<sup>a</sup>Experimental ( $4f^6 6s$ ) state.

<sup>b</sup>Estimated, see text.

<sup>c</sup>Calculated from  $B_0 = 0.352952 \text{ cm}^{-1}$  according to Bujin and Linton<sup>264</sup> and the  $\alpha_e$  value calculated from the Pekeris relation.

<sup>d</sup>Calculated from the Kratzer relation.

<sup>e</sup>Estimated state.

$$C_p^\circ / (\text{JK}^{-1} \text{mol}^{-1}) = 70.58620 - 20.2158 \times 10^{-3} (\text{T/K}) + 4.68050 \times 10^{-6} (\text{T/K})^2 - 0.243006 \times 10^{-9} (\text{T/K})^3 - 6.91999 \times 10^6 (\text{T/K})^{-2}$$

for the 1400–4000 K range.

#### 4.7.2. Enthalpy of formation

The results for the enthalpy of formation of  $\text{SmO}(\text{g})$  are presented in Table 41. Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurement of the weight loss for  $\text{Sm}_2\text{O}_3(\text{cr})$ . The results of these measurements were treated in this work under assumption of congruent vaporization of  $\text{Sm}_2\text{O}_3(\text{cr})$  according

to reaction



neglecting the possibility of formation of  $\text{Sm}(\text{g})$  atoms in the vapor (see Sec. 4.1). The enthalpy of formation of  $\text{SmO}(\text{g})$  thus calculated is slightly more negative in comparison with results of mass-spectrometric measurements for  $\text{Al}(\text{g}) + \text{SmO}(\text{g})$  oxygen exchange reaction.<sup>268</sup> Results of the latter work can be regarded as highly reliable, due to large number of measurements carried out using both vibrating-reed electrometer and pulse counting detection. It needs to be mentioned, however, that our treatment of data from that paper resulted in a considerable difference between second- and third-law values for the enthalpy of formation of  $\text{SmO}(\text{g})$ .

 TABLE 41. The enthalpy of formation of  $\text{SmO}(\text{g})$ , in  $\text{kJ mol}^{-1}$ 

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2333–2499	$\text{Sm}_2\text{O}_3 = 2\text{SmO}(\text{g}) + \text{O}(\text{g})$	1839.1	–113.6
	M	2360–2500	$\text{Sm}(\text{g}) + \text{YO}(\text{g}) = \text{SmO}(\text{g}) + \text{Y}(\text{g})$	135.2	–170.9
Dickson and Zare <sup>269</sup>	B	2155–2485	$\text{Sm}(\text{g}) + \text{NO}_2 = \text{SmO}(\text{g}) + \text{NO}(\text{g})$		$\leq -116.6$
Hildenbrand <sup>268</sup>	M	2087–2298	$\text{Al}(\text{g}) + \text{SmO}(\text{g}) = \text{AlO}(\text{g}) + \text{Sm}(\text{g})$	3.5	–109.0
	M	2110–2295	$\text{Al}(\text{g}) + \text{SmO}(\text{g}) = \text{AlO}(\text{g}) + \text{Sm}(\text{g})$	2.6	–101.6
Selected value:			$D_0(\text{SmO}) = 555.6 \pm 8$		–105.3 $\pm$ 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; B = beam-gas chemiluminescence.

<sup>b</sup>Recalculated by Ames *et al.*<sup>206</sup>

From the short wavelength cutoff of the Sm molecular beam and NO<sub>2</sub>(g) chemiluminescent spectra, the following lower boundary value to the ground state dissociation energy was obtained in the work of Dickson and Zare<sup>269</sup>:  $D_0(\text{SmO}) \geq (566.9 \pm 2.9)$  kJ mol<sup>-1</sup>. Comparison with both Knudsen effusion<sup>206</sup> and mass spectrometry<sup>268</sup> results reveals that this lower bound value is overestimated. The mass-spectrometric results of Ames *et al.*<sup>206</sup> seem to be erroneous.

The selected value

$$\Delta_f H^\circ(\text{SmO, g, 298.15K}) = -(105.3 \pm 8) \text{ kJ mol}^{-1}$$

is taken as a rounded average of two third-law values calculated from the data of Hildenbrand.<sup>268</sup> The selected enthalpy of formation corresponds to  $D_0(\text{SmO}) = (555.6 \pm 8)$  kJ mol<sup>-1</sup>.

## 4.8. EuO(g)

### 4.8.1. Heat capacity and entropy

The thermal functions of EuO(g) in the standard state have been calculated using the molecular constants presented in Table 42.

The electronic spectrum of EuO (single electronic transition) was investigated by McDonald,<sup>270</sup> but the data were not published. Some of the results for the ground state ( $\omega_e \approx 688$  cm<sup>-1</sup> and  $B_0 = 0.32624$  cm<sup>-1</sup> for <sup>153</sup>EuO) were cited by Dolg *et al.*<sup>271</sup> The values of the fundamental frequency of EuO were measured also in solid Ar and Kr matrices by Gabelnick *et al.*<sup>212</sup> (668 cm<sup>-1</sup>), Willson and Andrews<sup>213</sup> (667.8 cm<sup>-1</sup> and 633.5 cm<sup>-1</sup> for Eu<sup>16</sup>O and Eu<sup>18</sup>O, respectively), and Willson *et al.*<sup>246</sup> (668 cm<sup>-1</sup>). Taking into account the matrix shift, the obtained values are in agreement with the  $\omega_e$  value for EuO from gas phase. That permits to estimate the uncertainty of selected value of  $\omega_e$  to be within 2 cm<sup>-1</sup>.

The electronic structure of EuO was investigated by Carrete and Hocquet<sup>230</sup> and Dulick *et al.*<sup>247</sup> using Ligand field calculation and by Dolg *et al.*<sup>271</sup> by *ab initio* calculation. All calculations revealed the X<sup>8</sup>Σ(4f<sup>7</sup>) ground state and the first

excited state A<sup>8</sup>Σ(4f<sup>6</sup>6s). However the energy of the A<sup>8</sup>Σ state is not well defined (from 3300 up to 7937 cm<sup>-1</sup>). The theoretical calculations resulted also in relative energies for the other low-lying states of the 4f<sup>6</sup>6s configuration. In the present work we select the data obtained by Carrete and Hocquet<sup>230</sup> (all states of the 4f<sup>6</sup>(7F)6s subconfiguration with correction of the A<sup>8</sup>Σ energy (assumed 6000 cm<sup>-1</sup>). In Table 42 are also presented roughly estimated statistical weights (for 4f<sup>6</sup>6s, 4f<sup>6</sup>6d, and 4f<sup>6</sup>6p states) at fixed energies in the 15 000–40 000 cm<sup>-1</sup> interval.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (253.419 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are:

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 33.8838 + 7.70507 \times 10^{-3}(\text{T/K}) \\ &\quad - 7.38219 \times 10^{-6}(\text{T/K})^2 + 3.41476 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 2.52934 \\ &\quad \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1700 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= -79.4911 + 93.4647 \times 10^{-3}(\text{T/K}) \\ &\quad - 22.3194 \times 10^{-6}(\text{T/K})^2 + 1.86990 \\ &\quad \times 10^{-9}(\text{T/K})^3 + 5.27560 \\ &\quad \times 10^7(\text{T/K})^{-2} \end{aligned}$$

for the 1700–4000 K range.

### 4.8.2. Enthalpy of formation

The results for enthalpy of formation of EuO(g) are presented in Table 43. Ames *et al.*<sup>206</sup> carried out Knudsen effusion weight-loss measurements for Eu<sub>2</sub>O<sub>3</sub>(cr). Formal treatment of their data under the assumption of congruent evaporation with formation of EuO(g) and O(g) demonstrates the inadequacy of this approach due to high degree of EuO dissociation and the predominance of Eu(g) in the vapor phase. In this case, the results of the calculations of the enthalpy of formation are to be regarded as a lower boundary of the enthalpy of formation.

Dickson and Zare<sup>269</sup> studied the chemiluminescence resulting from the reaction of an europium molecular beam with NO<sub>2</sub>, N<sub>2</sub>O, and O<sub>3</sub> under single-collision conditions. From the short wavelength cutoff of the chemiluminescent spectra, the following lower boundary to the ground state dissociation energy was obtained from the Eu(g) + NO<sub>2</sub> study:  $D_0(\text{EuO}) = (549.8 \pm 2.9)$  kJ mol<sup>-1</sup>. This value was discarded in the paper by Murad and Hildenbrand,<sup>272</sup> in which a detailed discussion of the stability of EuO(g) was presented. The following values were calculated by Murad and Hildenbrand from the graph in the Dickson and Zare<sup>269</sup>: Eu + N<sub>2</sub>O,  $D_0(\text{EuO}) > 423$  kJ mol<sup>-1</sup>; Eu + O<sub>3</sub>,  $D_0(\text{EuO}) > 457$  kJ mol<sup>-1</sup> (not shown in Table 43). Both values are in agreement with mass-spectrometric data. Mass spectrometric measurements of isomolecular oxygen exchange reactions have been carried out by Murad and Hildenbrand,<sup>272</sup> and Balducci *et al.*<sup>273</sup> All results of these works are in reasonable agreement.

TABLE 42. Molecular constants of <sup>153</sup>Eu<sup>16</sup>O(g)

No.	State	T <sub>e</sub>	ω <sub>e</sub>	ω <sub>e</sub> x <sub>e</sub>	B <sub>e</sub>	α <sub>e</sub> 10 <sup>3</sup>	D <sub>e</sub> 10 <sup>7</sup>	r <sub>e</sub>	p <sub>i</sub>
					cm <sup>-1</sup>			pm	
0 <sup>d</sup>	X <sup>8</sup> Σ	0	688	3	0.3272 <sup>b</sup>	1.9	2.96 <sup>c</sup>	188.63	2
1 <sup>d</sup>	A <sup>8</sup> Σ	6000							8
2 <sup>d</sup>		7000							16
3 <sup>d</sup>		8000							18
4 <sup>d</sup>		9000							28
5 <sup>d</sup>		13 200							28
6 <sup>d</sup>		15 000							10
7 <sup>d</sup>		20 000							35
8 <sup>d</sup>		25 000							340
9 <sup>d</sup>		30 000							670
10 <sup>d</sup>		35 000							3300
11 <sup>d</sup>		40 000							3600

<sup>a</sup>Experimental (4f<sup>6</sup>6s) state.

<sup>b</sup>Calculated from B<sub>0</sub> = 0.32624 cm<sup>-1</sup> and the α<sub>e</sub> value calculated from the Pekeris relation.

<sup>c</sup>Calculated from the Kratzer relation.

<sup>d</sup>Estimated state.

TABLE 43. The enthalpy of formation of EuO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	1984–2188	$\text{Eu}_2\text{O}_3 = 2\text{EuO(g)} + \text{O(g)}$	1621.7	-142.7
Dickson and Zare <sup>269</sup>	B		$\text{Eu(g)} + \text{NO}_2 = \text{EuO(g)} + \text{NO(g)}$		<128.4
Murad and Hildenbrand <sup>272</sup>	M	2026–2255	$\text{Eu(g)} + \text{AlO(g)} = \text{EuO(g)} + \text{Al(g)}$	39.1	-47.4
	M	2009–2237	$\text{Eu(g)} + \text{BaO(g)} = \text{EuO(g)} + \text{Ba(g)}$	66.3	-60.5
	M	2044–2237	$\text{Eu(g)} + \text{TiO(g)} = \text{EuO(g)} + \text{Ti(g)}$	184.5	-62.9
Balducci <i>et al.</i> <sup>273</sup>	M	1920–2220	$\text{Eu(g)} + \text{WO}_3\text{(g)} = \text{EuO(g)} + \text{WO}_2\text{(g)}$	118.2	-54.4
Selected value:			$D_0(\text{EuO}) = 477.9 \pm 8$		-56.5 ± 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; B = beam-gas chemiluminescence.

The selected value

$$\Delta_f H^\circ(\text{EuO, g, 298.15 K}) = -(56.5 \pm 8) \text{ kJ mol}^{-1}$$

is taken as weighted average of the third-law values calculated from all mass-spectrometric works, with the weight of  $\text{WO}_3 + \text{Eu}$  result of 0.5, due to less accurate reference values. The selected enthalpy of formation corresponds to  $D_0(\text{EuO}) = 477.9 \text{ kJ mol}^{-1}$ .

## 4.9. GdO(g)

### 4.9.1. Heat capacity and entropy

The thermal functions of GdO(g) in the standard state have been calculated using the molecular constants presented in Table 44.

The first studies of the electronic spectrum of GdO were carried out at the low and moderate resolution and dealt with the vibrational structure of the spectrum (see Huber and Herzberg<sup>179</sup>), Yadav *et al.*,<sup>274,275</sup> Suarez and Grinfeld,<sup>276</sup> and the literature cited therein). The investigations revealed the highly complex structure of the spectrum, and the proposed vibrational analysis was doubtful. The values of the fundamental frequency of GdO were observed in solid inert gas matrices by Weltner and De Kock<sup>245</sup> in Ne and Ar (824 and 813 cm<sup>-1</sup>, respectively), by Willson and Andrews<sup>213</sup> in Ar (812.7 and 770.9 cm<sup>-1</sup> for Gd<sup>16</sup>O and Gd<sup>18</sup>O, respectively), and by Willson *et al.*<sup>246</sup> in Ar (812.8 cm<sup>-1</sup>).

Van Zee *et al.*<sup>277</sup> found by means of ESR spectroscopy in an Ar matrix at 4 K that the GdO ground state is  $^9\Sigma^-$ , and its vibrational quantum  $\Delta G_{1/2} = 824 \text{ cm}^{-1}$ . Taking into account the matrix shift, the values of the fundamental frequency observed in matrices are in agreement with the value  $\Delta G_{1/2} = 828.19 \text{ cm}^{-1}$  measured by Yadav *et al.*<sup>274</sup> for the lower state of the A system with multiple heads. Dmitriev *et al.*<sup>278</sup> analyzed the rotational spectrum of the transition connected with the ground state ( $^9\Pi_1\text{-X}^9\Sigma^-$ ). Later Dmitriev *et al.*<sup>279</sup> and Dmitriev<sup>280</sup> investigated the fluorescence spectra excited by the Ar<sup>+</sup> laser lines in the region of the  $^9\Sigma^-$ - $\text{X}^9\Sigma^-$  and  $^7\Sigma^-$ - $\text{a}^7\Sigma^-$  systems, obtained the data on the  $\nu = 0$ -8 levels of the ground state and for the  $\nu = 0$ -3 levels of the  $\text{a}^7\Sigma^-$  state, and found the excitation energy of the  $\text{a}^7\Sigma^-$  state from the intercombination transitions  $^9\Sigma^-$ - $\text{a}^7\Sigma^-$  and  $^7\Sigma^-$ - $\text{X}^9\Sigma^-$ . Till now the vibrational and rotational constants for these low-lying states and the energy of a  $^7\Sigma^-$  state are the most precise. The molecular constants for the  $\text{X}^9\Sigma^-$  and  $\text{a}^7\Sigma^-$  states (see Table 44) were reported by Gurvich *et al.*<sup>7,281</sup>. The studies of the laser induced fluorescence spectra by Carette *et al.*<sup>282</sup> and by Kaledin *et al.*<sup>258</sup> revealed new components [18.4]4 and [19.0]0<sup>-</sup> of the  $^9,7\Pi$  states. Kaledin *et al.*<sup>258</sup> showed that  $\Omega$ -assignment for the components of the  $^9\Pi$  state made by Dmitriev *et al.*<sup>278</sup> was not correct. At the same time the interpretation of the  $f^7(8S)p$  states proposed by Kaledin *et al.*<sup>258</sup> was not convincing. In any case it shows that the  $^9\Pi_5$  ([17.6]5) and  $^9\Pi_4$  ([18.4]4) components known experimentally belong to the states of different multiplicity.

TABLE 44. Molecular constants of  $^{158}\text{Gd}^{16}\text{O(g)}$ 

No	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		cm <sup>-1</sup>							pm
0 <sup>a</sup>	$\text{X}^9\Sigma^-$	0	831.19	2.619	0.35561	1.48	2.56	180.66	9
1 <sup>a</sup>	$\text{a}^7\Sigma^-$	1834.64	837.11	2.62	0.35678	1.47	2.59	180.37	7
2 <sup>a</sup>	$\text{A}^9\Delta?$	11 300							18
3 <sup>a</sup>		17 500							50
4 <sup>a</sup>		20 000							28
5 <sup>a</sup>	$^9\Sigma^-$	21 693 <sup>b</sup>							9
6 <sup>a</sup>	$^7\Sigma^-$	22 307 <sup>b</sup>							7
7 <sup>c</sup>		25 000							16
8 <sup>c</sup>		30 000							200
9 <sup>c</sup>		35 000							440
10 <sup>c</sup>		40 000							1050

<sup>a</sup>Experimental state.

<sup>b</sup>All the constants for these state from Dmitriev *et al.*<sup>279</sup> and Dmitriev.<sup>280</sup>

<sup>c</sup>Estimated state.



*Ab initio* calculations of GdO by Dolg *et al.*,<sup>271,283</sup> dealt with well-known electronic states  $X^9\Sigma^-$  and  $a^7\Sigma^-$  and resulted in theoretical constants in good agreement with the experiment. Ligand field calculation by Carrete and Hocquet,<sup>230</sup> Dulick *et al.*<sup>247</sup> and *ab initio* calculations by Kotzian *et al.*<sup>197</sup> and Sakai *et al.*<sup>284</sup> considered the electronic structure of the molecule. All calculations gave the  $X^9\Sigma^- f^7(8S)s$  ground state and the  $a^7\Sigma^- f^7(8S)s$  first excited state. The half filled  $4f$  shell along with the ground  $8S$  state and the other states of the Gd atom lying much higher ( $>30\,000\text{ cm}^{-1}$ ) explain the simple structure of the GdO molecule: only 12 states of the  $f^7(8S)s$ ,  $f^7(8S)d$ , and  $f^7(8S)p$  subconfigurations are expected up to  $30\,000\text{ cm}^{-1}$ . All the calculations except that by Dulick *et al.*,<sup>247</sup> who calculated only the  $f^7(8S)s$  states, are in agreement that the first excited configuration of GdO is  $f^7(8S)d$ . However the energy of the  $A^9\Delta$  state differs considerably (from  $8500$  to  $16700\text{ cm}^{-1}$ ). The study of the GdO<sup>-</sup> photoelectron spectrum by Klingeler *et al.*<sup>285</sup> revealed the group of states near  $11300 \pm 400\text{ cm}^{-1}$  which positions coincide with the  $A^9\Delta$  state calculated by Carrete and Hocquet.<sup>230</sup> The estimated energies and statistical weights of the unobserved states of the configuration mentioned above are presented in Table 44. It gives also the estimated statistical weights in the  $30\,000$ – $40\,000\text{ cm}^{-1}$  region (the  $f^7s$  and  $f^7d$  states).

The derived standard entropy at room temperature is:

$$S^\circ(298.15\text{ K}) = (253.495 \pm 0.03)\text{ J K}^{-1}\text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are:

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) &= 21.26451 + 40.9137 \times 10^{-3}(T/\text{K}) \\ &\quad - 30.1655 \times 10^{-6}(T/\text{K})^2 + 7.50616 \\ &\quad \times 10^{-9}(T/\text{K})^3 + 6.83276 \\ &\quad \times 10^4(T/\text{K})^{-2} \end{aligned}$$

for the  $298.15$ – $1300\text{ K}$  range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) &= 51.77714 - 10.0325 \times 10^{-3}(T/\text{K}) \\ &\quad + 2.57810 \times 10^{-6}(T/\text{K})^2 - 0.111668 \\ &\quad \times 10^{-9}(T/\text{K})^3 - 4.80380 \\ &\quad \times 10^6(T/\text{K})^{-2} \end{aligned}$$

for the  $1300$ – $4000\text{ K}$  range.

#### 4.9.2. Enthalpy of formation

The results for the enthalpy of formation of GdO(g) are presented in Table 45. Experimental data used in the analysis are based on Knudsen effusion<sup>206</sup> and mass spectrometric

measurements,<sup>260</sup> the results being in satisfactory agreement. At the same time, the mass-spectrometric results deserve more weight, because interpretation of Knudsen effusion data is not unambiguous due to possible formation of free gadolinium atoms in the vapors (see text on LaO(g) enthalpy of formation).

The selected value

$$\Delta_f H^\circ(\text{GdO, g, } 298.15\text{ K}) = -(68.0 \pm 8)\text{ kJ mol}^{-1}$$

is taken as a rounded average of the third-law values calculated from two mentioned works, with a low weight for the Knudsen effusion results. The selected enthalpy of formation corresponds to  $D_0(\text{GdO}) = (710.2 \pm 8)\text{ kJ mol}^{-1}$ .

### 4.10. TbO(g)

#### 4.10.1. Heat capacity and entropy

The thermal functions of TbO(g) in the standard state have been calculated using the molecular constants given in Table 46.

The electronic spectrum of TbO was investigated in emission and absorption by Kaledin and Shenyavskaya<sup>286,287</sup> (see Huber and Herzberg<sup>179</sup> for earlier works) and in the fluorescence by Kulikov *et al.*<sup>288,289</sup>, and Gurvich *et al.*<sup>281</sup> The fundamental frequency of TbO was observed in solid inert gas matrices by Weltner and De Kock<sup>245</sup> in Ne and Ar ( $824\text{ cm}^{-1}$ ), by Willson and Andrews<sup>213</sup> in Ar at  $10\text{ K}$  ( $823.9$  and  $781.4\text{ cm}^{-1}$  for  $\text{Tb}^{16}\text{O}$  and  $\text{Tb}^{18}\text{O}$ , respectively), and by Willson *et al.*<sup>246</sup> in Ar ( $823.9\text{ cm}^{-1}$ ). Taking into account the matrix shift the values of fundamental frequency obtained in matrices are in agreement with those from gas phase data. *Ab initio* calculations by Dolg and Stoll<sup>193</sup> dealt with the ground state of the TbO molecule and resulted in the theoretical spectroscopic constants.

The studies of the TbO electronic spectra revealed 14 low-lying states (including the X6.5 ground state), assigned to the  $f^8(7F)s$  subconfiguration, and 11 excited states with energies  $14\,899$ – $22\,300\text{ cm}^{-1}$ , mostly ascribed to the  $f^8(7F)d$  and  $f^8(7F)p$  subconfigurations. Ligand field calculation by Carrete and Hocquet<sup>230</sup> and Dulick *et al.*<sup>247</sup> showed that up to  $10\,000\text{ cm}^{-1}$  all the states belonged to the  $f^8(7F)s$  subconfiguration. The estimated energies and statistical weights of the unobserved states of the  $f^8(7F)s$  subconfiguration, presented in Table 46, are selected from Carrete and Hocquet.<sup>230</sup> Table 46 gives also the estimated statistical weights of the  $f^8(7F)d$  and  $f^8(7F)p$  states assuming the energies of the lowest states at  $12\,000$  and  $18\,000\text{ cm}^{-1}$ , respectively, and placing the other states of the  $f^8nl$  configurations higher than the corresponding  $f^8(7F)l$  states by approximately  $15\,000$ – $20\,000\text{ cm}^{-1}$ .

TABLE 45. The enthalpy of formation of GdO(g), in  $\text{kJ mol}^{-1}$

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15\text{ K})$	$\Delta_f H^\circ(298.15\text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2408–2546	$\text{Gd}_2\text{O}_3 = 2\text{GdO(g)} + \text{O(g)}$	1935.6	–78.5
Murad and Hildenbrand <sup>260</sup>	M	1957–2024	$\text{Gd(g)} + \text{TiO(g)} = \text{GdO(g)} + \text{Ti(g)}$	–43.5	–68.5
	M	1908–1977	$\text{Gd(g)} + \text{YO(g)} = \text{GdO(g)} + \text{Y(g)}$	5.7	–65.4
Selected value:			$D_0(\text{GdO}) = 710.2 \pm 8$		–68.0 $\pm$ 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry.

TABLE 46. Molecular constants of  $^{159}\text{Tb}^{16}\text{O}(\text{g})$ 

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		$\text{cm}^{-1}$					pm		
0 <sup>a</sup>	X(1)6.5	0	843.1 <sup>b</sup>	3.04 <sup>b</sup>	0.35325	1.50	2.48 <sup>c</sup>	181.213	2
1 <sup>a</sup>	(1)5.5	355							2
2 <sup>a</sup>	(1)4.5	750							2
3 <sup>a</sup>	(1)3.5	1183							2
4 <sup>a</sup>	(2)5.5	1624							2
5 <sup>a</sup>	(1)2.5	1654							2
6 <sup>a</sup>	(2)4.5	2101							2
7 <sup>a</sup>	(1)1.5	2160							2
8 <sup>a</sup>	(2)3.5	2610							2
9 <sup>a</sup>	(3)5.5	2900							2
10 <sup>a</sup>	(2)2.5	3119							2
11 <sup>a</sup>	(3)4.5	3188							2
12 <sup>a</sup>	(3)3.5	3506							2
13 <sup>a</sup>	(3)2.5	3830							2
14 <sup>d</sup>		4000							4
15 <sup>d</sup>		4500							8
16 <sup>d</sup>		5100							8
17 <sup>d</sup>		6100							16
18 <sup>d</sup>		7050							8
19 <sup>d</sup>		7950							10
20 <sup>d</sup>		9050							8
21 <sup>d</sup>		10 100							9
22 <sup>d</sup>		15 000							25
23 <sup>d</sup>		20 000							80
24 <sup>d</sup>		25 000							125
25 <sup>d</sup>		30 000							170
26 <sup>d</sup>		35 000							600
27 <sup>d</sup>		40 000							900

<sup>a</sup>Experimental state.<sup>b</sup>Calculated from  $\Delta G_{1/2} = 837.1 \text{ cm}^{-1}$  given by Kulikov *et al.*<sup>288</sup> and the assumed dissociation limit.<sup>c</sup>Calculated from the Kratzer relation.<sup>d</sup>Estimated state.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (245.758 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 30.97785 + 32.5225 \times 10^{-3} (\text{T/K}) \\ &\quad - 19.6568 \times 10^{-6} (\text{T/K})^2 + 4.14576 \\ &\quad \times 10^{-9} (\text{T/K})^3 - 1.18135 \\ &\quad \times 10^4 (\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1400 K range, and

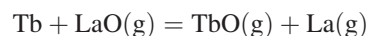
$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 64.48191 - 8.31957 \times 10^{-3} (\text{T/K}) \\ &\quad + 0.883659 \times 10^{-6} (\text{T/K})^2 \\ &\quad + 0.0270508 \times 10^{-9} (\text{T/K})^3 - 10.4723 \\ &\quad \times 10^6 (\text{T/K})^{-2} \end{aligned}$$

for the 1400–4000 K range.

#### 4.10.2. Enthalpy of formation

There is only one study of the determination of the enthalpy of formation of  $\text{TbO}(\text{g})$ . Ames *et al.*<sup>206</sup> have performed mass spectrometric measurements of isomolecular oxygen

exchange reaction (1830–2040 K)



for which we obtain  $\Delta_f H^\circ(298.15 \text{ K}) = 80.2 \text{ kJ mol}^{-1}$  by third-law analysis. This results in the following selected enthalpy of formation:

$$\Delta_f H^\circ(\text{TbO}, \text{g}, 298.15 \text{ K}) = -(84.7 \pm 10) \text{ kJ mol}^{-1}$$

This value corresponds to  $D_0(\text{TbO}) = (715.5 \pm 10) \text{ kJ mol}^{-1}$ .

### 4.11. DyO(g)

#### 4.11.1. Heat capacity and entropy

The thermal functions of  $\text{DyO}(\text{g})$  in the standard state have been calculated using the molecular constants given in Table 47.

The electronic spectrum of  $\text{DyO}$  was investigated in emission and absorption by Kaledin and Shenyavskaya<sup>286</sup> and Yadav *et al.*<sup>290</sup> (see Huber and Herzberg<sup>179</sup> for earlier works) and in the fluorescence by Linton *et al.*<sup>291</sup> and Cheng.<sup>292</sup> The values of the fundamental frequency of  $\text{DyO}$  were observed in solid inert gas matrices by Weltner and De Kock<sup>245</sup> in Ar ( $829 \text{ cm}^{-1}$ ), Willson and Andrews<sup>213</sup> in Ar at 10 K ( $829.0$  and

TABLE 47. Molecular constants of  $^{159}\text{Dy}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$					$r_e$ pm	$p_i$	
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$			
0 <sup>a</sup>	X(1)8	0	849 <sup>b</sup>	3.5 <sup>b</sup>	0.3593 <sup>c</sup>	2.0	2.57 <sup>d</sup>	179.4	2
1 <sup>a</sup>	(1)7	770							2
2 <sup>a</sup>	(1)6	1630							2
3 <sup>a</sup>	(2)7	1683							2
4 <sup>a</sup>	(2)6	2420							2
5 <sup>e</sup>		2180							2
6 <sup>e</sup>		3280							11
7 <sup>e</sup>		4400							10
8 <sup>e</sup>		5100							7
9 <sup>e</sup>		6100							8
10 <sup>e</sup>		7130							14
11 <sup>e</sup>		8565							14
12 <sup>e</sup>		9830							20
13 <sup>e</sup>		11 775							20
14 <sup>e</sup>		14 070							20
15 <sup>e</sup>		15 000							40
16 <sup>e</sup>		20 000							130
17 <sup>e</sup>		25 000							350
18 <sup>e</sup>		30 000							370
19 <sup>e</sup>		35 000							1500
20 <sup>e</sup>		40 000							1900

<sup>a</sup>Experimental state.<sup>b</sup>Calculated from  $\Delta G_{1/2} = 842 \text{ cm}^{-1}$  given by Linton *et al.*<sup>291</sup> and the assumed dissociation limit.<sup>c</sup>Calculated from  $B_0 = 0.358413 \text{ cm}^{-1}$  given by Linton *et al.*<sup>291</sup> and  $\alpha_e = 0.0020 \text{ cm}^{-1}$  found by Kaledin and Shenyavskaya.<sup>286</sup><sup>d</sup>Calculated from the Kratzer relation.<sup>e</sup>Estimated state.

786.1  $\text{cm}^{-1}$  for  $\text{Dy}^{16}\text{O}$ ) and  $\text{Dy}^{18}\text{O}$ , respectively), and Willson *et al.*<sup>246</sup> in Ar (828.7  $\text{cm}^{-1}$ ). Taking into account the matrix shift, the values of the fundamental frequency obtained in matrices are in agreement with those from gas phase data. *Ab initio* calculations by Dolg and Stoll<sup>193</sup> dealt with the ground state of the molecule and resulted in theoretical spectroscopic constants.

The studies of the DyO electronic spectra by Kaledin and Shenyavskaya<sup>286</sup> and Linton *et al.*<sup>291</sup> revealed 5 low-lying states (including the X8 ground state,) which were assigned to the  $f^9(6H)s$  subconfiguration, and 6 excited states with energies 17 070–19 000  $\text{cm}^{-1}$ , most of which could be assigned to the  $f^9(6H)d$  and  $f^9(6H)p$  configurations.

Carrete and Hocquet<sup>230</sup> and Dulick *et al.*<sup>247</sup> carried out Ligand field calculations of the DyO states belonging to the  $f^9(6H)s$  subconfiguration. Carrete and Hocquet<sup>230</sup> obtained all the  $f^9(6H)s$  states (total statistical weight 132), while Dulick *et al.*<sup>247</sup> obtained only states up to 10 000  $\text{cm}^{-1}$  with  $\Sigma p = 74$ . The data obtained by Dulick *et al.*<sup>247</sup> are closer to experimental data because of using the other parameter  $G_3 = 300 \text{ cm}^{-1}$  as compared with  $G_3 = 150 \text{ cm}^{-1}$  used by Carrete and Hocquet.<sup>230</sup> The deviations between the results are the largest for lower states. For that reason the estimated energies and statistical weights of the unobserved states of the  $f^9(6H)s$  subconfiguration up to 10 000  $\text{cm}^{-1}$  are selected from the Dulick *et al.*<sup>247</sup> and those in the interval 10 000–15 000  $\text{cm}^{-1}$  are selected from Carrete and Hocquet<sup>230</sup> with corrections to mean deviation. In Table 47 also the estimated statistical weights for the  $f^9(6FP)s$  states (>15 000  $\text{cm}^{-1}$ ) are given, for

the remaining  $f^9s$  states corresponding to the low multiplicity terms of the  $f^9$ -core (>21 000  $\text{cm}^{-1}$ ), for the  $f^9(6H)d$  and  $f^9(6H)p$  states assuming the energies of the lowest states at 12 500 and 18 500  $\text{cm}^{-1}$ , respectively, and for the other states of the  $f^9nl$  configurations as in case of the  $f^9s$  configuration. The upper limit for all configurations is assumed to be  $D_0 + IP$ , so all the estimated states are considered stable.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (242.208 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 19.08366 + 61.7856 \times 10^{-3}(\text{T/K}) \\ & - 37.4790 \times 10^{-6}(\text{T/K})^2 + 7.29509 \\ & \times 10^{-9}(\text{T/K})^3 + 2.19805 \\ & \times 10^4(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1300 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 68.49712 - 11.3191 \times 10^{-3}(\text{T/K}) \\ & + 1.99999 \times 10^{-6}(\text{T/K})^2 \\ & - 0.0418797 \times 10^{-9}(\text{T/K})^3 \\ & - 8.39005 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1300–4000 K range.

#### 4.11.2. Enthalpy of formation

Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurements of the weight loss of  $\text{Dy}_2\text{O}_3(\text{cr})$  in the temperature range 2440–2637 K. The results of these measurements were treated in this work under the assumption of congruent vaporization of  $\text{Dy}_2\text{O}_3(\text{cr})$  according to reaction



neglecting possible formation of Dy(g) atoms in the vapor (see Sec. 4.1). The DyO(g) enthalpy of formation thus calculated must be more negative than the correct value, the degree of deviation being dependent on amount of Dy atoms in the vapor. The third-law enthalpy of reaction  $\Delta_f H^\circ(298.15 \text{ K}) = 1953.4 \text{ kJ mol}^{-1}$ , yields  $\Delta_f H^\circ(\text{DyO}, \text{g}, 298.15 \text{ K}) = -79.2 \text{ kJ mol}^{-1}$ , which corresponds to  $D_0(\text{DyO}) = 609.8 \text{ kJ mol}^{-1}$ . Dulick *et al.*<sup>247</sup> have estimated the DyO dissociation energy using the crystal field model applied to diatomic molecules as  $D_0(\text{DyO}) = (602 \pm 1) \text{ kJ mol}^{-1}$ . In absence of additional information, this value is selected in this work for the enthalpy of formation of DyO(g):

$$\Delta_f H^\circ(\text{DyO}, \text{g}, 298.15 \text{ K}) = -(71 \pm 20) \text{ kJ mol}^{-1}.$$

### 4.12. HoO(g)

#### 4.12.1. Heat capacity and entropy

The thermal functions of HoO(g) in the standard state have been calculated using the molecular constants given in Table 48.

TABLE 48. Molecular constants of  $^{165}\text{Ho}^{16}\text{O}(\text{g})$ 

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		$\text{cm}^{-1}$						pm	
0 <sup>a</sup>	X8.5	0	848 <sup>b</sup>	3.5 <sup>b</sup>	0.358477	1.50	2.56 <sup>c</sup>	179.59	2
1 <sup>a</sup>	(1)7.5	603							2
2 <sup>a</sup>	(2)7.5	1130							2
3 <sup>a</sup>	(1)6.5	1853							2
4 <sup>d</sup>		1165							14
5 <sup>d</sup>		2275							12
6 <sup>d</sup>		6300							22
7 <sup>d</sup>		7100							8
8 <sup>d</sup>		10 050							26
9 <sup>d</sup>		13 500							26
10 <sup>d</sup>		15 000							50
11 <sup>d</sup>		20 000							90
12 <sup>d</sup>		25 000							200
13 <sup>d</sup>		30 000							250
14 <sup>d</sup>		35 000							700
15 <sup>d</sup>		40 000							1150

<sup>a</sup>Experimental state.<sup>b</sup>Calculated from  $\Delta G_{1/2} = 841.252 \text{ cm}^{-1}$  given by Linton and Liu<sup>294</sup> and the assumed dissociation limit.<sup>c</sup>Calculated from the Kratzer relation.<sup>d</sup>Estimated state.

The electronic spectrum of HoO was investigated in emission and absorption by Kaledin and Shenyavskaya<sup>286</sup> (see Huber and Herzberg<sup>179</sup> for earlier works), and in fluorescence by Liu *et al.*,<sup>293</sup> Linton and Liu,<sup>294</sup> and Cheng.<sup>292</sup> The values of fundamental frequency of HoO were measured in solid inert gas matrices by Weltner and De Kock<sup>245</sup> in Ar (829  $\text{cm}^{-1}$ ), Willson and Andrews<sup>213</sup> in Ar at 10 K (828.1 and 785.2  $\text{cm}^{-1}$  for Ho<sup>16</sup>O and Ho<sup>18</sup>O, respectively), and Willson *et al.*<sup>246</sup> in Ar (828.0  $\text{cm}^{-1}$ ). Taking into account the matrix shift, the values of fundamental frequency obtained in matrices are in agreement with those from the gas phase data. *Ab initio* calculations by Dolg and Stoll<sup>193</sup> dealt with the ground state of the molecule and gave its constants in good agreement with experiment.

The studies of the HoO electronic spectra revealed 4 low-lying states (including the X8.5 ground state) which were assigned to the  $f^{10}(5I)s$  subconfiguration and 6 excited states with energies 17 600–22 400  $\text{cm}^{-1}$  most of which could be assigned to the  $f^{10}(5I)d$  and  $f^{10}(5I)p$  superconfigurations. The Ligand field calculations were carried out by Carrete and Hocquet,<sup>230</sup> who considered all the  $f^{10}(5I)s$  states with the total statistical weight 130, and by Dulick *et al.*,<sup>247</sup> who did only the  $f^{10}(5I)s$  states up to 10 000  $\text{cm}^{-1}$  with  $\Sigma p = 74$ . The results obtained by Dulick *et al.*<sup>247</sup> were closer to experimental data because of using the parameter  $G_3 = 300 \text{ cm}^{-1}$  as compared with  $G_3 = 150 \text{ cm}^{-1}$  used by Carrete and Hocquet.<sup>230</sup> The deviations were the largest for lower states. That is why in Table 48 the estimated energies and statistical weights for the unobserved states of the  $f^{10}(5I)s$  subconfiguration up to 10 000  $\text{cm}^{-1}$  are taken from Dulick *et al.*,<sup>247</sup> and the data in the interval 10 000–15 000  $\text{cm}^{-1}$  are taken from Carrete and Hocquet,<sup>230</sup> with corrections for mean deviation. In Table 48 also the estimated statistical weights for the  $f^{10}(5SDFG)s$  states from 15 000  $\text{cm}^{-1}$  are given, for the remaining  $f^{10}s$  states higher than 21 500  $\text{cm}^{-1}$ , and for the  $f^{10}(5I)d$  and  $f^{10}(5I)p$  states assuming the energies

of the lowest states at 12 500 and 18 500  $\text{cm}^{-1}$ , respectively, and placing the other states of the  $f^{10}nl$  configurations as in case of the  $f^{10}s$  configuration. The upper limit for all configurations is assumed to be a sum of the dissociation energy and the ionization potential, so that all the estimated states are considered stable.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (244.590 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 48.31232 + 76.8332 \times 10^{-3}(\text{T/K}) \\ & - 155.715 \times 10^{-6}(\text{T/K})^2 + 76.9567 \\ & \times 10^{-9}(\text{T/K})^3 - 1.57926 \\ & \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–900 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 43.11154 - 5.49748 \times 10^{-3}(\text{T/K}) \\ & + 3.07490 \times 10^{-6}(\text{T/K})^2 - 0.350566 \\ & \times 10^{-9}(\text{T/K})^3 + 4.11955 \\ & \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 900–4000 K range.

#### 4.12.2. Enthalpy of formation

The results for the enthalpy of formation of HoO(g) are presented in Table 49. Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurements of the weight loss of Ho<sub>2</sub>O<sub>3</sub>(cr). The results of these measurements were treated in this work under the assumption of congruent vaporization of Ho<sub>2</sub>O<sub>3</sub>(cr) according to reaction





TABLE 49. The enthalpy of formation of HoO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2487–2711	Ho <sub>2</sub> O <sub>3</sub> = 2HoO(g) + O(g)	1988.4	-71.0
Murad and Hildenbrand <sup>260</sup>	M	1855–2178	Ho(g) + TiO(g) = HoO(g) + Ti(g)	65.5	-57.8
Selected value:			$D_0(\text{HoO}) = 604.1 \pm 10$		-57.8 ± 10

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; T = transpiration.

neglecting the possible formation of Ho(g) atoms in the vapor (see Sec. 4.1). The enthalpy of formation of HoO(g) so calculated must be more negative than the correct value, the degree of deviation being dependent on amount of Ho atoms in vapors. Comparison with the results of Murad and Hildenbrand<sup>260</sup> confirms this conclusion. The selected value

$$\Delta_f H^\circ(\text{HoO, g, 298.15 K}) = -(57.8 \pm 10) \text{ kJ mol}^{-1}$$

is taken as a rounded third-law value calculated from the latter work. The selected enthalpy of formation corresponds to  $D_0(\text{HoO}) = (604.1 \pm 10) \text{ kJ mol}^{-1}$ .

### 4.13. ErO(g)

#### 4.13.1. Heat capacity and entropy

Thermal functions of ErO(g) in the standard state have been calculated using the molecular constants presented in Table 50.

The electronic spectrum of ErO was investigated in emission and absorption by Kaledin and Shenyavskaya<sup>295</sup> (see Huber and Herzberg<sup>179</sup> for references to earlier works). The observed spectrum was highly complex. Only one rotational band at 5066 Å, observed in emission and absorption, was analyzed. It was supposed to be connected with the X8 ground state of the  $f^1(4I)s$  subconfiguration as predicted by Field.<sup>233</sup>

TABLE 50. Molecular constants of <sup>166</sup>Er<sup>16</sup>O(g)

No.	State	$\text{cm}^{-1}$					$r_e$ pm	$p_i$	
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$			
0 <sup>a</sup>	X0 <sup>-</sup>	0	849 <sup>b</sup>	3.54 <sup>b</sup>	0.359	1.9 <sup>c</sup>	2.57 <sup>d</sup>	179.4	1
1 <sup>c</sup>		25							2
2 <sup>c</sup>		95							2
3 <sup>c</sup>		205							2
4 <sup>c</sup>		350							2
5 <sup>c</sup>		525							7
6 <sup>c</sup>		730							4
7 <sup>c</sup>		1030							6
8 <sup>c</sup>		1550							4
9 <sup>c</sup>		2100							2
10 <sup>c</sup>		6115							14
11 <sup>c</sup>		6920							14
12 <sup>c</sup>		11 500							24
13 <sup>c</sup>		15 700							50
14 <sup>c</sup>		20 000							70
15 <sup>c</sup>		25 000							80
16 <sup>c</sup>		30 000							280
17 <sup>c</sup>		35 000							370
18 <sup>c</sup>		40 000							450

<sup>a</sup>Experimental state.

<sup>b</sup>Calculated from  $\Delta G_{1/2} = 841.5 \text{ cm}^{-1}$  (see text) and the assumed dissociation limit.

<sup>c</sup>Calculated from the Kratzer relation.

<sup>d</sup>Calculated from the Pekeris relation.

<sup>e</sup>Estimated state.

The values of the fundamental frequency of ErO were measured in solid inert gas matrices by Weltner and De Kock<sup>245</sup> in Ar (829 cm<sup>-1</sup>), Willson and Andrews<sup>213</sup> in Ar at 10 K (828.5 and 785.5 cm<sup>-1</sup> for Er<sup>16</sup>O and Er<sup>18</sup>O, respectively), and Willson *et al.*<sup>246</sup> in Ar (828.5 cm<sup>-1</sup>). Taking into account the matrix shift (~13 cm<sup>-1</sup>), the value 841.5 cm<sup>-1</sup> is obtained for  $\Delta G_{1/2}$  in the gas phase.

In the present work, the vibrational constants for ErO are calculated from above mentioned value  $\Delta G_{1/2} = 841.5 \text{ cm}^{-1}$  and the assumed value of the dissociation limit. Rotational constants are calculated using  $B_0 = 0.3583 \text{ cm}^{-1}$  from Kaledin and Shenyavskaya<sup>295</sup> and the well known Pekeris and Kratzer relations.

The molecular constants for all lanthanide monoxides were calculated *ab initio* using pseudopotentials having the 4f orbitals in the core by Dolg and Stoll<sup>193</sup> and compared with experimental data. Although there was no overall agreement between calculated and experimental data in the LnO series, the calculations indicated that the vibrational constants and internuclear distances of the lanthanide monoxides in electronic states of the same type of configurations changed very regularly. The selected values for  $r_e$  (ErO) = 179.4 pm and  $\omega_e = 849 \text{ cm}^{-1}$  can be compared with  $r_e(\text{DyO}) = 179.4 \text{ pm}$ ,  $r_e(\text{HoO}) = 179.59 \text{ pm}$ , and  $\omega_e(\text{DyO}) = 849 \text{ cm}^{-1}$  and  $\omega_e(\text{HoO}) = 848 \text{ cm}^{-1}$ .

According to Field,<sup>233</sup> the ground state subconfiguration of ErO is  $f^1(4I)s$  and the ground state is X8. Dulick *et al.*<sup>247</sup> and Carrete and Hocquet<sup>230</sup> carried out the Ligand field calculation of the subconfiguration  $f^1(4I)s$  and confirmed the X0<sup>-</sup> ground state. Carrete and Hocquet<sup>230</sup> considered all the  $f^1(4I)s$  states (total statistical weight 104), while Dulick *et al.*<sup>247</sup> gave only states up to 10 000 cm<sup>-1</sup> with  $\Sigma p = 60$ . The results obtained by Dulick *et al.*<sup>247</sup> and Carrete and Hocquet<sup>230</sup> were very close for three lowest states. In the present work we use the estimated energies of the  $f^1(4I)s$  states as recommended by Carrete and Hocquet.<sup>230</sup> Table 50 presents also the estimated statistical weights for the other  $f^1s$  states (higher 17 000 cm<sup>-1</sup>), and for the  $f^1d$ ,  $f^1p$ , and  $f^12$  states (higher 13 000 cm<sup>-1</sup>, 20 000 cm<sup>-1</sup>, and higher 40 000 cm<sup>-1</sup>, respectively).

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (256.473 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1},$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 43.23815 - 4.71231 \times 10^{-3} (\text{T/K}) \\ & + 0.571735 \times 10^{-6} (\text{T/K})^2 + 0.747515 \\ & \times 10^{-9} (\text{T/K})^3 - 2.55285 \\ & \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1300 K range, and

TABLE 51. The enthalpy of formation of ErO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2492–2687	$\text{Er}_2\text{O}_3 = 2\text{ErO(g)} + \text{O(g)}$	2017.1	–38.5
Murad and Hildenbrand <sup>260</sup>	M	1855–2165	$\text{Er(g)} + \text{TiO(g)} = \text{ErO(g)} + \text{Ti(g)}$	–76.8	–32.9
Selected value:			$D_0(\text{ErO}) = 593.7 \pm 8$		–32.9 ± 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry.

$$C_p^\circ/(\text{JK}^{-1} \text{mol}^{-1}) = 33.60868 + 3.85434 \times 10^{-3}(\text{T/K}) \\ - 0.382306 \times 10^{-6}(\text{T/K})^2 \\ + 0.0464068 \times 10^{-9}(\text{T/K})^3 + 2.52559 \\ \times 10^6(\text{T/K})^{-2}$$

for the 1300–4000 K range.

#### 4.13.2. Enthalpy of formation

The results for the enthalpy of formation of ErO(g) are presented in Table 51. Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurement of weight loss for several Ln<sub>2</sub>O<sub>3</sub>(cr) oxides. As in the case of Sm<sub>2</sub>O<sub>3</sub>(cr) and other lanthanide oxides, results of these measurements were treated under assumption of congruent vaporization of Er<sub>2</sub>O<sub>3</sub>(cr) according to reaction



neglecting the possibility of formation of Er(g) atoms in the vapor. In case of considerable degree of ErO dissociation this treatment will result in overestimated ErO stability. Mass spectrometric measurements of isomolecular oxygen exchange reaction have been carried out by Murad and Hildenbrand.<sup>260</sup> Results of both papers do not differ significantly, the enthalpy of formation of ErO(g) calculated from results of Ames *et al.* being only ~5 kJ mol<sup>-1</sup> more negative than that obtained from mass-spectrometric measurements of Murad and Hildenbrand. This closeness demonstrates that Er(g) does not predominate in the Er<sub>2</sub>O<sub>3</sub>(cr) vapor, in accordance with mass-spectrometric data on ErO<sup>+</sup>/Er<sup>+</sup> ratio.<sup>206</sup>

The selected value

$$\Delta_f H^\circ(\text{ErO, g, 298.15 K}) = -(32.9 \pm 8) \text{ kJ mol}^{-1}$$

is taken as the rounded third-law value from the work of Murad and Hildenbrand.<sup>260</sup> The selected enthalpy of formation corresponds to  $D_0(\text{ErO}) = (593.7 \pm 8) \text{ kJ mol}^{-1}$ .

### 4.14. TmO(g)

#### 4.14.1. Heat capacity and entropy

The thermal functions of TmO(g) in the standard state have been calculated using the molecular constants presented in Table 52.

The spectrum of TmO was measured only by infrared spectroscopy in solid inert gas matrices by Weltner and De Kock<sup>245</sup> and Willson *et al.*<sup>246</sup> in Ar (832 cm<sup>-1</sup>) and by Willson and Andrews<sup>213</sup> in Ar at 10 K (832.0 cm<sup>-1</sup> and 788.9 cm<sup>-1</sup> for Tm<sup>16</sup>O and Tm<sup>18</sup>O, respectively).

The molecular constants for all lanthanide monoxides were calculated *ab initio* by Dolg and Stoll<sup>193</sup> and compared with experimental data. Although there was no overall agreement between calculated and experimental data in the LnO series (see also Sec. 7.2.3), the calculations indicated that the vibrational constants and internuclear distances of the lanthanide monoxides in electronic states of the same type of configurations changed very regularly. In the present work we accept the extrapolated value of  $r_e(\text{TmO}) = 179.0 \text{ pm}$  considering  $r_e(\text{HoO}) = 179.59 \text{ pm}$  and  $r_e(\text{ErO}) = 179.4 \text{ pm}$ . The accepted value  $\Delta G_{1/2} = (845 \pm 5) \text{ cm}^{-1}$  for TmO is estimated from the matrix value  $832 \text{ cm}^{-1}$  taking into account matrix shift about  $13 \text{ cm}^{-1}$ . From this value and the dissociation limit according to Birge-Sponer extrapolation one gets  $\omega_e = 853.5 \text{ cm}^{-1}$  and  $\omega_e x_e = 4.24 \text{ cm}^{-1}$  (compare  $\omega_e(\text{HoO}) = 848 \text{ cm}^{-1}$ ).

According to Field<sup>233</sup> the ground state superconfiguration of TmO is  $f^{12}s$ . Dulick *et al.*<sup>247</sup> and Carrete and Hocquet<sup>230</sup> carried out the Ligand field calculation of the subconfiguration  $4f^{12}(3H)6s$ . Kotzian *et al.*<sup>197</sup> calculated the  $4f^{12}(3H)6s$  and some of the  $4f^{12}(3F)6s$  states using the Intermediate Neglect of

TABLE 52. Molecular constants of <sup>169</sup>Tm<sup>16</sup>O(g)

No.	State	$\text{cm}^{-1}$					$r_e$ pm	$p_i$	
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$a_e 10^3$			$D_e 10^7$
0 <sup>a</sup>	X(3.5)	0	853.5 <sup>b</sup>	4.24 <sup>b</sup>	0.360	2.0 <sup>c</sup>	2.56 <sup>d</sup>	179.0	2
1 <sup>c</sup>			24						2
2 <sup>c</sup>			190						2
3 <sup>c</sup>			320						2
4 <sup>c</sup>			420						2
5 <sup>c</sup>			510						2
6 <sup>c</sup>			650						4
7 <sup>c</sup>			860						2
8 <sup>c</sup>			1670						2
9 <sup>c</sup>			1910						2
10 <sup>c</sup>			2560						4
11 <sup>c</sup>			5230						4
12 <sup>c</sup>			6490						14
13 <sup>c</sup>			8100						14
14 <sup>c</sup>			8900						4
15 <sup>c</sup>			10 050						4
16 <sup>c</sup>			12 050						10
17 <sup>c</sup>			13 180						8
18 <sup>c</sup>			15 000						50
19 <sup>c</sup>			20 000						700
20 <sup>c</sup>			25 000						90
21 <sup>c</sup>			30 000						100
22 <sup>c</sup>			35 000						110
23 <sup>c</sup>			40 000						130

<sup>a</sup>Experimental state.

<sup>b</sup>Estimated (see text).

<sup>c</sup>Calculated from the Pekeris relation.

<sup>d</sup>Calculated from the Kratzer relation.

<sup>e</sup>Estimated state.

Differential Overlap model (INDO/S-CL). There is even no qualitative agreement between results of these calculations. Moreover, the ground state from the Ligand field calculations  $\Omega = 0.5$  while from INDO/S-CL calculation  $\Omega = 3.5$  (both states were derived from the same  $4f^{12}(3H)6s$  subconfiguration). However, for the calculation of the thermal function it does not change much: both states have the same statistical weight ( $pX = 2$ ). The most serious difference is in the arrangement of the states, mainly the first excited states. Kotzian *et al.*<sup>197</sup> calculated the first excited state at  $24 \text{ cm}^{-1}$  whereas the Ligand field calculation gave  $128$  or  $97 \text{ cm}^{-1}$ . In the present work we accept the data obtained by Kotzian *et al.*<sup>197</sup> and add the estimations of the states  $f^{12}d$  (according to Kotzian *et al.*<sup>197</sup> above  $13\,150 \text{ cm}^{-1}$ ),  $f^{12}p$  (higher  $20\,000 \text{ cm}^{-1}$ ), and  $f^{13}$  (higher  $40\,000 \text{ cm}^{-1}$ ).

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (255.041 \pm 0.15) \text{ JK}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{JK}^{-1} \text{ mol}^{-1}) = & 37.63499 + 4.06981 \times 10^{-3}(\text{T/K}) \\ & - 2.78859 \times 10^{-6}(\text{T/K})^2 + 0.967411 \\ & \times 10^{-9}(\text{T/K})^3 - 6.69507 \\ & \times 10^4(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1700 K range, and

$$\begin{aligned} C_p^\circ / (\text{JK}^{-1} \text{ mol}^{-1}) = & 39.61651 - 0.540323 \times 10^{-3}(\text{T/K}) \\ & + 1.26385 \times 10^{-6}(\text{T/K})^2 - 0.143823 \\ & \times 10^{-9}(\text{T/K})^3 - 1.21233 \\ & \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1700–4000 K range.

#### 4.14.2. Enthalpy of formation

The results for the enthalpy of formation of TmO(g) are presented in Table 53. Ames *et al.*<sup>206</sup> carried out Knudsen effusion measurement of weight loss for several Ln<sub>2</sub>O<sub>3</sub>(cr) oxides. As in the case of Sm<sub>2</sub>O<sub>3</sub>(cr) and other lanthanide oxides, the results of these measurements were treated under assumption of congruent vaporization of Tm<sub>2</sub>O<sub>3</sub>(cr) according to the reaction



neglecting possibility of formation of Tm(g) atoms in the vapor. In case of considerable degree of TmO dissociation this treatment will result in overestimated TmO stability.

Mass spectrometric measurements of isomolecular oxygen exchange reaction have been carried out by Murad and Hildenbrand.<sup>260</sup> Results of both papers seriously differ, and the enthalpy of formation of TmO(g) calculated from results of Ames *et al.* is about  $260 \text{ kJ mol}^{-1}$  more negative than that obtained from mass-spectrometric measurements by Murad and Hildenbrand. This difference can be explained by a large concentration of Tm atoms in the Tm<sub>2</sub>O<sub>3</sub>(cr) vapor in accordance with mass-spectrometric data for the TmO<sup>+</sup>/Tm<sup>+</sup> ratio.<sup>206</sup>

The selected value

$$\Delta_f H^\circ(\text{TmO, g, 298.15 K}) = -(13.6 \pm 8) \text{ kJ mol}^{-1}$$

is taken as a rounded third-law value from the work of Murad and Hildenbrand.<sup>260</sup> The selected enthalpy of formation corresponds to  $D_0(\text{TmO}) = (492.7 \pm 8) \text{ kJ mol}^{-1}$ .

## 4.15. YbO(g)

### 4.15.1. Heat capacity and entropy

The thermal functions of YbO(g) in the standard state have been calculated using the molecular constants presented in Table 54.

The electronic spectrum of YbO was investigated in emission by Melville *et al.*<sup>296</sup> and in laser excitation and fluorescence by McDonald *et al.*,<sup>297</sup> Linton *et al.*,<sup>227</sup> and Steimle *et al.*<sup>298</sup> The values of the fundamental frequency of YbO were observed in solid inert gas matrices by Willson and Andrews<sup>213</sup> in Ar at 10 K ( $660.0$  and  $625.8 \text{ cm}^{-1}$  for Yb<sup>16</sup>O and Yb<sup>18</sup>O, respectively), and by Willson *et al.*<sup>246</sup> in Ar ( $659.9 \text{ cm}^{-1}$ ). Taking into account the matrix shift, the values obtained in matrices are in agreement with the gas phase data.

*Ab initio* calculations for YbO carried out by Dolg and Stoll,<sup>193</sup> Dolg *et al.*,<sup>194</sup> and Cao *et al.*<sup>195</sup> revealed a marked disagreement with each other and the experimental data. The theoretical calculations performed by Liu *et al.*<sup>299</sup> favored a  $\Omega = 0^+$  ground state of a leading  $f^{14}\sigma 0$  configuration (in agreement with the interpretation of the experimental data) and predicted the 5 low-lying  $f^{13}s$  states.

The studies of the YbO electronic spectra revealed 7 low-lying states including the  $X^1\Sigma^+$  ( $f^{14}$ ) ground state, 5 of which were assigned to the  $f^{13}s$  and one to the  $f^{13}d$  configurations, and several excited states with energies  $16\,400$ – $24\,700 \text{ cm}^{-1}$ , assigned to the  $f^{13}d$  and  $f^{13}p$  configurations of YbO and to the  $f^{14}sp^5$  and  $f^{14}dp^5$  superconfigurations of Yb<sup>+</sup>O<sup>-</sup>. Ligand field calculations by McDonald *et al.*<sup>297</sup> and Carrete and Hocquet<sup>230</sup> resulted in all the  $f^{13}s$  states (the total statistical weight 28); Dulick *et al.*<sup>247</sup> obtained only the  $f^{13}s$  states up to  $10000 \text{ cm}^{-1}$

TABLE 53. The enthalpy of formation of TmO(g), in  $\text{kJ mol}^{-1}$

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2450-2641	$\text{Tm}_2\text{O}_3(\text{cr}) = 2\text{TmO}(\text{g}) + \text{O}(\text{g})$	1597.1	-270.4
Murad and Hildenbrand <sup>260</sup>	M	2249-2364	$\text{TmO}(\text{g}) + \text{Al}(\text{g}) = \text{Tm}(\text{g}) + \text{AlO}(\text{g})$	-16.3	-13.6
Selected value:			$D_0(\text{TmO}) = 492.7 \pm 8$		-13.6 ± 8

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry.

TABLE 54. Molecular constants of  $^{174}\text{Yb}^{16}\text{O}(\text{g})$ 

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$	$p_i$
		$\text{cm}^{-1}$					pm		
0 <sup>a</sup>	$X^1\Sigma^+$	0	689.9 <sup>b</sup>	3.49 <sup>b</sup>	0.352431	4.2	3.68 <sup>c</sup>	180.7	1
1 <sup>a</sup>	(2)0 <sup>-</sup>	839	832 <sup>d</sup>	3.4 <sup>d</sup>	0.355	1.9	2.6 <sup>d</sup>	180.1	1
2 <sup>a</sup>	(1)1	944							2
3 <sup>a</sup>	(1)2	2337							2
4 <sup>a</sup>	(2)2	2631							2
5 <sup>a</sup>	(1)3	4216							2
6 <sup>a</sup>	(3)0 <sup>+</sup>	4566							1
7 <sup>e</sup>		1557							1
8 <sup>e</sup>		5000							12
9 <sup>e</sup>		7500							16
10 <sup>e</sup>		10 000							24
11 <sup>e</sup>		15 000							40
12 <sup>e</sup>		20 000							50
13 <sup>e</sup>		25 000							55
14 <sup>e</sup>		30 000							44
15 <sup>e</sup>		35 000							35
16 <sup>e</sup>		40 000							35

<sup>a</sup>Experimental state.

<sup>b</sup>Calculated from  $\Delta G_{1/2} = 683.107 \text{ cm}^{-1}$  given by Linton *et al.*<sup>227</sup> and the assumed dissociation limit.

<sup>c</sup>Calculated from the Pekeris relation.

<sup>d</sup>Calculated from the Kratzer relation.

<sup>e</sup>Estimated state.

with  $\Sigma p = 17$ . The data obtained by McDonald *et al.*<sup>297</sup> are presented in Table 54; they were closer to the experimental data because of the use of adjustable parameters. In Table 54 also the estimated statistical weights for the  $f^{13}d$  states from the first observed ( $4637 \text{ cm}^{-1}$ ) are presented, as well as for the  $f^{13}p$  states (higher than  $19000 \text{ cm}^{-1}$ ), and for the  $\text{Yb}^+\text{O}^-$  states assuming the energies of the lowest state at  $15000 \text{ cm}^{-1}$ . The widths of configurations are estimated from the YbIII spectrum.

The accepted molecular constants for the  $X^1\Sigma^+$  state are derived from the high-resolution spectral data: the rotational constants from the data obtained by Melville *et al.*,<sup>296</sup> the vibrational constants ( $\Delta G_{1/2}$ ) from the data obtained by Linton *et al.*<sup>227</sup> The vibrational levels of the  $X^1\Sigma^+$  state were recorded in the latter work in the fluorescence spectrum up to  $v = 8$ , but with low accuracy ( $\sim 6 \text{ cm}^{-1}$ ). Moreover the ground state was perturbed in the region near  $v = 4$ . The molecular constants for low-lying states are also estimated from the low-resolution fluorescence spectral data. They are typical for the  $f^{N-1}s$  states of all lanthanide monoxides, and in the present work we use the average values. It should be noted that the ground state is the unique state of the  $f^{14}$  configuration. The constants for this state differ considerably from those for the other states. The low-lying  $f^{13}s$  states have practically identical potential curves

and therefore the same molecular constants ( $\Delta G_{1/2} \approx 825 \text{ cm}^{-1}$  and  $r_e \approx 180.0 \text{ pm}$ ). The  $X^1\Sigma^+$  state does not correlate with the Yb and O atoms in their ground states:  $^1S_g + ^3P_g$  give only triplet states  $^3\Pi$  and  $^3\Sigma^-$ . However, the components  $0^+$  of the  $^3\Pi$  and  $^3\Sigma^-$  states should affect the  $X^1\Sigma^+$  state, and because of the noncrossing rule the ground state should converge to the  $^1S_g + ^3P_g$  limit. The numerous  $f^{13}s$  states ( $\Sigma p = 28$ ) obviously have a higher dissociation limit. In the present work we assume the  $^3P_u + ^3P_g$  limit lying at  $17\,288 \text{ cm}^{-1}$  above the dissociation limit to normal atoms.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (238.521 \pm 0.10) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & 37.70801 + 48.5496 \times 10^{-3}(\text{T/K}) \\ & - 68.4668 \times 10^{-6}(\text{T/K})^2 + 30.5227 \\ & \times 10^{-9}(\text{T/K})^3 - 7.73176 \\ & \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1000 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & 32.77497 + 17.3649 \times 10^{-3}(\text{T/K}) \\ & - 5.08804 \times 10^{-6}(\text{T/K})^2 + 0.503717 \\ & \times 10^{-9}(\text{T/K})^3 + 1.98479 \\ & \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1000–4000 K range.

#### 4.15.2. Enthalpy of formation

There are no measurements of gas phase equilibria to derive the enthalpy of formation of  $\text{YbO}(\text{g})$ , due to very low concentration of  $\text{YbO}(\text{g})$  in the high-temperature systems. A lower limit estimate of the YbO dissociation energy (see Table 55) was obtained by Yokozeki and Menzinger<sup>300</sup> from the short wavelength chemiluminescence cutoffs in the  $\text{Yb} + \text{O}_3$  beam-gas experiment:  $D_0(\text{YbO}) > (394.1 \pm 6.3) \text{ kJ mol}^{-1}$ . The value  $D_0(\text{YbO}) = (413.8 \pm 4.8) \text{ kJ mol}^{-1}$  was found by crossing a thermal beam of Yb atoms with a supersonic seeded beam of  $\text{He} + \text{O}_2$  by Cosmovici *et al.*<sup>301</sup> As selected value we take the rounded result  $(414.0 \pm 10) \text{ kJ mol}^{-1}$  with increased uncertainty (Table 55), reflecting possible experimental errors in the work of Cosmovici *et al.*<sup>301</sup> The selected dissociation energy of YbO molecule corresponds to:

$$\Delta_f H^\circ(\text{YbO}, \text{g}, 298.15 \text{ K}) = -(16 \pm 10) \text{ kJ mol}^{-1}$$

 TABLE 55. The enthalpy of formation of  $\text{YbO}(\text{g})$ , in  $\text{kJ mol}^{-1}$ 

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Yokozeki and Menzinger <sup>300</sup>	C	<sup>b</sup>	$\text{Yb}(\text{g}) + \text{O}_3(\text{g}) = \text{YbO}(\text{g}) + \text{O}_2(\text{g})$	$> 394.1 \pm 6.3$	$< 3.9 \pm 6.3$
Cosmovici <i>et al.</i> <sup>301</sup>	MB	<sup>b</sup>	$\text{Yb}(\text{g}) + \text{O}_2(\text{g}) = \text{YbO}(\text{g}) + \text{O}(\text{g})$	$413.8 \pm 4.8$	$-16.2 \pm 10$
Selected value:			$D_0(\text{YbO}) = 414 \pm 10$		$-16 \pm 10$

<sup>a</sup>C = chemiluminescence; MB = crossed molecular beam.

<sup>b</sup>Not specified.



TABLE 56. Molecular constants of  $^{175}\text{Lu}^{16}\text{O}(\text{g})$ 

No	State	$\text{cm}^{-1}$					$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$		
0 <sup>a</sup>	$X^2\Sigma^+$	0 844.5	3.1	0.35894	1.76	2.78 <sup>b</sup>	179.0	2
1 <sup>a</sup>	$A^2\Pi_{1/2}$	19 392						2
2 <sup>a</sup>	$A^2\Pi_{3/2}$	21 470						2
3 <sup>a</sup>	$4^1\Pi ?^c$	19 520						8
4 <sup>a</sup>	$B^2\Sigma^+$	24 440						2
5 <sup>d</sup>		14 100						2
6 <sup>d</sup>		15 300						2
7 <sup>d</sup>		21 000						8
8 <sup>d</sup>		23 700						8
9 <sup>d</sup>		27 000						12
10 <sup>d</sup>		31 000						4
11 <sup>d</sup>		40 000						20

<sup>a</sup>Experimental state.<sup>b</sup> $D_0$ .<sup>c</sup>Unknown upper state in of the transition band at 5120 Å.<sup>d</sup>Estimated state.

## 4.16. LuO(g)

### 4.16.1. Heat capacity and entropy

The thermal functions of LuO(g) in the standard state have been calculated using the molecular constants presented in Table 56.

Three electronic transitions  $A^2\Pi_{1/2}-X^2\Sigma^+$ ,  $A^2\Pi_{3/2}-X^2\Sigma^+$ , and  $B^2\Sigma^+-X^2\Sigma^+$  were analyzed in the emission spectrum of LuO. Huber and Herzberg<sup>179</sup> reviewed the spectral data on LuO published till 1975. Later Bernard and Effantin<sup>302</sup> reanalyzed the known systems and found a new band at 5120 Å, which also was connected with the transition to the ground state. These are slightly different from values derived from the recent measurements of the pure rotational spectrum of  $^{175}\text{Lu}^{16}\text{O}$  in its ground electronic state ( $X^2\Sigma^+$ ) by Cooke *et al.*,<sup>303</sup> which were published after our analysis was completed. All the upper states of the transitions were perturbed that indicates many unobserved states. The values of the fundamental frequency of LuO were measured in solid inert gas matrices by Willson and Andrews<sup>213</sup> in Ar (829.3  $\text{cm}^{-1}$  and 786.2  $\text{cm}^{-1}$  for Lu<sup>16</sup>O and Lu<sup>18</sup>O, respectively) and by Weltner and De Kock<sup>245</sup> in Ar and Ne (825 and 836  $\text{cm}^{-1}$ , respectively). Taking into account the matrix shift, the values of the fundamental frequency of LuO obtained in matrices are in agreement with the gas phase data.

*Ab initio* calculations for the ground state of LuO carried out by Hong *et al.*,<sup>192</sup> Wang and Schwarz,<sup>304</sup> Kühle *et al.*,<sup>305</sup> Dolg and Stoll,<sup>193</sup> Dolg *et al.*,<sup>194</sup> and Cao *et al.*,<sup>195</sup> are in agreement with the experimental data. Kotzian *et al.*<sup>197</sup> calculated the excited states using the INDO model. According to this calculation, all the states below 45 000  $\text{cm}^{-1}$  arise from

superconfigurations  $5d(A^2\Delta, A^2\Pi, \text{ and } B^2\Sigma^+)$ ,  $6p(^2\Pi \text{ and } ^2\Sigma^+)$ , and  $6s(X^2\Sigma^+)$ , while the  $4f^{13}$  states have energies much higher (100 000  $\text{cm}^{-1}$ ). These results could not explain either the upper state of the band 5120 Å, or numerous perturbations. The electronic structures of LaO and LuO are similar. In the present work the upper state of the band 5120 Å is interpreted as the lowest component of quartet states, found by Schampsand *et al.*<sup>198</sup> for LaO. The estimates of the excited states are based on the above mentioned data.

The accepted molecular constants for the  $X^2\Sigma^+$  state are taken from the work by Bernard and Effantin.<sup>302</sup>

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (242.089 \pm 0.03) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 28.20132 + 20.4184 \times 10^{-3}(\text{T/K}) \\ &\quad - 16.5159 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 4.71909 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 1.19650 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1300 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 40.19679 - 0.758989 \times 10^{-3}(\text{T/K}) \\ &\quad - 0.623484 \times 10^{-6}(\text{T/K})^2 + 0.273798 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 2.75040 \\ &\quad \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1300–4000 K range.

### 4.16.2. Enthalpy of formation

The results for the enthalpy of formation of LuO(g) are presented in Table 57. Experimental data used in calculations consist of Knudsen effusion<sup>206</sup> and mass spectrometric measurements of isomolecular oxygen exchange reactions.<sup>206,260</sup> The results of all measurements are in fair agreement. At the same time, mass-spectrometric results are more reliable in comparison with the KE data, because interpretation of Knudsen effusion data is not unambiguous (see Sec. 4.1). Among the mass-spectrometric data, the results of Murad and Hildenbrand<sup>260</sup> are preferred. Absence of primary experimental data makes it difficult to recalculate results of mass-spectrometric measurements of Ames *et al.*<sup>206</sup>

The selected enthalpy of formation of LuO(g) is taken as the rounded value from the paper by Murad and Hildenbrand<sup>260</sup>:

$$\Delta_f H^\circ(\text{LuO, g, } 298.15 \text{ K}) = -(3.4 \pm 10) \text{ kJ mol}^{-1}.$$

TABLE 57. The enthalpy of formation of LuO(g), in  $\text{kJ mol}^{-1}$ 

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ames <i>et al.</i> <sup>206</sup>	K	2615–2700	$\text{Lu}_2\text{O}_3(\text{cr}) = 2\text{LuO}(\text{g}) + \text{O}(\text{g})$	2150.4	11.5
	M	2080–2214	$\text{Lu}(\text{g}) + \text{LaO}(\text{g}) = \text{LuO}(\text{g}) + \text{La}(\text{g})$	117.3	–5.8
Murad and Hildenbrand <sup>260</sup>	M	1782–1927	$\text{Lu}(\text{g}) + \text{YO}(\text{g}) = \text{LuO}(\text{g}) + \text{Y}(\text{g})$	45.1	3.4
Selected value:			$D_0(\text{LuO}) = 758.9 \pm 10$		3.4 ± 10

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry.

The selected enthalpy of formation corresponds to  $D_0(\text{LuO}) = (673.6 \pm 10) \text{ kJ mol}^{-1}$ .

## 5. The Actinide Oxides in Solid and Liquid State

### 5.1. $\text{Ac}_2\text{O}_3(\text{cr,l})$

#### 5.1.1. Polymorphism and melting point

Actinium sesquioxide has an A-type hexagonal sesquioxide structure (space group  $P\bar{3}m1$ ) at room temperature. Similar to the isostructural A-type lanthanide sesquioxides, high temperature transformations to the H- and X-type structures can be expected, but no experimental information exists. Considering the fact that  $\text{Ac}^{3+}$  has an ionic radius somewhat larger than  $\text{La}^{3+}$ , the transition temperatures are estimated to be slightly lower than those in  $\text{La}_2\text{O}_3$ , i.e.,  $A \rightarrow H$  at about 2300 K and  $H \rightarrow X$  around 2370 K. Similarly, the melting point of  $\text{Ac}_2\text{O}_3$  is estimated to be about 2600 K.

#### 5.1.2. Heat capacity and entropy

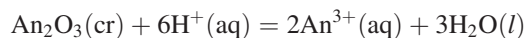
The standard entropy of  $\text{Ac}_2\text{O}_3$  has been estimated from the trends in the lanthanide and actinide compounds as

$$S^\circ(298.15 \text{ K}) = (138.3 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

according to the method proposed by Konings *et al.*<sup>127</sup> This is slightly lower than the value suggested by Konings *et al.*<sup>306</sup> which did not take into account the structural discontinuity in the  $\text{Ln}_2\text{O}_3$  series, as evidenced by the data for monoclinic  $\text{Gd}_2\text{O}_3$ .

#### 5.1.3. Enthalpy of formation

The enthalpy of formation of  $\text{Ac}_2\text{O}_3$  was estimated by Konings *et al.*<sup>306</sup> from the enthalpy of the idealised dissolution reaction:



relating the quantity  $\{\Delta_f H^\circ(\text{MO}_{1.5}) - \Delta_f H^\circ(\text{M}^{3+})\}$  to the molar volume. They thus obtained

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1107 \pm 15) \text{ kJ mol}^{-1}$$

## 5.2. $\text{ThO}_2(\text{cr,l})$

### 5.2.1. Melting point

$\text{ThO}_2$  has a fluorite crystal structure (space group  $Fm\bar{3}m$ ), which is stable up to the melting point. The melting point of  $\text{ThO}_2$  has been measured by several authors, as summarized in Table 58. The reported values vary from  $T = 3323 \text{ K}$  to  $T = 3808 \text{ K}$ , but the more recent ones agree on a melting point around  $T = 3600 \text{ K}$ . We here consider the value measured by Ronchi and Hiernaut,<sup>307</sup>  $T_{\text{fus}} = (3651 \pm 17) \text{ K}$ , as the most

TABLE 58. Temperature of melting of thorium dioxide

Authors	$T_{\text{fus}}/\text{K}$	
	Reported	ITS-90
Ruff <i>et al.</i> <sup>308</sup>	3323	
Wartenberg and Reusch <sup>309</sup>	3803	
Geach and Harper <sup>310</sup>	$3323 \pm 25$	$3328 \pm 25$
Lambertson <i>et al.</i> <sup>311</sup>	$3573 \pm 100$	$3578 \pm 100$
Benz <sup>312</sup>	$3663 \pm 100$	$3668 \pm 100$
Chikalla <i>et al.</i> <sup>107</sup>	3573	3572
Sibieude and Foex <sup>313</sup>	3473	3472
Ronchi and Hiernaut <sup>307</sup>	$3651 \pm 17$	$3651 \pm 17$
Manara <i>et al.</i> <sup>52</sup>	3620	3620
Selected value:		$3651 \pm 17$

accurate one as it was determined on a well-defined sample ( $O/\text{Th}$  ratio = 2.00) and with a well-defined technique.

### 5.2.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{ThO}_2$  has been measured by Osborne and Westrum Jr.<sup>314</sup> These measurements were used in the CODATA Key Values selection,<sup>315</sup> and selected here without change:

$$S^\circ(298.15 \text{ K}) = (65.23 \pm 0.20) \text{ J K}^{-1} \text{ mol}^{-1}.$$

Magnani *et al.*<sup>316</sup> reported heat capacity data that are in good agreement, but the numerical details of this measurement are not published and the analytical technique is less accurate.

The high-temperature enthalpy increment of  $\text{ThO}_2$  has been measured by Jaeger and Veenstra,<sup>317</sup> Southard,<sup>318</sup> Hoch and Johnston,<sup>319</sup> Victor and Douglas,<sup>320</sup> Pears *et al.*,<sup>58</sup> Springer *et al.*,<sup>321</sup> Springer and Langedrost,<sup>322</sup> Fischer *et al.*,<sup>323,324</sup> and Dash *et al.*<sup>325</sup> The data cover the temperature range from 500 to 3400 K as shown in Fig. 15. Up to 2500 K the results are in fair agreement, except in the low-temperature region where the data of Victor and Douglas<sup>320</sup> and Springer *et al.*<sup>321</sup> deviate significantly due to small inaccuracies in temperature or enthalpy that become prominent close to 298.15 K when using the  $\{H^\circ(T) - H^\circ(298.15 \text{ K})\}/(T - 298.15)$  function. The data listed by Springer *et al.* have been corrected for obvious typographical errors. The results of Pears *et al.*<sup>58</sup> are very scattered and are evidently not accurate enough. Direct heat

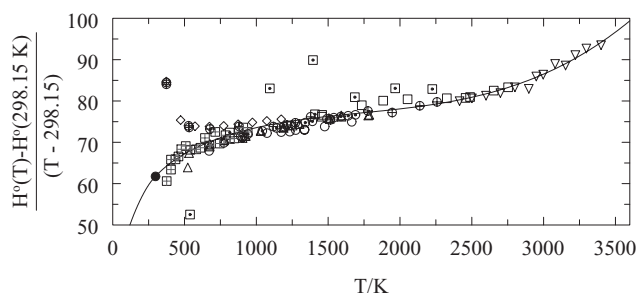


FIG. 15. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{ThO}_2$ ;  $\circ$ , Jaeger and Veenstra<sup>317</sup>;  $\triangle$ , Southard<sup>318</sup>;  $\square$ , Hoch and Johnston<sup>319</sup>;  $\square$ , Pears *et al.*<sup>58</sup>;  $\diamond$ , Victor and Douglas<sup>320</sup>;  $\oplus$ , Springer *et al.*<sup>321</sup>;  $\nabla$ , Fischer *et al.*<sup>323</sup>;  $\boxplus$ , Agarwal *et al.*<sup>324</sup>;  $\odot$ , Dash *et al.*<sup>325</sup>;  $\bullet$ , Osborne and Westrum Jr.<sup>314</sup>; the curve shows the recommended equation.

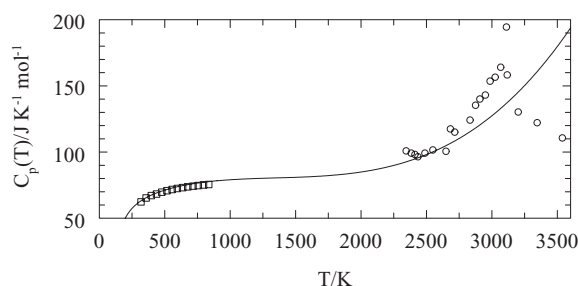


FIG. 16. The heat capacity (in  $\text{J K}^{-1} \text{mol}^{-1}$ ) of  $\text{ThO}_2$ ;  $\circ$ , Ronchi and Hiernaut<sup>307</sup>;  $\square$  Dash *et al.*<sup>325</sup>; the curve shows the recommended equation. Note that the data of Ronchi and Hiernaut<sup>307</sup> indicate a value of about  $600 \text{ J K}^{-1} \text{mol}^{-1}$  (not shown in the graph) at the maximum of the anomalie.

capacity measurements of  $\text{ThO}_2$  have been reported by Dash *et al.*,<sup>325</sup> in good agreement with the enthalpy measurements (Fig. 16).

Above  $T = 2500 \text{ K}$ ,  $\text{ThO}_2$  exhibits an excess enthalpy, like many high-melting refractory oxides (e.g.,  $\text{UO}_2$ ,  $\text{PuO}_2$ , and  $\text{ZrO}_2$ ). Fischer *et al.*<sup>323</sup> suggested that this effect is due to a phase transformation and this possibility was studied in detail by Ronchi and Hiernaut<sup>307</sup> using a thermal arrest technique. From their results, Ronchi and Hiernaut concluded that a premelting transition occurs at  $3090 \text{ K}$ , which was attributed to order-disorder anion displacements in the oxygen sublattice (Frenkel defects).

For the recommended heat capacity equation the results of Southard,<sup>318</sup> Hoch and Johnston<sup>319</sup> and Fischer *et al.*<sup>323</sup> have been combined and fitted to the equation ( $298.15\text{--}3500 \text{ K}$ ):

$$C_p^\circ / (\text{J K}^{-1} \text{mol}^{-1}) = 55.9620 + 51.2579 \times 10^{-3} (T/\text{K}) \\ - 36.8022 \times 10^{-6} (T/\text{K})^2 \\ + 9.2245 \times 10^{-9} (T/\text{K})^3 \\ - 5.740310 \times 10^5 (T/\text{K})^{-2}$$

constrained to  $C_p^\circ(298.15 \text{ K}) = 61.76 \text{ J K}^{-1} \text{mol}^{-1}$ , as derived from the low-temperature heat capacity measurements by Osborne and Westrum Jr.<sup>314</sup> The  $C_p^\circ$  derived from this function, of course, does not reproduce the heat capacity data by Ronchi and Hiernaut<sup>307</sup> around the order-disorder transition (see Fig. 16), with a maximum value of  $600 \text{ J K}^{-1} \text{mol}^{-1}$  (not shown in the figure).

No experimental data are available for liquid thorium dioxide. Fink *et al.*<sup>326</sup> estimated  $C_p^\circ(\text{liq}) = 61.76 \text{ J K}^{-1} \text{mol}^{-1}$ , which is adopted here as

$$C_p^\circ(\text{ThO}_2, \text{liq}, T) = 61.8 \text{ J K}^{-1} \text{mol}^{-1}.$$

The entropy of fusion is assumed to be identical to that of  $\text{UO}_2$  ( $24 \text{ J K}^{-1} \text{mol}^{-1}$ ), yielding

$$\Delta_{\text{fus}} H^\circ = (88 \pm 6) \text{ kJ mol}^{-1}.$$

### 5.2.3. Enthalpy of formation

The enthalpy of formation of  $\text{ThO}_2$  is a CODATA Key Value for Thermodynamics,<sup>315</sup> and is based on the enthalpy of

combustion of thorium metal by Huber Jr. *et al.*:<sup>327</sup>

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1226.4 \pm 3.5) \text{ kJ mol}^{-1}.$$

Earlier measurements by Roth and Becker<sup>328</sup> gave  $-(1226 \pm 5) \text{ kJ mol}^{-1}$ , which is in good agreement with the selected value.

## 5.3. $\text{PaO}_2(\text{cr,l})$

### 5.3.1. Melting point

$\text{PaO}_2$  has a fluorite crystal structure (space group  $Fm\bar{3}m$ ). No information exists about the high temperature behavior of  $\text{PaO}_2$ , but it can be presumed that it is similar to the neighboring  $\text{ThO}_2$  and  $\text{UO}_2$  compounds, i.e. the fluorite structure is stable up to the melting. The melting point of  $\text{PaO}_2$  is estimated from the trend in the actinide dioxide series, as discussed in Sec. 7.2.2, and we estimate  $T_{\text{fus}} = (3200 \pm 60) \text{ K}$ .

### 5.3.2. Heat capacity and entropy

The standard entropy of  $\text{PaO}_2$  was estimated by Konings<sup>329</sup> from the trends in actinide dioxide series:

$$S^\circ(298.15 \text{ K}) = (81.1 \pm 5.0) \text{ J K}^{-1} \text{mol}^{-1}.$$

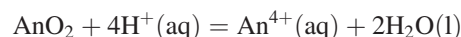
The high temperature heat capacity of  $\text{PaO}_2$  has been estimated as

$$C_p^\circ(T) / (\text{J K}^{-1} \text{mol}^{-1}) = 58.0078 + 50.9087 \times 10^{-3} (T/\text{K}) \\ - 35.9277 \times 10^{-6} (T/\text{K})^2 + 9.5704 \\ \times 10^{-9} (T/\text{K})^3 - 0.51080 \\ \times 10^6 (T/\text{K})^{-2}$$

using the approach outlined in Konings and Benes<sup>330</sup>.

### 5.3.3. Enthalpy of formation

The enthalpy of formation of  $\text{PaO}_2$  was estimated by Konings *et al.*<sup>306</sup> from the enthalpy of the idealised dissolution reaction:



relating this quantity to the molar volume. They thus obtained

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1107 \pm 15) \text{ kJ mol}^{-1}.$$

## 5.4. $\gamma\text{-UO}_3$

### 5.4.1. Polymorphism

$\gamma\text{-UO}_3$  is one of the many crystallographic modifications of  $\text{UO}_3$ , but probably the thermodynamic stable one above room temperature.<sup>331,332</sup> It has an orthorhombic cell (space group  $Fddd$ ) at  $298.15 \text{ K}$ . It transforms at  $\sim 373 \text{ K}$  to a closely related tetragonal structure (space group  $I4_1/amd$ ).

### 5.4.2. Heat capacity and entropy

Low-temperature heat capacity measurements of  $\text{UO}_3$  have been reported by Jones *et al.*<sup>333</sup> from 15 to 300 K. Based on the sample preparation and the color of the product, it is generally believed that it had the  $\gamma$  structure. Cordfunke and Westrum Jr.<sup>334</sup> measured the low-temperature heat capacity of a well-characterised  $\gamma$ - $\text{UO}_3$  sample from 5 to 350 K, yielding somewhat lower values. The selected entropy value is taken from that study

$$S^\circ(298.15 \text{ K}) = (96.11 \pm 0.40) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high temperature heat capacity of  $\text{UO}_3$  of unknown crystallographic structure has been measured by Popov *et al.*<sup>335</sup> from 392 to 673 K. The high-temperature enthalpy increment of  $\text{UO}_3$  has been measured by Moore and Kelley<sup>336</sup> from 416 to 886 K on a sample of unspecified crystallographic structure, but since it was identical to that used by Jones *et al.*<sup>333</sup> for the low-temperature measurements, the results most likely refer to  $\gamma$ - $\text{UO}_3$ . Cordfunke and Westrum Jr.<sup>334</sup> measured the high-temperature enthalpy increment from 347 to 691 K on a well-defined sample having the  $\gamma$  structure. These measurements do not reveal the phase transition at  $\sim 373$  K, which means that the entropy of transition is negligible.

The results of the two enthalpy studies agree well, and although the latter study is made on a better characterised sample, our recommended heat capacity equation is based on a fit of all results, since the study by Moore and Kelley<sup>336</sup> covers a wider temperature range. The enthalpy fit is constrained to  $C_p^\circ(298.15 \text{ K}) = 81.67 \text{ J K}^{-1} \text{ mol}^{-1}$  from the low-temperature measurements and yields for the heat capacity:

$$C_p / (\text{J K}^{-1} \text{ mol}^{-1}) = 90.2284 + 13.85332 \times 10^{-3} (T/\text{K}) - 1.12795 \times 10^6 (T/\text{K})^{-2}.$$

### 5.4.3. Enthalpy of formation

The enthalpy of formation of  $\gamma$ - $\text{UO}_3$  is a CODATA Key Value for Thermodynamics,<sup>315</sup> and is based on the enthalpies of dissolution of uranium oxides in Ce(IV) solutions corrected to stoichiometric  $\text{UO}_3$  by Fitzgibbon *et al.*,<sup>337</sup> and on the dissolution of  $\gamma$ - $\text{UO}_3$  and  $\text{UF}_6(\text{cr})$  in HF solutions by Johnson and O'Hare.<sup>338</sup> The value is in agreement with the decomposition pressures measured by Cordfunke and Aling<sup>332</sup>:

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1223.8 \pm 2.0) \text{ kJ mol}^{-1}.$$

## 5.5. $\text{U}_3\text{O}_8(\text{cr})$

### 5.5.1. Polymorphism and melting point

At room temperature  $\alpha$ - $\text{U}_3\text{O}_8(\text{cr})$  has an orthorhombic structure (space group  $C2mm$ ). It transforms to a hexagonal structure (space group  $P\bar{6}2m$ ) at 483 K. This transition is revealed as a clear  $\lambda$  peak in the heat capacity studies.<sup>339,340</sup> They indicated two additional peaks at 568 and 850 K (see

below), whose origin is not established but presumably involves further changes in ordering in the lattice.

At atmospheric pressure  $\text{U}_3\text{O}_8$  decomposes before melting. The melting temperature of  $\text{U}_3\text{O}_8$  was determined by Manara *et al.*<sup>341</sup> as  $(2010 \pm 30)$  K under a pressure of 1 kbar pure helium.

### 5.5.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{U}_3\text{O}_8$  has been measured by Westrum Jr. and Grønvold<sup>342</sup> from 5 to 350 K, which revealed a  $\lambda$  type transition at 25.3 K, probably of magnetic origin. The standard entropy derived from this work, also accepted as CODATA Key Value for Thermodynamics,<sup>315</sup> is

$$S^\circ(298.15 \text{ K}) = (282.55 \pm 0.50) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature heat capacity of  $\text{U}_3\text{O}_8$  has been measured by Popov *et al.*<sup>343</sup> from 350 to 875 K, Girdhar and Westrum Jr.<sup>339</sup> from 303 to 529 K, and Inaba *et al.*<sup>340</sup> from 310 to 970 K. The high-temperature enthalpy increment has been measured by Maglic and Herak<sup>344</sup> from 312 to 927 K, Marchidan and Ciopec<sup>345</sup> from 273 to 1000 K, and Cordfunke whose results have not been published (cited by Cordfunke and Konings<sup>8</sup>). The heat capacity studies by Girdhar and Westrum Jr.<sup>339</sup> and Inaba *et al.*<sup>340</sup> reveal a  $\lambda$  peak at 483 K. The latter authors found two further  $\lambda$  peaks at 568 and 850 K. The enthalpies of the transitions of the peak at 483 K derived from these studies are, however, very different. Girdhar and Westrum Jr.<sup>339</sup> derive  $\Delta_{trs} H^\circ = 171 \text{ J mol}^{-1}$ , Inaba *et al.*<sup>340</sup>  $\Delta_{trs} H^\circ = 405 \text{ J mol}^{-1}$ . As can be seen in Fig. 17, the difference mainly arises from the fact that the peak is much broader in the measurements of Inaba *et al.*<sup>340</sup>. We consider the measurement of Girdhar and Westrum Jr.<sup>339</sup> more precise and select the value derived from that work. For the transitions at 568 and 850 K we have taken the values from the work of Inaba *et al.*,<sup>340</sup> although they may be somewhat too high:

$$\Delta_{trs} H^\circ(483\text{K}) = 171 \text{ J mol}^{-1},$$

$$\Delta_{trs} H^\circ(568\text{K}) = 444 \text{ J mol}^{-1},$$

$$\Delta_{trs} H^\circ(850\text{K}) = 942 \text{ J mol}^{-1}.$$

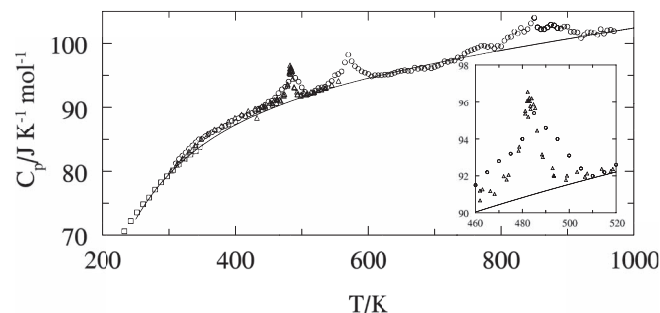


FIG. 17. The heat capacity of  $\text{UO}_{2.667}$ ;  $\square$ , Inaba *et al.*<sup>340</sup>;  $\square$ , Westrum Jr. and Grønvold<sup>342</sup>;  $\triangle$ , Girdhar and Westrum Jr.<sup>339</sup>



The transitions are not detected in the enthalpy drop studies since these enthalpy effects are too small.

The experimental studies indicate that the baseline heat capacity of the three phases can be represented by a single curve as a function of temperature. The selected heat capacity curve is taken from Cordfunke and Konings<sup>8</sup> which includes unpublished results by Cordfunke. It is subject to the constraint of  $C_p^\circ(298.15\text{ K}) = 237.94\text{ J K}^{-1}\text{ mol}^{-1}$ , as given by Westrum Jr. and Grønvold<sup>342</sup>:

$$C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) = 279.267 + 27.480 \times 10^{-3}(T/\text{K}) - 4.3116 \times 10^6(T/\text{K})^2.$$

It is recommended to use this equation in combination with the transition enthalpies.

### 5.5.3. Enthalpy of formation

The enthalpy of formation of  $\text{U}_3\text{O}_8$  is a CODATA Key Value for Thermodynamics,<sup>315</sup> and is based on the enthalpy of combustion of uranium measured by Huber and Holley Jr.<sup>346</sup>:

$$\Delta_f H^\circ(298.15\text{ K}) = -(3574.8 \pm 2.5)\text{ kJ mol}^{-1}.$$

This value is in close agreement with, but more precise than earlier values by Huber Jr. and Holley Jr.<sup>98</sup> and Popov and Ivanov.<sup>347</sup>

## 5.6. $\text{U}_4\text{O}_9(\text{cr})$

### 5.6.1. Polymorphism

$\text{U}_4\text{O}_9$ , which has a stability range between  $2.234 < \text{O/U} < 2.245$ , has three polymorphic modifications. The  $\alpha$  form has probably a rhombohedrally distorted fluorite structure. At 348 K it transforms into the  $\beta$  modification which has a body-centered cubic (space group  $I\bar{4}3d$ ). This temperature is the maximum of the  $\lambda$  peak in the heat capacity measured by Westrum Jr. *et al.*<sup>348</sup> and Grønvold *et al.*<sup>349</sup> Inaba and Naito<sup>350</sup> and Naito *et al.*<sup>351</sup> found that the temperature of this transition slightly increases with decreasing O/U ratio of the sample.

The  $\beta$  modification transforms to  $\gamma\text{-U}_4\text{O}_9$  at about 893 K, as derived from electrical conductivity and X-ray diffraction measurements by Inaba and Naito.<sup>350</sup> The structure of this phase is not known. A disordered  $\text{UO}_{2.25}$  phase is stable above 1400 K.<sup>352</sup> Essentially this may be regarded as an order-disorder transformation at a fixed composition, although strictly it is peritectoid decomposition from  $\text{U}_4\text{O}_9$  (ordered) to  $\text{UO}_{2+x}$  (disordered,  $x \sim 0.25$ ) and a small amount of  $\text{U}_3\text{O}_{8-y}$ .

### 5.6.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{U}_4\text{O}_9$  has been measured by Osborne *et al.*<sup>353</sup> from 5 to 310 K, by Westrum Jr. *et al.*<sup>348</sup> from 190 to 399 K, and Flotow *et al.*<sup>354</sup> from 1.6 to 24 K. The results are in good agreement. The standard entropy

derived from these measurements is<sup>354</sup>

$$S^\circ(298.15\text{ K}) = (334.1 \pm 0.7)\text{ J K}^{-1}\text{ mol}^{-1}.$$

In addition to the results of Westrum Jr. *et al.*,<sup>348</sup> which extend to 399 K, three more high-temperature heat capacity studies have been reported. Gotoo and Naito<sup>355</sup> reported measurements from 297 to 515 K, Grønvold *et al.*<sup>349</sup> from 303 to 997 K, and Inaba and Naito<sup>350</sup> from 190 to 470 K. MacLeod<sup>352</sup> measured the enthalpy increments of  $\text{U}_4\text{O}_9$  from 845 to 1568 K. The heat capacity studies show that the  $\alpha \rightarrow \beta$  transition is associated with a strong  $\lambda$ -type peak. The  $\beta \rightarrow \gamma$  transition is only evident from a spread in the  $C_p^\circ$  results of Grønvold *et al.*<sup>349</sup> We have selected the heat capacity equation given by Cordfunke and Konings,<sup>8</sup> which describes the three phases with one polynomial, treating the  $\lambda$  transition as an enthalpy discontinuity only

$$C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) = 319.163 + 49.691 \times 10^{-3}(T/\text{K}) - 3.9602 \times 10^6(T/\text{K})^{-2}.$$

MacLeod<sup>352</sup> has analysed the integrated enthalpy for the  $\alpha \rightarrow \beta$  transition as  $\Delta_{irs} H^\circ(\alpha-\beta) = 2594\text{ J}\cdot\text{mol}^{-1}$ . The  $\beta$  to  $\gamma$  transition has been assumed to have a negligible associated enthalpy. Although MacLeod analysed his data to suggest  $\Delta_{irs} H^\circ(1395\text{ to }1405\text{ K}) = 9.372\text{ kJ mol}^{-1}$ , this calculation seems to be in error - the enthalpy difference between the disordered  $\text{UO}_{2+x}$  and the  $\gamma$ -phase at 1400 K, from Eqs. (5) and (6) is in fact  $11.90\text{ kJ mol}^{-1}$ .

### 5.6.3. Enthalpy of formation

The selected enthalpy of formation of  $\text{U}_4\text{O}_9$  is

$$\Delta_f H^\circ(298.15\text{ K}) = -(4512 \pm 7)\text{ kJ mol}^{-1}$$

based on the enthalpies of solution of  $\text{UO}_2$ ,  $\text{U}_4\text{O}_9$  and  $\gamma\text{-UO}_3$  in aqueous Ce(IV) solutions measured by Fitzgibbon *et al.*<sup>337</sup> This value is supported by the less precise work of Burdese and Abbatista<sup>356</sup> using dissolution in nitric acid.

## 5.7. $\text{UO}_2(\text{cr,l})$

### 5.7.1. Melting point

$\text{UO}_2$  has a fluorite crystal structure (space group  $Fm\bar{3}m$ ), which is stable up to the melting point. The reported melting temperatures are listed in Table 59, which shows considerable variation. This is due to the fact that deviations from stoichiometry and the relatively high vapor pressure have significant effects, in addition to interactions of liquid with the container material. We select the value  $T_{fus} = (3130 \pm 20)\text{ K}$ . The exact congruent melting composition at atmospheric pressure is still controversial. Most probably, it lies between  $\text{UO}_{1.98}$  and  $\text{UO}_{2.00}$ .<sup>357,358</sup>

TABLE 59. Temperature of melting of uranium dioxide

Authors	$T_{\text{fus}}/\text{K}$	
	Reported	ITS-90
Ackermann <sup>359</sup>	2680 ± 19	
Lambertson and Handwerk <sup>360</sup>	3323 ± 20	
Wisnyi and Pijanowski <sup>361</sup>	3033 ± 30	
Ehlert and Margrave <sup>362</sup>	3133 ± 45	
Pijanowski and DeLuca <sup>363</sup>	3033 ± 30	3050 ± 30
Chikalla <sup>364</sup>	3003 ± 30	
Lyon and Bailey <sup>365</sup>	3046 ± 21	
Hausner <sup>366</sup>	3078 ± 15	
Lyon and Bailey <sup>367</sup>	3113 ± 20	
Latta and Fryxell <sup>368</sup>	3138 ± 15	3142 ± 15
Tachibana <i>et al.</i> <sup>369</sup>	3118 ± 25	3120 ± 25
Ronchi and Sheindlin <sup>370</sup>	3110 ± 10	3110 ± 10
Manara <i>et al.</i> <sup>371</sup>	3147 ± 20	3147 ± 20
Kato <i>et al.</i> <sup>372</sup>	3123	3123
Selected value:		3130 ± 20

### 5.7.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{UO}_2$  has been measured by Jones *et al.*<sup>333</sup> from 15 to 300 K and Hunzicker and Westrum Jr.<sup>373</sup> from 5 to 330 K. These measurements reveal a transition at 30.4 K from a low-temperature antiferromagnetic state to a high-temperature paramagnetic state. The selected standard entropy is the CODATA Key Value,<sup>315</sup> which is solely based on the results of Hunzicker and Westrum Jr.<sup>373</sup> that refer to a better characterised sample:

$$S^\circ(298.15 \text{ K}) = (77.03 \pm 0.20) \text{ J K}^{-1} \text{ mol}^{-1}.$$

The high-temperature data for  $\text{UO}_2$  extend into the liquid range (up to 8000 K) and have been obtained as heat capacity<sup>335,349,379–387</sup> and enthalpy increment values.<sup>336,374–378,388–390</sup> The results are shown in Figs. 18 and 19. There is in general good agreement between the different studies, with exception of the early heat capacity measurements.<sup>335,383–385</sup>

It is now well established that the heat capacity of  $\text{UO}_2$  shows an anomalous increase above 1800 K. Neutron scattering measurements by Hutchings<sup>391,392</sup> and Clausen *et al.*<sup>393</sup> reveal that thermally induced disorder as a result of Frenkel pair formation on the oxygen lattice occurs at temperatures above 2000 K. These studies also showed that excitation of the electronic levels of  $\text{U}^{4+}$

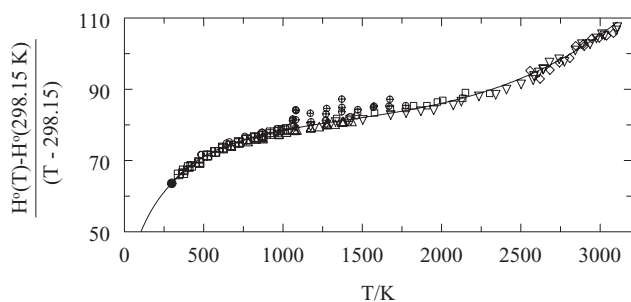


FIG. 18. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{UO}_2$ ;  $\circ$ , Moore and Kelley<sup>336</sup>;  $\square$ , Ogard and Leary<sup>374</sup>;  $\triangle$ , Fredrickson and Chasanov<sup>375</sup>;  $\nabla$ , Hein and Flagella<sup>376</sup>;  $\diamond$ , Leibowitz *et al.*<sup>377</sup>;  $\boxplus$ ,<sup>378</sup>;  $\oplus$ , Mills *et al.*<sup>379</sup>;  $\bullet$ , value derived from the low-temperature measurements by Hunzicker and Westrum Jr.<sup>373</sup>; the curve shows the recommended equation.

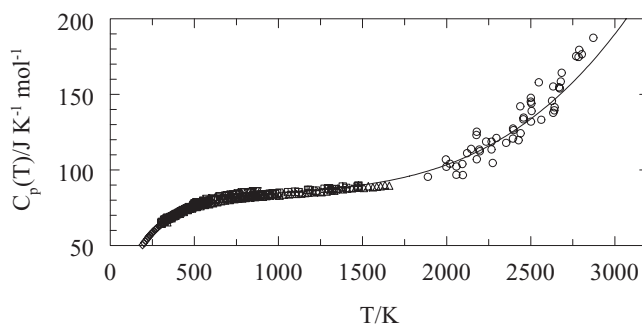


FIG. 19. The heat capacity of  $\text{UO}_2$ ;  $\circ$ , Ronchi *et al.*<sup>380</sup>;  $\square$ , Grønvdal *et al.*<sup>349</sup>;  $\triangle$ , Amaya *et al.*<sup>381</sup>;  $\nabla$ , Popov *et al.*<sup>335</sup>;  $\diamond$ , Hunzicker and Westrum Jr.<sup>373</sup>;  $\boxplus$ , Inaba *et al.*<sup>382</sup>; the curve shows the recommended equation.

contribute to the anomalous behavior. According to Ronchi and Hyland<sup>394</sup> the former dominate. The heat capacity measurements in the pre-melting range by Hiernaut *et al.*<sup>386</sup> showed a order-disorder transition with a peak at  $(2670 \pm 30) \text{ K}$ .<sup>380</sup> The heat capacity maximum of this peak reaches values above  $215 \text{ J K}^{-1} \text{ mol}^{-1}$ . Above this transition also the role of Schottky effects must be taken into account.<sup>394</sup>

Numerous evaluations of the heat capacity of solid and liquid  $\text{UO}_2$  have been reported. The listing for the CODATA Key values<sup>315</sup> is based on the evaluation in Glushko *et al.*,<sup>395</sup> which is also adopted for the Equation of State description of  $\text{UO}_2$  by Ronchi *et al.*<sup>396</sup> That recommendation does not include the heat capacity data by Ronchi and coworkers<sup>380,386,387</sup> in the premelting and liquid range. For that reason we have re-fitted the experimental data in a combined treatment of enthalpy increments and heat capacity, to give

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 66.7437 + 43.1393 \times 10^{-3} (T/\text{K}) \\ & - 35.640 \times 10^{-6} (T/\text{K})^2 \\ & + 11.655 \times 10^{-9} (T/\text{K})^3 \\ & - 1.16863 \times 10^6 (T/\text{K})^{-2}. \end{aligned}$$

This single equation describes the data below and above the  $\lambda$  transition, since no change in the slope of the heat capacity or enthalpy curves were observed.

The enthalpy increment of liquid  $\text{UO}_2$  has been measured by Hein and Flagella<sup>376</sup> (four temperatures up to 3270 K) and by Leibowitz *et al.*<sup>390</sup> (six temperatures between 3173 and 3523 K). These measurements suggest a constant heat capacity value in this temperature range. The heat capacity measurements for liquid  $\text{UO}_2$  by<sup>387</sup> show a decrease from about  $120 \text{ J K}^{-1} \text{ mol}^{-1}$  near the melting point to about  $84 \text{ J K}^{-1} \text{ mol}^{-1}$  at 4500 K, followed by an increase to the maximum temperature of the measurements, 8200 K. Since it is not possible to fit these data into a single polynomial equation of the type used in the present assessment, we have fitted the heat capacity data by Ronchi *et al.*<sup>387</sup> from the melting point to  $T = 5000 \text{ K}$  to the following equation:

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 1365.4956 - 0.85866 (T/\text{K}) \\ & + 191.305 \times 10^{-6} (T/\text{K})^2 \\ & - 14.1608 \times 10^{-9} (T/\text{K})^3. \end{aligned}$$

The enthalpy of fusion of slightly hypostoichiometric  $\text{UO}_2$  samples was derived from enthalpy increment measurements as  $76 \text{ kJ mol}^{-1}$  by Hein and Flagella<sup>376</sup> and  $74 \text{ kJ mol}^{-1}$  by Leibowitz *et al.*<sup>377</sup> We select

$$\Delta_{\text{fus}}H^\circ = (75 \pm 3) \text{ kJ mol}^{-1}.$$

The selected enthalpy of fusion together with the heat capacity equations reproduce the enthalpy data for liquid  $\text{UO}_2$  by Hein and Flagella<sup>376</sup> and Leibowitz *et al.*<sup>390</sup> within 2%.

### 5.7.3. Enthalpy of formation

The enthalpy of formation of  $\text{UO}_2$  is a CODATA Key value for Thermodynamics,<sup>315</sup> which has been accepted here. This value is based on the combustion calorimetric experiments work by Huber and Holley Jr.<sup>346</sup> and Johnson and Cordfunke<sup>397</sup>:

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1085.0 \pm 1.0) \text{ kJ mol}^{-1}$$

## 5.8. $\text{Np}_2\text{O}_5(\text{cr,l})$

### 5.8.1. Crystal structure

$\text{Np}_2\text{O}_5$  has a monoclinic structure (space group P2/c). It decomposes to  $\text{NpO}_2$  and  $\text{O}_2$  at about 700 K.<sup>398</sup>

### 5.8.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Np}_2\text{O}_5$  has not been measured. Merli and Fuger<sup>399</sup> have estimated the entropy at room temperature as  $S^\circ(298.15 \text{ K}) = (186 \pm 15) \text{ J K}^{-1} \text{ mol}^{-1}$ , and Lemire<sup>400</sup> as  $S^\circ(298.15 \text{ K}) = (163 \pm 23) \text{ J K}^{-1} \text{ mol}^{-1}$ . We select

$$S^\circ(298.15 \text{ K}) = (186 \pm 15) \text{ J K}^{-1} \text{ mol}^{-1}.$$

as this value is in agreement with the variation of the standard entropy of the uranium oxides.

The high-temperature heat capacity of  $\text{Np}_2\text{O}_5$  has been measured by drop calorimetry from 350 to 750 K by Belyaev *et al.*,<sup>401</sup> who gave the following equation:

$$C_p^\circ(T)/(\text{J K}^{-1} \text{ mol}^{-1}) = 99.2 + 98.6 \times 10^{-3}(T/\text{K}).$$

### 5.8.3. Enthalpy of formation

The enthalpy of formation of  $\text{Np}_2\text{O}_5$  has been measured by Belyaev *et al.*<sup>402</sup> and by Merli and Fuger<sup>399</sup> by solution calorimetry. The results of the two studies,  $-2148 \text{ kJ mol}^{-1}$  and  $-(2162.7 \pm 9.3) \text{ kJ mol}^{-1}$  respectively, are in poor agreement. We select the value derived by Merli and Fuger<sup>399</sup> as it was based on a well-defined sample and a quick and well-defined dissolution reaction. It remains unchanged when recalculated

$$\Delta_f H^\circ(298.15 \text{ K}) = -(2162.7 \pm 9.3) \text{ kJ mol}^{-1}.$$

## 5.9. $\text{NpO}_2(\text{cr,l})$

### 5.9.1. Melting point

$\text{NpO}_2$  has a face-centered cubic crystal structure (space group  $\text{Fm}\bar{3}\text{m}$ ) which is stable up to the melting point. The melting point of  $\text{NpO}_2$  was reported to be  $(2833 \pm 50) \text{ K}$  by Chikalla *et al.*,<sup>403</sup> which becomes  $(2836 \pm 50) \text{ K}$  on ITS-90. However, recent work by Böhler *et al.*<sup>404</sup> using self-crucible laser melting gave a value of  $(3072 \pm 66) \text{ K}$ , which is selected here. Due to the high oxygen pressure of neptunium dioxide at temperatures close to melting, it cannot be excluded that the congruent melting composition of this compound be slightly hypostoichiometric.

### 5.9.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{NpO}_2$  has been measured by Westrum Jr. *et al.*<sup>405</sup> by adiabatic calorimetry from 5 to 300 K, yielding for the standard entropy:

$$S^\circ(298.15 \text{ K}) = (80.3 \pm 0.4) \text{ J K}^{-1} \text{ mol}^{-1}.$$

Magnani *et al.*<sup>316</sup> reported heat capacity data that are in excellent agreement, but the numerical details of this measurement have not been reported.

The high-temperature enthalpy increment of  $\text{NpO}_2$  has been measured by Arkhipov *et al.*<sup>406</sup> from 350 to 1100 K, by Nishi *et al.*<sup>407</sup> from 334 to 1071 K and by Beneš *et al.*<sup>408</sup> from 376 to 1770 K. The latter two studies were made on very small samples (less than 100 mg). The results by Arkhipov *et al.*<sup>406</sup> are in poor agreement with the low-temperature data and are significantly higher than the other two studies (Fig. 20). The results by Nishi *et al.*<sup>407</sup> and Beneš *et al.*<sup>408</sup> are in excellent agreement and fit the low temperature data very well (Fig. 20). Several estimates of the heat capacity of  $\text{NpO}_2$  have been also reported. Yamashita *et al.*<sup>409</sup> and Serizawa *et al.*<sup>410</sup> calculated the lattice heat capacity from the phonon and dilatation contributions using Debye temperature, thermal expansion and Grüneisen constants and the electronic contributions from crystal field energies. As shown in Fig. 20 they are in fair agreement with the experimental data.

Above 2000 K, it is likely that the heat capacity of  $\text{NpO}_2$  exhibits an excess component due to defect formation

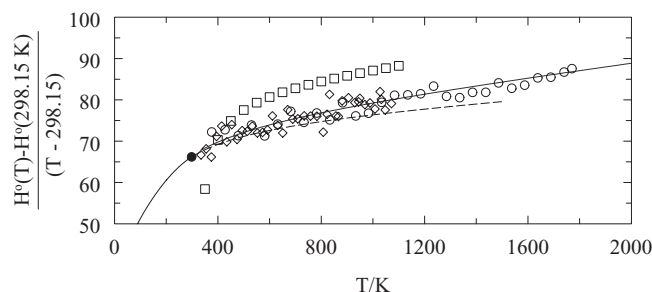


FIG. 20. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{NpO}_2$ ;  $\square$ , Arkhipov *et al.*<sup>406</sup>;  $\diamond$ , Nishi *et al.*<sup>407</sup>;  $\circ$ , Beneš *et al.*<sup>408</sup>;  $\bullet$ , value derived from the low-temperature measurements by Westrum Jr. *et al.*<sup>405</sup>; the dashed curve shows the recommended equation based on the estimates of Serizawa *et al.*<sup>410</sup>

(principally oxygen Frenkel pairs). No information on this effects exists for  $\text{NpO}_2$ , but Konings and Beneš<sup>330</sup> estimated this contribution from the heat capacity values for  $\text{ThO}_2$ ,  $\text{UO}_2$ , and  $\text{PuO}_2$  by interpolating the enthalpy of oxygen Frenkel pair formation. By adding this contribution to the representation of the experimental results they obtained

$$C_p^\circ(T)/(\text{J K}^{-1} \text{mol}^{-1}) = 64.7712 + 43.8574 \times 10^{-3}(T/\text{K}) - 35.0695 \times 10^{-6}(T/\text{K})^2 + 13.1917 \times 10^{-9}(T/\text{K})^3 - 0.78932 \times 10^6(T/\text{K})^{-2}$$

which has been constrained to  $66.20 \text{ J K}^{-1} \text{mol}^{-1}$ , as derived from the low-temperature heat capacity measurements by Westrum Jr. *et al.*<sup>405</sup>

No experimental data are available for liquid neptunium dioxide. We estimate

$$C_p^\circ(\text{NpO}_2, \text{liq}, T) = 66 \text{ J K}^{-1} \text{mol}^{-1}.$$

The entropy of fusion is assumed to be identical to that of  $\text{UO}_2$  ( $24 \text{ J K}^{-1} \text{mol}^{-1}$ ), yielding

$$\Delta_{\text{fus}}H^\circ = (70 \pm 6) \text{ kJ mol}^{-1}.$$

### 5.9.3. Enthalpy of formation

The enthalpy of formation of neptunium dioxide has been measured by Huber and Holley<sup>411</sup> by oxygen bomb calorimetry, the product being stoichiometric  $\text{NpO}_2$ :

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1078.5 \pm 2.7) \text{ kJ mol}^{-1}.$$

## 5.10. $\text{PuO}_2(\text{cr}, \text{l})$

### 5.10.1. Melting point

$\text{PuO}_2$  has a face-centered fluorite structure (space group  $\text{Fm}\bar{3}\text{m}$ ) up to its melting point. The various measurements of the melting point of  $\text{PuO}_2$  are summarized in Table 60. Because  $\text{PuO}_2$ , similar to  $\text{CeO}_2$ , starts to lose oxygen according

TABLE 60. The melting point of  $\text{PuO}_2(\text{cr})$

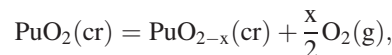
Authors	$T_{\text{fus}}/\text{K}$	
	Reported	ITS-90
Pijanowski and DeLuca <sup>363</sup>	$2569 \pm 30^{\text{a}}$	$2586 \pm 30$
Russel <sup>415</sup>	$2673^{\text{b}}$	
Chikalla <sup>364</sup>	$2553 \pm 30^{\text{a}}$	$2556 \pm 30$
Freshley and Mattys <sup>416</sup>	2523	
Lyon and Bailey <sup>365</sup>	$2511 \pm 135^{\text{c}}$	$2513 \pm 135$
Lyon and Bailey <sup>367</sup>	$2663 \pm 20^{\text{c}}$	$2666 \pm 20$
Aitken and Evans <sup>417</sup>	$2663^{\text{a}}$	
Riley <sup>412</sup>	$2673 \pm 20^{\text{b}}$	$2682 \pm 20$
Kato <i>et al.</i> <sup>372</sup>	2843	2843
De Bruycker <i>et al.</i> <sup>413</sup>		$3017 \pm 28$
Selected value:		$3017 \pm 28$

<sup>a</sup>in He.

<sup>b</sup>in Ar.

<sup>c</sup> $\text{O}_2$ .

to the reaction:



the melting temperature determinations are difficult to interpret. For example, Chikalla<sup>364</sup> found that samples of stoichiometric  $\text{PuO}_2$  had a O/Pu ratio near 1.62 after melting in inert gas atmosphere. Riley<sup>412</sup> realised that the melting point of stoichiometric plutonium dioxide could only be defined under a high oxygen pressure. However, his flame melting experiments were still imprecise and the samples were still affected by high temperature reduction despite the controlled atmosphere. Also the interaction of the liquid with container materials affects the results, as demonstrated by Kato *et al.*<sup>372</sup> They obtained a melting point for pure  $\text{PuO}_2$  about 200 K higher than the earlier values, when using rhenium as container material instead of tungsten. This was further confirmed by De Bruycker *et al.*<sup>413,414</sup> who employed a containerless laser melting technique in oxygen and found an even higher melting temperature for stoichiometric  $\text{PuO}_2$ ,  $T_{\text{fus}} = (3017 \pm 28) \text{ K}$ , which is our selected value.

### 5.10.2. Heat capacity and entropy

Low-temperature heat capacity measurements have been performed on samples of  $^{239}\text{PuO}_2$  by Sandenaw<sup>418</sup> from 15 to 325 K and Kruger and Savage<sup>419</sup> from 192 to 320 K, as well as the less radioactive  $^{242}\text{PuO}_2$  and  $^{244}\text{PuO}_2$  from 12 to 350 K and 4 to 25 K, respectively, by Flotow *et al.*<sup>420</sup> In the latter samples the effects of accumulation of radiation damage are less significant and more accurate results in the very low temperature range can be obtained. For that reason our selected value for the entropy is solely based on the results by Flotow *et al.*<sup>420</sup>:

$$S^\circ(298.15 \text{ K}) = (66.13 \pm 0.30) \text{ J K}^{-1} \text{mol}^{-1}.$$

High-temperature heat capacity of  $\text{PuO}_2$  has been measured by Engel<sup>384</sup> from 300 to 1100 K, and high-temperature enthalpy increments have been measured by Kruger and Savage<sup>419</sup> from 298 to 1404 K, Ogard<sup>421</sup> from 1500 to 2715 K, and Oetting<sup>422</sup> from 353 to 1610 K (Fig. 21). Ogard's measurements suggest a rapid increase in  $C_p$  above 2370 K. This has been attributed to partial melting of  $\text{PuO}_2$  through

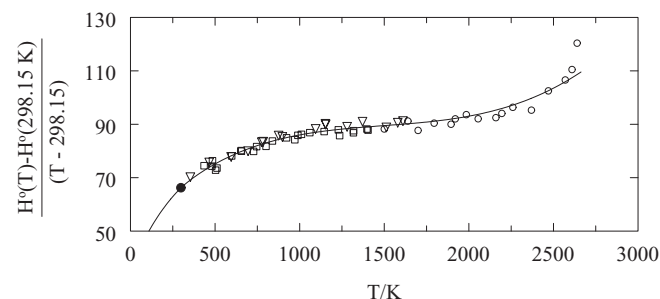


FIG. 21. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{mol}^{-1}$ ) of  $\text{PuO}_2$ ;  $\circ$ , Ogard<sup>421</sup>;  $\square$ , Kruger and Savage<sup>419</sup>;  $\nabla$ , Oetting<sup>422</sup>;  $\bullet$ , value derived from the low-temperature measurements by Flotow *et al.*<sup>420</sup>; the curve shows the recommended equation.



interaction with the tungsten container,<sup>422,423</sup> although no evidence exists for this interaction. A similar rapid increase is found in the heat capacities of ThO<sub>2</sub>, UO<sub>2</sub>, and ZrO<sub>2</sub>, which has been attributed to the formation of Frenkel and Schottky lattice defects at high temperatures. For that reason we have fitted all experimental data to a polynomial equation constrained to  $C_p^\circ(298.15\text{ K}) = 66.25\text{ J K}^{-1}\text{ mol}^{-1}$ , as derived by Flotow *et al.*,<sup>420</sup> though it proved to be very difficult to fit the results to the standard form due to strong variation between room temperature and melting temperature. The following equation gives an acceptable description up to 2300 K and includes the upward trend, though the results of Ogard<sup>421</sup> in the 2470–2640 K range, which were given a lower weight, could not be reproduced accurately,

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) &= 35.2952 + 0.15225(\text{T/K}) \\ &\quad - 127.255 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 36.289 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 3.47593 \times 10^5(\text{T/K})^{-2}. \end{aligned}$$

No data for the heat capacity or enthalpy of liquid PuO<sub>2</sub> are known, except a single enthalpy measurement by Ogard.<sup>421</sup> The enthalpy of fusion is thus estimated, assuming that the entropy of fusion is identical to that of UO<sub>2</sub> (22.4 J K<sup>-1</sup> mol<sup>-1</sup>), yielding

$$\Delta_{fus}H^\circ = (64 \pm 6)\text{ kJ mol}^{-1}.$$

We have also estimated for the heat capacity of liquid PuO<sub>2</sub> from the value for UO<sub>2</sub> as

$$C_p^\circ = 70\text{ J K}^{-1}\text{ mol}^{-1}.$$

### 5.10.3. Enthalpy of formation

The enthalpy of formation of PuO<sub>2</sub> has been determined by Popov *et al.*,<sup>424</sup> Holley *et al.*<sup>425</sup> and Johnson *et al.*<sup>426</sup> by oxygen combustion calorimetry starting from pure plutonium metal. The results are in very good agreement, as shown in Table 61. We select

$$\Delta_f H^\circ(298.15\text{ K}) = -(1055.8 \pm 1.0)\text{ kJ mol}^{-1}$$

which is based to the appreciably more precise value by Johnson *et al.*,<sup>426</sup> who carefully analysed the metal for impurities and applied appropriate corrections.

TABLE 61. The enthalpy of formation of plutonium dioxide

Authors	Method <sup>a</sup>	$\Delta_f H^\circ(298.15\text{ K})/\text{kJ mol}^{-1}$
Popov <i>et al.</i> <sup>424</sup>	C	-1056.0 ± 4.6
Holley <i>et al.</i> <sup>425</sup>	C	-1058.0 ± 1.6
Johnson <i>et al.</i> <sup>426</sup>	C	-1055.8 ± 1.0
Selected value:		-1055.8 ± 1.0

<sup>a</sup>C = combustion calorimetry.

## 5.11. Pu<sub>2</sub>O<sub>3</sub>(cr,l)

### 5.11.1. Polymorphism and melting point

Pu<sub>2</sub>O<sub>3</sub> has a hexagonal type-A rare-earth sesquioxide structure (space group P $\bar{3}$ m1). It is likely that Pu<sub>2</sub>O<sub>3</sub> may exhibit a similar high-temperature behavior as the light lanthanide sesquioxides, with the eventual appearance of H-type and the X-type structures before melting. This is however not confirmed experimentally. The melting point of stoichiometric Pu<sub>2</sub>O<sub>3</sub> was measured by Chikalla *et al.*<sup>427</sup> and Riley<sup>412</sup> being in excellent agreement (Table 62), but significantly lower than the value reported by Holley *et al.*<sup>425</sup> for a less well-characterised sample. We select (2352 ± 10) K, based on the results of Riley,<sup>412</sup> corrected to ITS-90, which is, however, relatively low compared to the lanthanide sesquioxides as well as Am<sub>2</sub>O<sub>3</sub> and Cm<sub>2</sub>O<sub>3</sub> (see Sec. 7.2.2).

### 5.11.2. Heat capacity and entropy

Flotow and O'Hare<sup>428</sup> have measured the heat capacity of a sample of <sup>244</sup>Pu<sub>2</sub>O<sub>3</sub> from 8 K to 350 K. Their results revealed a  $\lambda$ -type anomaly in the heat capacity at 17.65 K, associated with an antiferromagnetic transition. The standard entropy derived from this work is:

$$S^\circ(298.15\text{ K}) = (163.02 \pm 0.65)\text{ J K}^{-1}\text{ mol}^{-1}$$

There are no measurements of the heat capacity or enthalpy of Pu<sub>2</sub>O<sub>3</sub> at high temperatures. We have here estimated the following equation, based on comparison of actinide and lanthanide oxides, which fits  $C_p^\circ(298.15\text{ K}) = 116.98\text{ J K}^{-1}\text{ mol}^{-1}$  from the low-temperature heat capacity measurements.<sup>428</sup>

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1}\text{ mol}^{-1}) &= 130.6670 + 18.4357 \times 10^{-3}(\text{T/K}) \\ &\quad - 1.70530 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

Previously estimated values have been presented by various authors,<sup>395,429</sup> but these are about 10 kJ mol<sup>-1</sup> higher at 2000 K.

We assume that Pu<sub>2</sub>O<sub>3</sub> transforms to the H-type structure at  $T_{trs} = (2300 \pm 50)\text{ K}$ , like in the lanthanide sesquioxides and estimate for the enthalpies of transition and fusion:

$$\begin{aligned} \Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (32 \pm 10)\text{ kJ mol}^{-1} \\ \Delta_{fus}H^\circ &= (71 \pm 10)\text{ kJ mol}^{-1} \end{aligned}$$

TABLE 62. Temperature of melting of plutonium sesquioxide

Authors	$T_{fus}/\text{K}$	
	Reported	ITS-90
Holley <i>et al.</i> <sup>425</sup>	2513 ± 33	2516 ± 33
Chikalla <i>et al.</i> <sup>427</sup>	2358 ± 25	2361 ± 25
Riley <sup>412</sup>	2348 ± 5	2352 ± 5
Selected value:		2352 ± 10

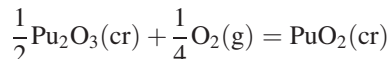
For the heat capacity of the high temperature modifications we assume the same values as for  $\text{Sm}_2\text{O}_3$ :

$$C_p^\circ(\text{H}, T) = 165 \text{ J K}^{-1} \text{ mol}^{-1}$$

$$C_p^\circ(\text{liq}, T) = 179 \text{ J K}^{-1} \text{ mol}^{-1}$$

### 5.11.3. Enthalpy of formation

The assessment of the enthalpy of formation of  $\text{Pu}_2\text{O}_3$  is not straightforward, as direct measurements do not exist. The value must be based on the analysis of the reaction:



This was extensively discussed in various reviews,<sup>8,395,430</sup> in which similar results have been derived. These studies generally refer to the assessment by Markin and Rand<sup>431</sup> of the oxygen potential measurements by the same authors. As discussed by Lemire *et al.*,<sup>430</sup> the work of Chereau *et al.*<sup>432</sup> suggests, however, that the partial enthalpies and entropies from the work of Markin and Rand might need some adjustment. Guéneau *et al.*,<sup>357,433</sup> presented a consistent analysis of the phase diagram and oxygen potential data in the Pu-O system using the CALPHAD method. The results of that analysis are in reasonable agreement with the analysis by Rand,<sup>8,430</sup> though a slight change in the enthalpy of formation of  $\text{PuO}_2$  has been suggested, which is not consistent with the present review. If we take the Gibbs energy of the above reaction, and combine it with the selected values from this review, we obtain  $\Delta_f H^\circ(298.15 \text{ K}) = -1647 \text{ kJ mol}^{-1}$ , somewhat lower than the value by Rand,<sup>8,430</sup>  $\Delta_f H^\circ(298.15 \text{ K}) = -(1656 \pm 10) \text{ kJ mol}^{-1}$ , or Glushko *et al.*,<sup>395</sup>  $\Delta_f H^\circ(298.15 \text{ K}) = -(1670 \pm 20) \text{ kJ mol}^{-1}$ . We select

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1647 \pm 10) \text{ kJ mol}^{-1}.$$

## 5.12. $\text{AmO}_2(\text{cr,l})$

### 5.12.1. Melting point

Americium dioxide has a fcc fluorite structure (space group  $\text{Fm}\bar{3}\text{m}$ ). The melting point of  $\text{AmO}_2$  has been measured by McHenry.<sup>434</sup> His experiments were hindered by the effect of dissociation, but McHenry concluded from measurements at different heating rates that the melting point of the dioxide is about 2383 K. However, this value is probably not referring to stoichiometric  $\text{AmO}_2$ , but to  $\text{AmO}_{2-x}$  of undefined composition. Stoichiometric melting of  $\text{AmO}_2$  is unlikely to occur at atmospheric pressure, as follows from the phase diagram.<sup>435</sup> Upon heating the oxide starts to lose oxygen and transforms into  $\text{AmO}_{2-x}$ , which has a wide range of composition.

### 5.12.2. Heat capacity and entropy

No experimental study of the low-temperature heat capacity of  $\text{AmO}_2$  has been reported. The estimate by Westrum Jr. and Grönvold<sup>436</sup> for the standard entropy,  $S^\circ(298.15 \text{ K}) = 83.7$

$\text{J K}^{-1} \text{ mol}^{-1}$ , is somewhat too high, as has been demonstrated by Konings,<sup>329,437</sup> who analysed the systematics in the entropies of actinide(IV) compounds, deriving  $S^\circ(298.15 \text{ K}) = 75.5 \text{ J K}^{-1} \text{ mol}^{-1}$ , from a lattice contribution ( $63.0 \text{ J K}^{-1} \text{ mol}^{-1}$ ) and an excess contribution ( $12.5 \text{ J K}^{-1} \text{ mol}^{-1}$ ) from the  $^6\text{H}_{5/2}$  electronic state of the  $\text{Am}^{4+}$  ion. This value is adopted here with an estimated uncertainty

$$S^\circ(298.15 \text{ K}) = (75.5 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

This value is also consistent with preliminary calculations of the vaporization data for americium oxides dissolved in plutonium oxides.<sup>438</sup>

The enthalpy increment of  $\text{AmO}_2$  has been measured by Nishi *et al.*<sup>439</sup> on a sample of about 50 mg encapsulated in a platinum container. As a result, the scatter in the data is relatively large, and the values at the lowest temperatures are somewhat uncertain. Consequently the fit of the data by Nishi *et al.*<sup>439</sup> yields too low values near room temperature (Fig. 22). For that reason we have refitted the enthalpy increment data constrained to  $C_p^\circ(298.15 \text{ K}) = 64.3 \text{ J K}^{-1} \text{ mol}^{-1}$ , as derived from the trend in the  $\text{AnO}_2$  series by Konings.<sup>329</sup> We thus obtain

$$C_p^\circ / \text{J K}^{-1} \text{ mol}^{-1} = 78.9718 + 3.8365 \times 10^{-3}(T/\text{K}) - 1.40591 \times 10^6(T/\text{K})^{-2}.$$

Above about 1200 K, this equation has been fitted to the heat capacity estimated by Thiriet and Konings<sup>440</sup> taking  $\text{ThO}_2$  as lattice and calculating the excess from the free ion energy levels.

### 5.12.3. Enthalpy of formation

The enthalpy of formation of  $^{243}\text{AmO}_2$  has been measured by Morss and Fuger<sup>441</sup> who determined the enthalpy of solution of a well-characterized sample in a  $\{\text{H}_2\text{SO}_4 + \text{KI}\}$  solution using a micro-calorimeter. The thermochemical cycle used in that work, was based on the dissolution of  $\text{AmCl}_3$  in the same medium. Their resulting value, which remains unchanged after recalculation, has been selected here as

$$\Delta_f H^\circ(298.15 \text{ K}) = -(932.2 \pm 3.0) \text{ kJ mol}^{-1}.$$

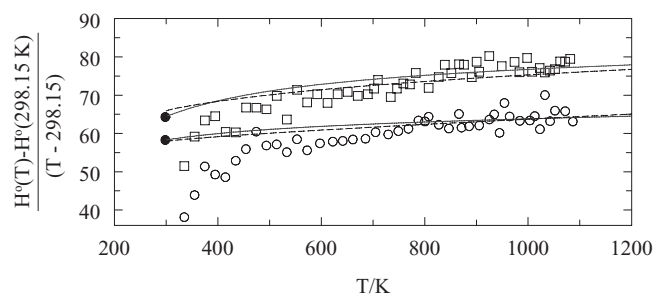


FIG. 22. The reduced enthalpy increment (in  $\text{J K}^{-1} \text{ mol}^{-1}$ ) of  $\text{AmO}_2$  ( $\square$ ) and  $\text{AmO}_{1.5}$  ( $\circ$ ) by Nishi *et al.*<sup>439</sup>; the solid curve shows the recommended equations, the dashed curves the estimates based on comparison with other lanthanide and actinide dioxides and sesquioxides.

Eyring *et al.*<sup>80</sup> measured the enthalpy of solution of  $^{241}\text{AmO}_2$  in a  $\{\text{HNO}_3+\text{HBF}_4\}$  solution and derived for the enthalpy of formation  $\Delta_f H^\circ(298.15\text{ K}) = -(1003.7\text{ kJ mol}^{-1})$ , substantially more negative. The thermochemical cycle used in that study is based on (estimated) extrapolation of the result to infinite dilution, which introduces large uncertainties.

### 5.13. $\text{Am}_2\text{O}_3(\text{cr,l})$

#### 5.13.1. Polymorphism and melting point

The stable crystal structure of americium sesquioxide at room temperature is not fully established. The early studies report a bcc C-type rare-earth structure (space group Ia3) at room temperatures. However, since  $\text{Pu}_2\text{O}_3$  has a A-type hexagonal and  $\text{Cm}_2\text{O}_3$  a B-type monoclinic structure, this is not likely. Considering the ionic radius of  $\text{Am}^{3+}$  the B-type structure is more probable (see also Sec. 7.2.2). This would be consistent with the transformation to the rare-earth type-A  $\text{La}_2\text{O}_3$  structure at a temperature between 1073 and 1173 K as observed by Wallmann.<sup>442</sup> However, most experimental studies on  $\text{Am}_2\text{O}_3$  have been made on the hexagonal form.

The melting point of  $\text{Am}_2\text{O}_3$  was found as  $(2478 \pm 15)\text{ K}$  by Chikalla *et al.*,<sup>443,444</sup> which is  $T_{fus} = (2481 \pm 15)\text{ K}$  on ITS-90.

#### 5.13.2. Heat capacity and entropy

The low-temperature heat capacity of  $\text{Am}_2\text{O}_3$  has not been measured but several estimates have been made. Westrum Jr. and Grønvdal<sup>436</sup> estimated the value  $S^\circ(298.15\text{ K}) = 158.2\text{ J K}^{-1}\text{ mol}^{-1}$  (specified as cubic), by describing the entropy as the sum of the lattice entropy and an excess contribution. A similar approach, but using a more sound basis of spectroscopic and calorimetric information, was used by Konings.<sup>127,445</sup> The value was composed only of a lattice part, obtained by extrapolating the trend in the lanthanide sesquioxides to the actinide sesquioxides. The excess part for the  $\text{Am}^{3+}$  ion is zero as the  $^7F_0$  ground state degeneracy is 1, and the first excited state  $^7F_1$  does not contribute at room temperature. This value is accepted here

$$S^\circ(298.15\text{ K}) = (134.2 \pm 5.0)\text{ J K}^{-1}\text{ mol}^{-1}.$$

The high-temperature enthalpy increment of hexagonal (A-type)  $\text{Am}_2\text{O}_3$  has been measured by Nishi *et al.*<sup>439</sup> on a sample of about 50 mg encapsulated in a platinum container. As a result, the scatter in the data is relatively large, and the values at the lowest temperatures are somewhat uncertain (Fig. 22). Consequently the fit of the data by Nishi *et al.*<sup>439</sup> yields too low values near room temperature (e.g.,  $C_p^\circ(298.15\text{ K}) = 88.0\text{ J K}^{-1}\text{ mol}^{-1}$ ) considering the results for the  $\text{Ln}_2\text{O}_3$  compounds, for which the lattice heat capacity at 298.15 K is between 110 and 100  $\text{J K}^{-1}\text{ mol}^{-1}$ , and the value for  $\text{Pu}_2\text{O}_3$  ( $116.98\text{ J K}^{-1}\text{ mol}^{-1}$ ). For that reason we have refitted the enthalpy increment data above 700 K, constrained to  $C_p^\circ = 116.5\text{ J K}^{-1}\text{ mol}^{-1}$ , as derived from comparison with the

$\text{Ln}_2\text{O}_3$  series and the value for  $\text{Pu}_2\text{O}_3$ . We thus obtain

$$C_p^\circ/\text{J K}^{-1}\text{ mol}^{-1} = 126.0084 + 8.0097 \times 10^{-3}(\text{T/K}) - 1.05752 \times 10^6(\text{T/K})^{-2}.$$

The equation has been fitted at high temperature to the heat capacity estimated on the basis of comparison to  $\text{Pu}_2\text{O}_3$  and the lanthanide sesquioxides (Fig. 22).

The transition to the H-type structure is assumed to take place at  $T_{trs} = (2350 \pm 50)\text{ K}$ , and we estimate for the enthalpies of transition and fusion:

$$\begin{aligned}\Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (33 \pm 10)\text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (10 \pm 5)\text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (74 \pm 10)\text{ kJ mol}^{-1}.\end{aligned}$$

For the heat capacity of the high temperature modifications we assume the same values as for  $\text{Eu}_2\text{O}_3$ :

$$C_p^\circ(\text{H}, \text{T}) = 141\text{ J K}^{-1}\text{ mol}^{-1}, C_p^\circ(\text{liq}, \text{T}) = 156\text{ J K}^{-1}\text{ mol}^{-1}.$$

#### 5.13.3. Enthalpy of formation

Morss and Sonnenberger<sup>446</sup> have derived the enthalpy of formation of hexagonal  $\text{Am}_2\text{O}_3$  from enthalpy-of-solution measurements in  $6\text{ mol dm}^{-3}$  hydrochloric acid using a microcalorimeter. This value remains unchanged after recalculation. The thermochemical cycle used in that work, was based on the dissolution of  $\text{Am}(\text{cr})$  in the same medium, based on the work of Fuger *et al.*<sup>447</sup>

$$\Delta_f H^\circ(298.15\text{ K}) = -(1690.4 \pm 7.9)\text{ kJ mol}^{-1}.$$

### 5.14. $\text{Cm}_2\text{O}_3(\text{cr})$

#### 5.14.1. Polymorphism and melting point

At room temperature stoichiometric curium sesquioxide has a monoclinic B-type crystal structure. Upon heating, three high-temperature transformations have been identified.<sup>92,403</sup> At  $(1888 \pm 15)\text{ K}$  the monoclinic form transforms into the A-type hexagonal structure, as was observed by high-temperature X-ray diffraction.<sup>448</sup> Above 2000 K transitions at  $(2273 \pm 20)\text{ K}$  and at  $(2383 \pm 20)\text{ K}$  have been observed, which are most likely transformations to the H and X structures that are known for the lanthanide sesquioxides.

There are several reports of the melting point of  $\text{Cm}_2\text{O}_3$ , as summarized in Table 63. The values are in good agreement,

TABLE 63. The melting point of  $\text{Cm}_2\text{O}_3(\text{cr})$

Authors	$T_{fus}/\text{K}$	
	Reported	ITS-90
McHenry <sup>434</sup>	2223	
Smith <sup>449</sup>	$2538 \pm 20$	$2545 \pm 20$
Gibby <i>et al.</i> <sup>92a</sup>	$2548 \pm 25$	$2551 \pm 25$
	2538	2541
Baybarz <sup>450</sup>	$2533 \pm 20$	$2532 \pm 20$
Selected value		$2542 \pm 25$

<sup>a</sup>Also reported by Chikalla *et al.*<sup>403</sup>

except for the results of McHenry,<sup>434</sup> who used CmO<sub>2</sub> as starting material and assumed that it was converted to the sesquioxide during the measurements. For all values a small correction to the ITS-90 temperature scale is needed. Smith<sup>449</sup> used the melting points of Al<sub>2</sub>O<sub>3</sub> and Tm<sub>2</sub>O<sub>3</sub> as reference. For the former, a value about 7 K lower than the currently accepted value was obtained. For Tm<sub>2</sub>O<sub>3</sub> no recommended value is available. We correct the value by Smith therefore with +7 K. Regarding the results of Gibby *et al.*,<sup>92</sup> Chikalla *et al.*<sup>403</sup> we think it is likely that they refer to IPTS-48. The selected value for the melting point is  $T_{fus} = (2542 \pm 25)$  K.

### 5.14.2. Heat capacity and entropy

The low-temperature heat capacity of Cm<sub>2</sub>O<sub>3</sub> has not been measured but several estimates have been made. Westrum Jr. and Grønvold<sup>436</sup> have estimated the value  $S^\circ(298.15 \text{ K}) = 160.7 \text{ J K}^{-1} \text{ mol}^{-1}$  (specified as cubic), by describing the entropy as the sum of the lattice entropy and an excess contribution. A similar approach, but using a more sound basis of spectroscopic and calorimetric information, was used by Konings<sup>127,445</sup> who obtained  $(167.0 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$  for the monoclinic modification. The value was composed of a lattice part ( $133.5 \text{ J K}^{-1} \text{ mol}^{-1}$ ), obtained by extrapolating the trend in the lanthanide sesquioxides to the actinide sesquioxides, and an excess part ( $34.58 \text{ J K}^{-1} \text{ mol}^{-1}$ ), calculated from the ground state degeneracy of the Cm<sup>3+</sup> ion. This value is accepted here

$$S^\circ(298.15 \text{ K}) = (168.1 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}.$$

Estimated values for the high-temperature heat capacity of monoclinic Cm<sub>2</sub>O<sub>3</sub> have been presented by Gibby *et al.*,<sup>92</sup> who used them to convert the thermal diffusivity measurements to thermal conductivity data, which agree reasonably with direct measurements (see below). As discussed by Konings<sup>437</sup> Gibby's values are rather high compared to the experimental values for the lanthanide sesquioxides and Pu<sub>2</sub>O<sub>3</sub>. He estimated the following equation:

$$C_p^\circ(T)/\text{J K}^{-1} \text{ mol}^{-1} = 123.532 + 14.550 \times 10^{-3}(T/\text{K}) - 1.3489 \times 10^6(T/\text{K})^{-2}.$$

Data on the heat capacity of the H, X, and liquid (L) phases and enthalpies of transition for C → H, H → X, and X → L are not available. Although rather speculative we have estimated the properties of the high temperature phases in a similar manner as for the lanthanide sesquioxides, and derive for Cm<sub>2</sub>O<sub>3</sub>:

$$\begin{aligned}\Delta_{trs}H^\circ(\text{B} \rightarrow \text{A}) &= (6 \pm 3) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{A} \rightarrow \text{H}) &= (32 \pm 6) \text{ kJ mol}^{-1}, \\ \Delta_{trs}H^\circ(\text{H} \rightarrow \text{X}) &= (10 \pm 5) \text{ kJ mol}^{-1}, \\ \Delta_{fus}H^\circ &= (66 \pm 10) \text{ kJ mol}^{-1}.\end{aligned}$$

For the heat capacity of the high temperature H modification and liquid Cm<sub>2</sub>O<sub>3</sub> we estimate

$$\begin{aligned}C_p^\circ(\text{A}, T) &= 142 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{H}, T) &= C_p^\circ(\text{X}, T) = 130 \text{ J K}^{-1} \text{ mol}^{-1}, \\ C_p^\circ(\text{liq}, T) &= 140 \text{ J K}^{-1} \text{ mol}^{-1}.\end{aligned}$$

### 5.14.3. Enthalpy of formation

The enthalpy of solution of curium sesquioxide in 6 mol dm<sup>-3</sup> HCl(aq) has been measured by Morss *et al.*<sup>451</sup> by solution calorimetry using milligram samples of the monoclinic form. Combining this result with the enthalpy of solution of Cm metal in this solvent, estimated from the value for the dissolution in 1.0 mol dm<sup>-3</sup> HCl(aq), the enthalpy of formation is calculated as

$$\Delta_f H^\circ(298.15 \text{ K}) = -(1684 \pm 14) \text{ kJ mol}^{-1}.$$

## 5.15. BkO<sub>2</sub>(cr) and Bk<sub>2</sub>O<sub>3</sub>(cr)

### 5.15.1. Polymorphism and melting point

BkO<sub>2</sub> has a fcc fluorite structure (space group Fm $\bar{3}$ m). The stable crystallographic modification of Bk<sub>2</sub>O<sub>3</sub> is the bcc C-type rare-earth structure (space group Ia3).<sup>452</sup> Baybarz<sup>450</sup> studied the high temperature polymorphism and found that C-Bk<sub>2</sub>O<sub>3</sub> irreversibly transforms to a monoclinic B structure at  $(1473 \pm 50)$  K. The monoclinic-to-hexagonal transition was observed at about 2022 K and the melting point at  $(2193 \pm 25)$  K. Correction to ITS-90 would involve a correction of -1 K if the measurements would refer to IPTS-68 and +2 K in case of IPTS-48. Since this is not clear from the paper, we choose to retain the original value.

### 5.15.2. Heat capacity and entropy

No experimental data exist on the heat capacity and entropy of the berkelium oxides. Konings *et al.*<sup>306</sup> gave the following estimates for the standard entropy based on the trends in the actinide and lanthanide series:

$$S^\circ(\text{BkO}_2, \text{ cr}, 298.15 \text{ K}) = (83 \pm 5) \text{ J K}^{-1} \text{ mol}^{-1},$$

$$S^\circ(\text{Bk}_2\text{O}_3, \text{ cr}, 298.15 \text{ K}) = (173.8 \pm 5) \text{ J K}^{-1} \text{ mol}^{-1}.$$

### 5.15.3. Enthalpy of formation

The enthalpies of formation of BkO<sub>2</sub> and Bk<sub>2</sub>O<sub>3</sub>(cr) have not been determined. Konings *et al.*<sup>306</sup> gave the following estimated values:

$$\Delta_f H^\circ(\text{BkO}_2, \text{ cr}, 298.15 \text{ K}) = -(1023 \pm 9) \text{ kJ mol}^{-1},$$

$$\Delta_f H^\circ(\text{Bk}_2\text{O}_3, \text{ cr}, 298.15 \text{ K}) = -(1694 \pm 20) \text{ kJ mol}^{-1}$$

based on the correlation between the difference in the enthalpy of formations of the oxides and the aqueous ions with molar volume for the AnO<sub>2</sub> compounds.

## 5.16. CfO<sub>2</sub>(cr) and Cf<sub>2</sub>O<sub>3</sub>(cr)

### 5.16.1. Polymorphism and melting point

CfO<sub>2</sub> has a fcc fluorite structure (space group Fm $\bar{3}$ m). The stable crystallographic modification of Bk<sub>2</sub>O<sub>3</sub> is the bcc



C-type rare-earth structure (space group Ia3). It transforms to cubic or hexagonal forms due to self-irradiation.<sup>450</sup> The cubic (cr) to monoclinic (B) transformation has been observed at about 1673 K.<sup>453</sup> The melting point of Cf<sub>2</sub>O<sub>3</sub> was determined to be (2023 ± 25) K by Baybarz<sup>450</sup> in vacuum and in helium. Correction to ITS-90 would involve a correction of -1 K if the measurements would refer to ITPS-68 and +2 K in case of ITPS-48. Since this is not clear from the paper, we choose to retain the original value.

### 5.16.2. Heat capacity and entropy

No experimental data exist on the heat capacity and entropy of the berkelium oxides. Konings *et al.*<sup>306</sup> gave the following estimated values for the standard entropy based on the trends in the actinide and lanthanide series:

$$S^\circ(\text{CfO}_2, \text{ cr}, 298.15 \text{ K}) = (87 \pm 5) \text{ J K}^{-1} \text{ mol}^{-1},$$

$$S^\circ(\text{Cf}_2\text{O}_3, \text{ cr}, 298.15 \text{ K}) = (173.8 \pm 5) \text{ J K}^{-1} \text{ mol}^{-1}.$$

### 5.16.3. Enthalpy of formation

The enthalpy of formation of CfO<sub>2</sub> has not been determined. Konings *et al.*<sup>306</sup> gave the following estimated value:

$$\Delta_f H^\circ(\text{CfO}_2, \text{ cr}, 298.15 \text{ K}) = -(857 \pm 14) \text{ kJ mol}^{-1}$$

based on the correlation between the difference in the enthalpy of formation of the oxides and the aqueous ions with molar volume for the AnO<sub>2</sub> compounds. The enthalpy of formation of cubic Cf<sub>2</sub>O<sub>3</sub>(cr) has been determined by Morss *et al.*<sup>454</sup> using solution microcalorimetry on milligram samples in 6.0 mol dm<sup>-3</sup> HCl(aq), to give

$$\Delta_f H^\circ(\text{Cf}_2\text{O}_3, \text{ cr}, 298.15 \text{ K}) = -(1652.6 \pm 10.3) \text{ kJ mol}^{-1}$$

using a thermochemical cycle based on the solution of Cf(cr) for which the enthalpy was estimated from the experimental value in 1.0 mol dm<sup>-3</sup> HCl(aq) by Fuger *et al.*<sup>455</sup>

## 6. The Gaseous Actinide Oxides

### 6.1. AcO(g)

#### 6.1.1. Heat capacity and entropy

The thermal functions of AcO(g) in the standard state have been calculated using the data given in Table 64. Because no spectral data on the AcO molecule exists, the molecular constants were obtained from B3LYP calculations by Kovács<sup>456</sup> and estimations using the trends revealed in the spectral data of the lanthanide and actinide monoxides. The electronic structure is assumed to be analogous to that of LaO. The energies of the low-lying states taken from the LaO data are approximated to hundreds cm<sup>-1</sup>. The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (245.545 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

TABLE 64. Molecular constants of AcO(g)

No.	State	cm <sup>-1</sup>					<i>r<sub>e</sub></i> pm	<i>p<sub>i</sub></i>	
		<i>T<sub>e</sub></i>	<i>ω<sub>e</sub></i>	<i>ω<sub>e</sub>x<sub>e</sub></i>	<i>B<sub>e</sub></i>	<i>α<sub>e</sub>10<sup>3</sup></i>			<i>D<sub>e</sub>10<sup>7</sup></i>
0 <sup>a</sup>	X <sup>2</sup> Σ <sup>+</sup>	0	764	2.3	0.34	1.5	2.63	194	2
1 <sup>b</sup>	A' <sup>2</sup> Δ <sub>3/2</sub>	7500							2
2 <sup>b</sup>	A' <sup>2</sup> Δ <sub>5/2</sub>	8200							2
3 <sup>b</sup>	A <sup>2</sup> Π	13 100							4
4 <sup>b</sup>	B <sup>2</sup> Σ	17 900							2
5 <sup>b</sup>	C <sup>2</sup> Π	22 700							4
6 <sup>b</sup>	D <sup>2</sup> Σ	27 000							2
7 <sup>b</sup>	F <sup>2</sup> Σ	28 000							4
8 <sup>b</sup>		17 200							8
9 <sup>b</sup>		18 500							8
10 <sup>b</sup>		21 200							8
11 <sup>b</sup>		24 000							20
12 <sup>b</sup>		25 900							8
13 <sup>b</sup>		30 200							12
14 <sup>b</sup>		35 000							30
15 <sup>b</sup>		40 000							36

<sup>a</sup>Computed state.

<sup>b</sup>Estimated state

and the coefficients of the equations for the heat capacity are

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 26.7681 + 26.00447 \times 10^{-3}(\text{T/K})$$

$$- 24.23966 \times 10^{-6}(\text{T/K})^2$$

$$+ 8.10964 \times 10^{-9}(\text{T/K})^3$$

$$- 0.83961 \times 10^5(\text{T/K})^{-2}$$

for the 298.15–1100 K range, and

$$C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = 47.1310 - 13.09869 \times 10^{-3}(\text{T/K})$$

$$+ 6.02184 \times 10^{-6}(\text{T/K})^2 - 0.628330$$

$$\times 10^{-9}(\text{T/K})^3 - 2.91011$$

$$\times 10^6(\text{T/K})^{-2}$$

for the 1100–4000 K range.

### 6.1.2. Enthalpy of formation

No experimental data exists for the derivation of the enthalpy of formation of AcO(g). We have estimated the dissociation energy as *D*<sub>0</sub> = (860 ± 20) kJ mol<sup>-1</sup>. This value corresponds to:

$$\Delta_f H^\circ(\text{AcO}, \text{ g}, 298.15 \text{ K}) = -(193 \pm 20) \text{ kJ mol}^{-1}.$$

### 6.2. ThO<sub>2</sub>(g)

#### 6.2.1. Heat capacity and entropy

The thermodynamic functions of ThO<sub>2</sub>(g) in the standard state have been calculated using the data given in Table 65. There are no experimental data for the structure of free gaseous ThO<sub>2</sub> molecules, and the ThO<sub>2</sub> structure is determined only from infrared spectra in the solid matrices. In an early work Kaufman *et al.*<sup>457</sup> found that a ThO<sub>2</sub> molecular beam exhibits some deflection in an inhomogeneous electric field, indicating the possibility of a bent structure for this molecule. The first quantitative information on the bent structure of ThO<sub>2</sub> was

TABLE 65. The molecular parameters for ThO<sub>2</sub>(g)

Parameter	Value
Ground electronic state	<sup>1</sup> A <sub>1</sub>
Symmetry group	C <sub>2v</sub>
Symmetry number, σ	2
I <sub>A</sub> I <sub>B</sub> I <sub>C</sub> (g <sup>3</sup> cm <sup>6</sup> ) <sup>a</sup>	1110 × 10 <sup>-117</sup>
Vibrational frequencies (cm <sup>-1</sup> )	808, 160, 757
Electronic states (cm <sup>-1</sup> ) <sup>b</sup>	0(1), 17909(2), 18933(2), 22220(2), 23569(2), 24637(2)

<sup>a</sup>Product of moments of inertia.<sup>b</sup>Numbers in parentheses represent the statistical weights.

obtained from infrared spectroscopy. Gabelnick *et al.*,<sup>212</sup> Zhou and Andrews,<sup>458</sup> and Andrews *et al.*<sup>459</sup> have studied the infrared spectra of ThO<sub>2</sub> isotopomers in argon and neon matrices. On the basis of <sup>18</sup>O isotope shifts a nonlinear structure has been derived for ThO<sub>2</sub> with the bond angle (122.5 ± 2)°. Quantum-chemical calculations at DFT (Density Functional Theory),<sup>458–460</sup> Dirac-Hartree-Fock,<sup>461</sup> and SO-CASPT2 level<sup>462</sup> resulted in a bent structure for ThO<sub>2</sub> as well. The quantum chemical calculations resulted in Th–O bond lengths between 188.1–192.3 pm. The probably most accurate bond distance was suggested on the basis of the frequency–bond distance relationship derived by Kovács and Konings<sup>460</sup> utilizing the experimental matrix-IR data. The product of the principal moments of inertia of ThO<sub>2</sub> (see Table 65) was calculated using the values  $r_e(\text{Th–O}) = (191 \pm 5)$  pm and O–Th–O = (122.5 ± 2)°. The  $r_e(\text{Th–O})$  value is an average of three bond distances (190.6, 191.1, and 189.2 pm) obtained in the mentioned theoretical calculations.

The accepted vibrational frequencies  $\nu_1$  and  $\nu_3$  (see Table 65) were taken from IR spectra obtained by Gabelnick *et al.*,<sup>212</sup> Zhou and Andrews,<sup>458</sup> and Andrews *et al.*<sup>459</sup> for the neon matrix. ThO<sub>2</sub> bands detected in the Ne matrix are blue-shifted 22 cm<sup>-1</sup> from the solid argon data. In comparison with Ar, the Ne matrix is less polarizable, and matrix shifts in this case are substantially less. The deformation frequency  $\nu_2$  was estimated from calculations by Zhou and Andrews,<sup>458</sup> Dyall,<sup>461</sup> Andrews *et al.*<sup>459</sup> and Kovács and Konings.<sup>460</sup>

No experimental investigations of the electronic spectra of ThO<sub>2</sub> were reported. Quantum chemical calculations on the molecule agree in the X<sup>1</sup>A<sub>1</sub> ground state.<sup>458–463</sup> This state corresponds to the closed shell configuration and no low-lying electronic states may be expected for this molecule. The high electronic energy levels of ThO<sub>2</sub> given in Table 65 are estimated to be the same as for isoelectronic UO<sub>2</sub><sup>2+</sup> from *ab initio* calculations by Pierloot *et al.*<sup>464</sup>

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (285.233 \pm 2.0) \text{ JK}^{-1} \text{ mol}^{-1}.$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{JK}^{-1} \text{ mol}^{-1}) &= 36.83878 + 55.95391 \times 10^{-3} (\text{T/K}) \\ &\quad - 56.33462 \times 10^{-6} (\text{T/K})^2 + 20.31019 \\ &\quad \times 10^{-9} (\text{T/K})^3 - 1.940496 \\ &\quad \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

for the 298.15–900 K range, and

$$\begin{aligned} C_p^\circ / (\text{JK}^{-1} \text{ mol}^{-1}) &= 56.72155 + 2.163537 \times 10^{-3} (\text{T/K}) \\ &\quad - 1.189606 \times 10^{-6} (\text{T/K})^2 \\ &\quad + 0.2274986 \times 10^{-9} (\text{T/K})^3 \\ &\quad - 1.407937 \times 10^6 (\text{T/K})^{-2} \end{aligned}$$

for the 900–4000 K range.

## 6.2.2. Enthalpy of formation

The value of ThO<sub>2</sub>(g) enthalpy of formation is based on Langmuir and Knudsen effusion measurements of ThO<sub>2</sub>(cr) enthalpy of sublimation and mass spectrometric measurements of equilibrium constants for isomolecular oxygen exchange reactions (see Table 66). In the early Langmuir and Knudsen effusion measurements, the weight loss of thoria samples was used for calculation of the vapor pressure under assumption of ThO<sub>2</sub>(g) as the only thorium-bearing vapor species.<sup>465–467</sup> Knudsen effusion studies were carried out using substantially inert tungsten effusion cells. The only exception is the work by Hoch and Johnston<sup>466</sup> in which a tantalum cell was used. Ackermann *et al.*<sup>468</sup> have performed a combined study of the thorium-oxygen system evaporation behavior in the temperature range 2000–3000 K. It was found that solid thorium dioxide evaporates congruently. Above 2800 K, a thermodynamically insignificant substoichiometry ThO<sub>1.998</sub> was found. A combination of mass effusion and mass-spectrometric examination of the vapor composition has led to the conclusion that ThO and ThO<sub>2</sub> gaseous species are of comparable importance in the thoria vapor. More detailed study of thermodynamics of thoria vaporization has been carried out by Ackermann and

TABLE 66. The enthalpy of sublimation of ThO<sub>2</sub>(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Δ <sub>sub</sub> H°(298.15 K)
Shapiro <sup>465</sup>	L	2062–2257	791.3 ± 15.0
Hoch and Johnston <sup>466</sup>	K	2389–2676	(730.6)
Darnell and McCollum <sup>467</sup>	K	2268–2593	788.7 ± 12.0
Ackermann <i>et al.</i> <sup>468</sup>	K+M	2180–2871	787.9 ± 10.0
Shchukarev and Semenov <sup>472</sup>	K+M	2600–3000	802.9 ± 15.0
Semenov <sup>473b</sup>	K+M	2200–2700	779.0 ± 12.0
Ackermann and Rauh <sup>469c</sup>	K+M	2400–2800	797.1 ± 10.0
Hildenbrand and Murad <sup>470d</sup>	M	2160–2176	782.1 ± 15.0
Ackermann and Rauh <sup>471e</sup>	M	2320–2650	797.1 ± 15.0
Selected value:			790.8 ± 12.0

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry; L = Langmuir effusion.<sup>b</sup>Extended and reassessed data of Shchukarev and Semenov.<sup>472</sup><sup>c</sup>Recalculated from the Ackermann *et al.*<sup>468</sup> total pressure of Th-bearing species.<sup>d</sup>Calculated from the enthalpy of the gaseous reaction Th + ThO<sub>2</sub> = 2ThO.<sup>e</sup>Calculated from the enthalpy of the gaseous reaction ZrO<sub>2</sub> + ThO = ThO<sub>2</sub> + ZrO.

Rauh.<sup>469</sup> The congruently vaporizing composition of thoria was found  $\text{ThO}_{1.994}$ , instead of previous value  $\text{ThO}_{1.998}$ . Using the reassessed results of their previous investigation,<sup>468</sup> Ackermann and Rauh<sup>469</sup> have found equations for the temperature dependence of the  $\text{ThO}(\text{g})$  and  $\text{ThO}_2(\text{g})$  partial pressures over stoichiometric thoria. As follows from these equations,  $p(\text{ThO})/p(\text{ThO}_2)$  value linearly rises from 0.20 to 0.36 in the temperature range 2000–2800 K.

The enthalpy of sublimation values presented in Table 66 were obtained using partial pressures of  $\text{ThO}_2(\text{g})$  species calculated from the total vapor pressure over slightly substoichiometric thoria. All values of the enthalpy of sublimation are in good agreement. The only exception is the work by Hoch and Johnston.<sup>466</sup> Much higher values of weight loss found in this work are obviously connected with use of tantalum effusion cells for vaporization of thoria, leading to intensive formation of  $\text{ThO}(\text{g})$ . The enthalpy of sublimation values calculated from the results of mass-spectrometric measurements of gaseous equilibria<sup>470,471</sup> are in agreement with Knudsen effusion and Langmuir measurements.

The selected value is taken as a rounded weighted average from all works except Hoch and Johnston<sup>466</sup>:  $\Delta_{\text{sub}}H^\circ(298.15 \text{ K}) = (790.8 \pm 12.0) \text{ kJ mol}^{-1}$ . This value corresponds to:

$$\Delta_f H^\circ(\text{ThO}_2, \text{g}, 298.15) = -(435.6 \pm 12.6) \text{ kJ mol}^{-1}.$$

The corresponding energy of atomization is  $D_0(\text{ThO}_2) = (1531.3 \pm 13.5) \text{ kJ mol}^{-1}$ .

### 6.3. ThO(g)

#### 6.3.1. Heat capacity and entropy

The thermal functions of  $\text{ThO}(\text{g})$  in the standard state have been calculated using the molecular constants presented in Table 67.

The spectroscopic properties of ThO were studied since the beginning of the last century. The data published up to 1975 were reviewed by Huber and Herzberg<sup>179</sup> and up to 1980 by Gurvich *et al.*<sup>180</sup> Later Edvinsson and Lagerqvist<sup>474–479</sup>, Edvinsson and Jonsson,<sup>480</sup> published a number of papers concerning the ThO emission spectrum. The accurate Th-O

TABLE 67. Molecular constants of  $^{232}\text{Th}^{16}\text{O}(\text{g})$

No.	State	$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$	$r_e$ pm	$p_i$
0 <sup>a</sup>	$X^1 \Sigma_0^+$	0	895.77	2.39	0.33246	1.3	1.83 <sup>b</sup>	184.018613(24)	1
1 <sup>a</sup>	(1)3	5335.9							2
2 <sup>a</sup>	(1)2	6146.6							2
3 <sup>a</sup>		9315							4
4 <sup>a</sup>	[10.6]0	10626							2 <sup>c</sup>
5 <sup>a</sup>	[11.1]1	11156							2
6 <sup>a</sup>	[14.5]1	14525							4 <sup>c</sup>
7 <sup>a</sup>	[15.9]1	15975							3 <sup>c</sup>
8 <sup>a</sup>	[16.3]0	16354							1
9 <sup>a</sup>	[18.1]2	18053							2
10 <sup>a</sup>	[18.4]0	18406							1
11 <sup>a</sup>	[19.1]0	19068							1
12 <sup>a</sup>	[19.5]1	19586							2
13 <sup>a</sup>	[20.1]3	20112							7 <sup>c</sup>
14 <sup>a</sup>	[21.7]1	21756							6 <sup>c</sup>
15 <sup>a</sup>	[22.6]1	22685							8 <sup>c</sup>
16 <sup>a</sup>	[23.1]0	23155							1
17 <sup>a</sup>	[23.2]0	23199							2
18 <sup>a</sup>	[24.0]3	24084							4 <sup>c</sup>
19 <sup>a</sup>	[24.9]1	24886							6 <sup>c</sup>
20 <sup>a</sup>	[25.1]1	25184							3 <sup>c</sup>
21 <sup>a</sup>	[25.8]1	25850							8 <sup>c</sup>
22 <sup>a</sup>	[27.7]1	27756							6 <sup>c</sup>
23 <sup>a</sup>	[28.0]0	28071							2 <sup>c</sup>
24 <sup>a</sup>	[28.3]2	28360							2
25 <sup>a</sup>	[28.6]1	28614							2
26 <sup>a</sup>	[30.3]1	30346							14 <sup>c</sup>
27 <sup>a</sup>	[32.7]2	32770							10 <sup>c</sup>
28 <sup>d</sup>		35000							14
29 <sup>d</sup>		40000							20

<sup>a</sup>Experimental state.

<sup>b</sup> $D_0$ .

<sup>c</sup>Added statistical weights of near-lying predicted states.

<sup>d</sup>Estimated state.

TABLE 68. The enthalpy of formation of ThO(g), in kJ mol<sup>-1</sup>

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ackermann <i>et al.</i> <sup>468</sup>	K+M	2337–2381	ThO <sub>2</sub> (cr) = ThO(g) + O(g)	1434.4	-41.2
Shchukarev and Semenov <sup>472</sup>	M	2573–2973	ThO <sub>2</sub> (cr) = ThO(g) + O(g)	1393.2	-82.3
Ackermann and Rauh <sup>469</sup>	M	2400–2800	ThO <sub>2</sub> (cr) = ThO(g) + O(g)	1447.6	-28.0
	K	2080–2214	$\frac{1}{2}\text{Th}(\text{cr}) + \frac{1}{2}\text{ThO}_2(\text{cr}) = \text{ThO}(\text{g})$	599.8	-13.4
	M	1930–2280	ThO(g) + Y(g) = Th(g) + YO(g)	-169.5	-29.1
Hildenbrand and Murad <sup>470</sup>	M	1782–1927	$\frac{1}{2}\text{Th}(\text{cr}) + \frac{1}{2}\text{ThO}_2(\text{cr}) = \text{ThO}(\text{g})$	593.1	-20.1
	M	2064–2176	ThO(g) + Si(g) = Th(g) + SiO(g)	77.3	-24.1
Ackermann and Rauh <sup>469</sup>	M	2200–2550	ThO(g) + Hf(g) = Th(g) + HfO(g)	73.5	-21.4
Neubert and Zmbov <sup>496</sup>	M	1759–1961	ThO(g) + La(g) = Th(g) + LaO(g)	62.8	-11.7
Murad and Hildenbrand <sup>497</sup>	M	2288	ThO(g) + Zr(g) = Th(g) + ZrO(g)	110.9	-24.4
Selected value:			$D_0(\text{ThO}) = 868.6 \pm 5.0$		-21.5 ± 5.0

<sup>a</sup>K = Knudsen effusion; M = mass spectrometry.

bond distance in the ground-state molecule is known from a recent microwave spectroscopic study.<sup>481</sup> The harmonic vibrational frequency and other rotational and vibrational constants given in Table 67 were obtained from rotationally-resolved electronic emission spectra.<sup>474</sup> The IR measurements of ThO isolated in low-temperature inert gas matrices<sup>212,458,459,482,483</sup> revealed the matrix-shift effects on the fundamental frequency.

At present, 27 states (see Table 67) classified as belonging to Hund's coupling case "c" are rotationally analyzed, and 25 of them are arranged in the term scheme by Edvinsson *et al.*<sup>474–479,484</sup> The analyses by Behere *et al.*<sup>485,486</sup> are not convincing. The ground state of the ThO molecule is  $X0_0^+$  ( $X^1\Sigma_0^+$ ), as confirmed experimentally by the studies of the electronic spectrum of ThO<sup>179,180,474–480</sup> and also in the numerous theoretical calculations by Paulovic *et al.*,<sup>487,488</sup> Watanabe and Matsuoka,<sup>489–491</sup> Kühle *et al.*,<sup>492</sup> Kaledin *et al.*,<sup>258</sup> Marian *et al.*,<sup>493</sup> Seijo *et al.*,<sup>494</sup> Andrews *et al.*,<sup>459</sup> Goncharov and Heaven,<sup>484</sup> Buchachenko,<sup>495</sup> and Infante *et al.*<sup>462</sup>

Kaledin *et al.*<sup>258</sup> carried out the Ligand field calculations and assigned all 17 states observed till that time to the electron configurations  $7s^2$ ,  $6d7s$ ,  $7s7p$ , and  $5f7s$ , and calculated unobserved states of these and  $6d^2$  configurations. The calculations showed that the lowest unobserved states are  $\Omega = 3$  (probably W-state) and  $\Omega = 2$  (probably lower state of the band at 5579 Å) of the  $6d7s$  configuration. In the present work the statistical weights of predicted but unobserved states are attributed to the near-lying experimental states for simplification of the input data. The  $5f6d$ ,  $6d7p$ , and  $5f7p$  states are taken into account in the interval 25000 - 45000 cm<sup>-1</sup>.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (240.071 \pm 0.03) \text{ J K}^{-1} \text{ mol}^{-1},$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & 30.10768 + 13.9388 \times 10^{-3}(\text{T/K}) \\ & - 11.5407 \times 10^{-6}(\text{T/K})^2 \\ & + 4.92964 \times 10^{-9}(\text{T/K})^3 \\ & - 1.86435 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1500 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & -13.55069 + 44.5401 \times 10^{-3}(\text{T/K}) \\ & - 8.99062 \times 10^{-6}(\text{T/K})^2 + 0.583589 \\ & \times 10^{-9}(\text{T/K})^3 + 1.48587 \\ & \times 10^7(\text{T/K})^{-2} \end{aligned}$$

for the 1500–4000 K range.

### 6.3.2. Enthalpy of formation

The results of the determination of the enthalpy of formation of ThO(g) are presented in Table 68. The value is based on Knudsen effusion and mass spectrometric studies of ThO<sub>2</sub>(cr) and Th(cr) + ThO<sub>2</sub>(cr) system evaporation<sup>468–470,472</sup> and mass spectrometric measurements of isomolecular oxygen exchange reactions.<sup>469,470,496,497</sup> With the exception of the results of Ackermann *et al.*<sup>468</sup> and Shchukarev and Semenov,<sup>472</sup> all values of the enthalpy of formation of ThO(g) are in good agreement. The selected value is taken as a rounded average from all sources except the two mentioned papers:

$$\Delta_f H^\circ(\text{ThO, g, 298.15K}) = -(21.5 \pm 5.0) \text{ kJ mol}^{-1}.$$

This value corresponds to  $D_0(\text{ThO}) = (868.6 + 5.0) \text{ kJ mol}^{-1}$ .

## 6.4. PaO<sub>2</sub>(g)

### 6.4.1. Heat capacity and entropy

The thermodynamic functions of PaO<sub>2</sub>(g) in the standard state have been calculated using the data given in Table 69. Experimental data on molecular structure and spectra of PaO<sub>2</sub> are not available. Recently, the molecular geometry, vibrational properties and low-lying electronic states have been computed by Infante *et al.*,<sup>462</sup> and Kovács and Konings,<sup>460</sup> Kovács *et al.*,<sup>498</sup> and Kovács.<sup>456</sup> The calculations agree on a linear structure of  $D_{\infty h}$  symmetry in the ground electronic state, similar to all AnO<sub>2</sub> molecules (except ThO<sub>2</sub> which is bent). The moment of inertia has been evaluated using the SO-CASPT2 bond distance of 181.6 pm of the linear molecule Infante *et al.*<sup>462</sup> The vibrational frequencies were taken from Ref. 460.



TABLE 69. The molecular parameters for PaO<sub>2</sub>(g)

Parameter	Value
Ground electronic state	<sup>2</sup> Σ <sub>1/2g</sub>
Symmetry group	D <sub>∞h</sub>
Symmetry number, σ	2
I (g cm <sup>2</sup> ) <sup>a</sup>	17.52 × 10 <sup>-39</sup>
Vibrational frequencies (cm <sup>-1</sup> ) <sup>b</sup>	852, 82(2), 899
Electronic states (cm <sup>-1</sup> ) <sup>c</sup>	0(2), 3961(2), 8252(2), 8924(2), 11290(2), 14430(2), 15488(2), 27933(2), 28075(2), 28617(2), 29739(2), 29914(2)

<sup>a</sup>Moment of inertia.<sup>b</sup>Numbers in parentheses represent degeneracy.<sup>c</sup>Numbers in parentheses represent the statistical weights.

The electronic ground state of PaO<sub>2</sub> is <sup>2</sup>Σ<sub>1/2g</sub>.<sup>462</sup> The vertical excitation energies to the low-lying states were calculated recently at the SO-CASPT2 level by Kovács.<sup>456</sup> The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (279.711 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 35.4855 + 69.7567 \times 10^{-3}(\text{T/K}) \\ &\quad - 71.6190 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 27.8213 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 0.58791 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–900 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 57.8335 + 4.88325 \times 10^{-3}(\text{T/K}) \\ &\quad + 6.76950 \times 10^{-8}(\text{T/K})^2 \\ &\quad - 1.07986 \times 10^{-10}(\text{T/K})^3 \\ &\quad - 1.40951 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 900–4000 K range.

#### 6.4.2. Enthalpy of formation

Kleinschmidt and Ward<sup>499</sup> have measured partial pressures of Pa(g), PaO(g), and PaO<sub>2</sub>(g) above the Pa(l) + PaO<sub>2-y</sub>(cr) two-phase system and the congruently vaporizing composition PaO<sub>2-x</sub> (*x* and *y* values were not specified). From the latter data Kleinschmidt and Ward calculated the enthalpy of sublimation Δ<sub>sub</sub>H<sup>o</sup>(PaO<sub>2</sub>, cr, 298.15 K) = (595 ± 7) kJ mol<sup>-1</sup> as an average of the second- and third-law values (thermodynamic functions of all three gaseous species were estimated; however, molecular constants and numerical values of functions were not presented in the paper). The enthalpy of formation for PaO<sub>2</sub>(g) was derived by Kleinschmidt and Ward assuming the enthalpy of formation of solid substoichiometric protactinium dioxide equal to that for PaO<sub>2</sub>(cr). Following this suggestion, we obtain for the enthalpy of formation Δ<sub>f</sub>H<sup>o</sup>(298.15 K) = -(514 ± 17) kJ mol<sup>-1</sup>.

To acquire some insight into reliability of the enthalpy of formation of PaO<sub>2</sub>(g) obtained with indeterminate thermo-

dynamic functions, we calculated the third-law enthalpy of sublimation using the thermodynamic properties for a related neighbor compound, uranium dioxide, adopted in this work: Δ<sub>sub</sub>H<sup>o</sup>(298.15 K) = 619 kJ mol<sup>-1</sup>. The 24 kJ mol<sup>-1</sup> difference between two enthalpy of sublimation values can serve as an estimation of possible uncertainty in the enthalpy of formation of PaO<sub>2</sub>(g) found by Kleinschmidt and Ward.<sup>499</sup> On the basis of calculations performed, we select the value derived from the paper by Kleinschmidt and Ward<sup>499</sup> as a provisional recommendation, with an extended uncertainty range

$$\Delta_f H^\circ(\text{PaO}_2, \text{g}, 298.15) = -(514.0 \pm 30) \text{ kJ mol}^{-1}.$$

To the selected PaO<sub>2</sub>(g) enthalpy of formation corresponds the atomization energy value D<sub>0</sub>(PaO<sub>2</sub>) = (1576.0 ± 30) kJ mol<sup>-1</sup>.

## 6.5. PaO(g)

### 6.5.1. Heat capacity and entropy

The thermal functions of PaO(g) in the standard state have been calculated using the data given in Table 70. The molecular constants were estimated using the trends revealed in the spectral data of the lanthanide monoxides as there is no spectral data on the PaO molecule. The electronic structure is assumed to be analogous to that of PrO. The energies of the low-lying states taken from the PrO data are approximated to hundreds cm<sup>-1</sup>.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (250.778 \pm 10.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 22.73634 + 18.0345 \times 10^{-3}(\text{T/K}) \\ &\quad + 28.8250 \times 10^{-6}(\text{T/K})^2 - 20.1289 \\ &\quad \times 10^{-9}(\text{T/K})^3 \\ &\quad + 2.78485 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

TABLE 70. Molecular constants of <sup>231</sup>Pa<sup>16</sup>O(g)

No.	State	cm <sup>-1</sup>						r <sub>e</sub> pm	p <sub>i</sub>
		T <sub>e</sub>	ω <sub>e</sub>	ω <sub>e</sub> x <sub>e</sub>	B <sub>e</sub>	α <sub>e</sub> 10 <sup>3</sup>	D <sub>e</sub> 10 <sup>7</sup>		
0	X <sup>8</sup> Σ	0	845	2.69	0.335	1.5	2.1	183.5	2
1			200						2
2			2000						8
3			3000						12
4			4000						8
5			5500						18
6			7000						8
7			10 000						85
8			15 000						150
9			20 000						180
10			25 000						185
11			30 000						190
12			35 000						195
13			40 000						215

for the 298.15–1100 K range, and

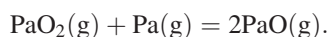
$$C_p^{\circ}/(\text{J K}^{-1} \text{ mol}^{-1}) = 84.42095 - 29.0423 \times 10^{-3}(\text{T/K}) \\ + 8.96146 \times 10^{-6}(\text{T/K})^2 - 0.876655 \\ \times 10^{-9}(\text{T/K})^3 - 1.36244 \\ \times 10^7(\text{T/K})^{-2}$$

for the 1100–4000 K range.

### 6.5.2. Enthalpy of formation

Kleinschmidt and Ward<sup>499</sup> have measured the partial pressures of Pa(g), PaO(g), and PaO<sub>2</sub>(g) above the Pa(l) + PaO<sub>2-y</sub>(cr) two-phase system and the congruently vaporizing composition PaO<sub>2-x</sub> (*x* and *y* values were not specified). From this data, the third-law enthalpy of formation of PaO<sub>2</sub>(g) was found as explained above and the enthalpy of formation of Pa(g). Thermodynamic functions of all three gaseous species were estimated. However, neither molecular constants nor numerical values of functions have been presented in the paper.

The vapor pressure measurements in the two-phase region Pa(l) + PaO<sub>2-y</sub>(cr) resulted in the value  $\Delta_f H^{\circ}(298.15 \text{ K}) = -(18 \pm 13) \text{ kJ mol}^{-1}$  for the reaction



Combining of this enthalpy of reaction with the thermochemical values for PaO<sub>2</sub>(g) and Pa(g), leads to the enthalpy of formation of PaO(g):

$$\Delta_f H^{\circ}(\text{PaO}, \text{g}, 298.15 \text{ K}) = (8 \pm 30) \text{ kJ mol}^{-1}$$

This value corresponds to  $D_0(\text{PaO}) = (789 \pm 30) \text{ kJ mol}^{-1}$ . This is in good agreement with the value  $(801 \pm 59) \text{ kJ mol}^{-1}$  estimated by Marçalo and Gibson<sup>500</sup> on the basis of gas phase oxidation reactions involving the ionised species and known ionisation energies.

## 6.6. UO<sub>3</sub>(g)

### 6.6.1. Heat capacity and entropy

The thermodynamic functions of UO<sub>3</sub>(g) in the standard state have been calculated using the data given in Table 71. Experimental data for the structure of free gaseous UO<sub>3</sub> molecule does not exist in the literature. From analysis of the infrared spectra of matrix-isolated UO<sub>3</sub> and its oxygen-18 isotopomers, Gabelnick *et al.*<sup>501</sup> and Green *et al.*<sup>502</sup> have determined a planar T-shaped (*C*<sub>2v</sub>) geometry for the UO<sub>3</sub> molecule, with a O–U–O linear fragment. Three low-frequency and two stretching modes observed in matrix-isolated UO<sub>3</sub> are inconsistent with a pyramidal (*C*<sub>3v</sub>) or planar (*D*<sub>3h</sub>) geometry. The investigation of infrared spectra of UO<sub>3</sub> in solid argon by Hunt and Andrews<sup>503</sup> is in agreement with the data of Gabelnick *et al.*<sup>501</sup> and Green *et al.*<sup>502</sup> The results of *ab initio* calculations by Pyykkö *et al.*<sup>504</sup> support the symmetry assignment from experiments of Gabelnick *et al.*<sup>501</sup> and Green *et al.*<sup>502</sup> and Hunt and Andrews<sup>503</sup>; it has been shown

that *D*<sub>3h</sub> structure is a saddle point, 49 kJ mol<sup>-1</sup> above the *C*<sub>2v</sub> minimum. Pyykkö *et al.*<sup>504</sup> obtained  $\angle\text{O–U–O} = 161^{\circ}$ , instead of the linear O–U–O fragment found by Gabelnick *et al.*<sup>501</sup> and Green *et al.*<sup>502</sup> Similar results have been obtained later by Privalov *et al.*<sup>505</sup> using a similar theoretical (SCF) level. Quantum chemical calculations at the more reliable B3LYP DFT level were performed by Zhou *et al.*<sup>506</sup> for the geometry and vibrational frequencies. They retain the main structural features obtained by Pyykkö *et al.*<sup>504</sup> and by Privalov *et al.*,<sup>505</sup> but with reduced difference between the shorter U–O (in the O–U–O fragment) and the longer U–O' bonds. The product of the principal moments of inertia of UO<sub>3</sub> (see Table 71) is calculated with the selected molecular constants  $r_e(\text{U–O}) = (181 \pm 5) \text{ pm}$ ,  $r_e(\text{U–O}') = (185.3 \pm 5) \text{ pm}$ , and  $\angle\text{O–U–O} = (158.8 \pm 5)^{\circ}$  from the DFT calculations by Zhou *et al.*<sup>506</sup>

Experimental data on vibrational frequencies were obtained by IR spectroscopy in argon matrices,<sup>501–503</sup> and in neon matrix.<sup>506</sup> With the exception of  $\nu_1$ , all vibrational modes were observed. The accepted vibrational frequencies  $\nu_2$  and  $\nu_4$  (see Table 71) were taken from the IR spectra obtained by Zhou *et al.*<sup>506</sup> in an neon matrix, ensuring minimal matrix shift of the frequencies. The low frequencies  $\nu_3$ ,  $\nu_5$ , and  $\nu_6$  were accepted from infrared spectra obtained by Green *et al.*<sup>502</sup> The unobserved  $\nu_1$  frequency is accepted as an average of two values obtained in calculations of Green *et al.*<sup>502</sup> and Zhou *et al.*<sup>506</sup>

The experimental and theoretical investigations of electronic spectra of UO<sub>3</sub> are unknown. *X*<sup>1</sup>*A*<sub>1</sub> ground state is accepted for UO<sub>3</sub> molecule from calculations of Zhou *et al.*<sup>506</sup> This state corresponds to the closed shell configuration and no low-lying electronic states may be expected for this molecule. Additional electronic energy levels of UO<sub>3</sub> given in Table 71 are accepted to be the same as for the isoelectronic UO<sub>2</sub><sup>2+</sup> from *ab initio* calculations by Pierloot *et al.*<sup>464</sup>

The derived standard entropy at room temperature is

$$S^{\circ}(298.15 \text{ K}) = (310.648 \pm 3) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$C_p^{\circ}/(\text{J K}^{-1} \text{ mol}^{-1}) = 46.69199 + 94.69427 \times 10^{-3}(\text{T/K}) \\ - 94.91701 \times 10^{-6}(\text{T/K})^2 + 34.1585 \\ \times 10^{-9}(\text{T/K})^3 - 2.793843 \\ \times 10^5(\text{T/K})^{-2}$$

TABLE 71. The molecular parameters for UO<sub>3</sub>(g)

Parameter	Value
Ground electronic state	<i>X</i> <sup>1</sup> <i>A</i> <sub>1</sub>
Symmetry group	<i>C</i> <sub>2v</sub>
Symmetry number, $\sigma$	2
$I_A I_B I_C$ (g <sup>3</sup> cm <sup>6</sup> ) <sup>a</sup>	$4210 \times 10^{-117}$
Vibrational frequencies (cm <sup>-1</sup> )	860, 760, 186, 865, 212, 152
Electronic states (cm <sup>-1</sup> ) <sup>b</sup>	0(1), 17909(2), 18933(2), 22220(2), 23569(2), 24637(2)

<sup>a</sup>Product of moments of inertia.

<sup>b</sup>Numbers in parentheses represent the statistical weights.

for the 298.15–900 K range, and

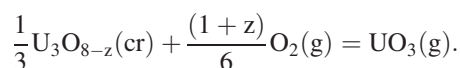
$$\begin{aligned} C_p^{\circ}/(\text{JK}^{-1} \text{mol}^{-1}) = & 81.70962 + 2.009478 \times 10^{-3}(\text{T/K}) \\ & - 1.110505 \times 10^{-6}(\text{T/K})^2 \\ & + 0.2162739 \times 10^{-9}(\text{T/K})^3 \\ & - 2.580355 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 900–4000 K range.

### 6.6.2. Enthalpy of formation

For determination of the enthalpy of formation of  $\text{UO}_3(\text{g})$  the results of transpiration and mass-spectrometric measurements were used. Transpiration experiments with  $\text{U}_3\text{O}_8$  with oxygen employed as a carrier gas were carried out by Ackermann *et al.*,<sup>507</sup> Alexander,<sup>508</sup> Dharwadkar *et al.*,<sup>509</sup> Krikorian *et al.*,<sup>510</sup> and Alexander.<sup>511</sup> In all works except of Alexander,<sup>511</sup> pure oxygen at 1 atm was used as carrier gas; in the latter work dry air at 1 atm total pressure was employed. It was shown by Ackermann *et al.*<sup>507</sup> that variation of the oxygen pressure in the carrier gas established the oxygen-uranium ratio equal to three for the volatile uranium oxide molecule. By a comparison with entropies of sublimation of molybdenum and tungsten trioxides it was inferred that gaseous monomeric uranium trioxide is the principal species produced by the reaction of  $\text{U}_3\text{O}_8$  with oxygen. To mention is that, unlike  $(\text{MoO}_3)_n$  and  $(\text{WO}_3)_n$ ,  $(\text{UO}_3)_n$  gaseous associates till now are not detected in the U-O system.

Taking into account that the  $\text{U}_3\text{O}_8$  phase has a homogeneity range, and that at high temperatures some deviation from stoichiometry develops, the process of transpiration has to be described according to the equation



Detailed thermodynamic characterization of the  $\text{U}_3\text{O}_{8-z}$  phase was carried out by Ackermann and Chang.<sup>512</sup> The  $\text{U}_3\text{O}_{8-z}$  composition equilibrated at controlled temperature and oxygen pressure values was experimentally studied in this work. The equilibrium composition of the  $\text{U}_3\text{O}_{8-z}$  phase becomes progressively substoichiometric with increasing temperature at constant oxygen pressure. At the highest temperature of experiments, 1445 K, and the oxygen pressure 1 atm, the initial composition  $\text{UO}_{2.667}$  changes to  $\text{UO}_{2.636}$ . In air, at an oxygen pressure  $\approx 0.2$  atm, the composition changes to  $\approx 2.627$ . In spite of these findings, we adopted a simplified approach in the treatment of the transpiration data, without using information on deviation from stoichiometry. This approach is justified by a weak dependence of the free energy of formation of  $\text{U}_3\text{O}_{8-z}(\text{cr})$  on the  $z$  value at constant temperature, as was shown by Ackermann and Chang.<sup>512</sup> This feature becomes apparent in absence of any systematic temperature drift of the third-law enthalpy values calculated during our assessment of experimental data for the reaction

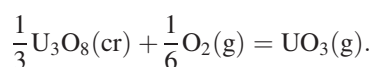


TABLE 72. The enthalpy of formation of  $\text{UO}_3(\text{g})$ , in  $\text{kJ mol}^{-1}$

Authors	Method <sup>a</sup>	T/K	$\Delta_f H^{\circ}(298.15 \text{ K})$
DeMaria <i>et al.</i> <sup>513</sup>	M	2200–2322	$-769.9 \pm 20$
Ackermann <i>et al.</i> <sup>507</sup>	T	1200–1370	$-799.5 \pm 6$
Alexander <sup>508</sup>	T	1415–1760	$-792.8 \pm 8$
Pattoret <i>et al.</i> <sup>515b</sup>	M	2080–2500	$-805.2 \pm 15$
Dharwadkar <i>et al.</i> <sup>509</sup>	T	1525–1675	$-797.2 \pm 6$
Younés <i>et al.</i> <sup>221</sup>	M	1916–2336	$-785.1 \pm 15$
Krikorian <i>et al.</i> <sup>510</sup>	T	1173–1573	$-789.3 \pm 10$
Alexander <sup>511</sup>	T	1410–1815	$-792.9 \pm 8$
Selected value:			$-795.0 \pm 10$

<sup>a</sup>M = mass spectrometry; T = transpiration.

<sup>b</sup>Results of treatment of experimental data from Drowart *et al.*<sup>514</sup>

Results of all investigations are summarized in Table 72. All transpiration experiments gave very close results, an average of five works being  $-794.3 \text{ kJ mol}^{-1}$ . Results of mass-spectrometric measurements<sup>221,513–515</sup> are close to this value but show more marked scatter. The most probable reason for that is the use of different estimates for the ionization cross sections of molecules and different estimates of fragmentation degree under electron impact, the latter being especially important for  $\text{UO}_3$  molecules. We have derived the selected value for  $\text{UO}_3(\text{g})$  enthalpy of formation as a weighted average of all results shown in Table 72 (except the results of DeMaria *et al.*<sup>513</sup> with the largest deviation from the mean):

$$\Delta_f H^{\circ}(\text{UO}_3, \text{g}, 298.15) = (-795.0 \pm 10) \text{ kJ mol}^{-1}.$$

## 6.7. $\text{UO}_2(\text{g})$

### 6.7.1. Heat capacity and entropy

The thermodynamic functions of  $\text{UO}_2(\text{g})$  in the standard state have been calculated using the data given in Table 73.

The investigations of infrared spectra of matrix-isolated uranium oxide species by Gabelnick *et al.*,<sup>501</sup> Lue *et al.*,<sup>516</sup> and Zhou *et al.*<sup>506</sup> strongly indicated a linear geometry of  $D_{\infty h}$  symmetry for  $\text{UO}_2$ . This is consistent with the absence in spectra of peaks attributable to  $\nu_1$  symmetric stretch for the  $\text{U}^{16}\text{O}_2$  and  $\text{U}^{18}\text{O}_2$  isotopomers. Consistent with the experiment results, theoretical calculations for  $\text{UO}_2$  predict a linear symmetric ground state.<sup>460,506,517–519</sup> There are no experimental data on U–O bond distances in  $\text{UO}_2(\text{g})$ . The principal moment of inertia of  $\text{UO}_2$  is calculated using  $r_e(\text{U–O}) = (178 \pm 5) \text{ pm}$ . This value is an average of five bond distances (177, 178.4, 179.5, 176.4, and 180 pm) obtained from *ab initio* and DFT calculations;<sup>506,517–520</sup> it is close to  $r(\text{U–O}) = 179 \text{ pm}$  estimated by Green *et al.*<sup>502</sup> and the value derived on the basis of the frequency-bond distance relationship by Kovács and Konings<sup>460</sup> utilizing an estimated “gas-phase” value for the asymmetric stretching fundamental (vide infra).

Experimental data for  $\nu_2$  bending and  $\nu_3$  asymmetric vibrations were obtained by infrared spectroscopy in noble gas matrices.<sup>501,503,506</sup> An interesting feature of  $\text{UO}_2$  is the large red shift (about  $130 \text{ cm}^{-1}$ ) in the asymmetric stretch found when replacing a neon matrix by an argon matrix. This feature has been explained and supported by computational and

TABLE 73. The molecular parameters for UO<sub>2</sub>(g)

Parameter	Value
Ground electronic state	<sup>3</sup> Φ <sub>2u</sub>
Symmetry group	D <sub>∞h</sub>
Symmetry number, σ	2
I (g cm <sup>2</sup> ) <sup>a</sup>	17.0 × 10 <sup>-39</sup>
Vibrational frequencies (cm <sup>-1</sup> ) <sup>b</sup>	860, 120(2), 915
Electronic states (cm <sup>-1</sup> ) <sup>c</sup>	0(2), 360(2), 2231(2), 2588(2), 5047(2), 6148(2), 6501(2), 7081(1), 7152(2), 7431(2), 7867(2), 8268(2), 8746(2), 10089(1), 10914(2), 11221(2), 11436(1), 11510(2), 12310(1), 12564(2), 12700(1), 12958(2), 13458(2), 13919(2), 14104(2), 14654(2), 14995(2), 15196(1), 15408(2), 15455(2), 15502(2), 15860(1), 15945(1), 16625(2), 16786(2), 16949(2), 17058(2), 17127(1), 17340(2), 17516(2), 17606(1), 18355(1), 18596(2), 18913(2), 19105(2), 19317(2), 19491(2), 21300(16), 24700(22), 30100(39)

<sup>a</sup>Moment of inertia.<sup>b</sup>Numbers in parentheses represent the degeneracies.<sup>c</sup>Numbers in parentheses represent the statistical weights.

further experimental studies<sup>506,521,522</sup> that the more polarizable argon and krypton stabilize the low-lying <sup>3</sup>H<sub>g</sub> state becoming the electronic ground state in these matrices. In neon the <sup>3</sup>Φ<sub>2u</sub> state found in the gaseous phase remains the electronic ground state.

The adopted value for ν<sub>3</sub> asymmetric vibrational mode of U<sup>16</sup>O<sub>2</sub> is taken from the investigation of the infrared spectra in solid neon matrix<sup>506</sup> because neon is substantially less polarizable than argon and matrix shifts in this case are substantially less. The value of infrared inactive ν<sub>1</sub> mode is calculated by normal coordinate analysis using the experimental data for ν<sub>3</sub> of UO<sub>2</sub>. These values are in excellent agreement with the recent assessment by Kovács and Konings,<sup>460</sup> who derived the gas-phase ν<sub>3</sub> value by correction of 5 cm<sup>-1</sup> for the matrix-shift of neon and deriving the ν<sub>1</sub> value from the frequency-bond distance relationship.

The bending frequency ν<sub>2</sub> = 120 cm<sup>-1</sup> was observed using the resonantly enhanced multiphoton ionization (REMPI) technique in the gas phase.<sup>523</sup> This bending frequency is lower than the values predicted by calculations. Majumdar *et al.*<sup>517</sup> reported ν<sub>2</sub> frequency values in the range of 149–222 cm<sup>-1</sup> using three different computational methods. DFT calculation by Zhou *et al.*<sup>506</sup> yielded 138 cm<sup>-1</sup>. The gas phase bending frequency is also lower than the ΔG<sub>1/2</sub> = 225.2 cm<sup>-1</sup> reported for UO<sub>2</sub> isolated in a solid Ar matrix by Green *et al.*<sup>502</sup> In spite of this discrepancy, the experimental value ν<sub>2</sub> = 120 cm<sup>-1</sup> derived for free UO<sub>2</sub> molecule by Han *et al.*<sup>523</sup> is selected in our work for calculations of the UO<sub>2</sub>(g) thermodynamic functions.

Thermodynamic functions have been calculated with the inclusion of data for a wide range of excited electronic states. Experimental data on the UO<sub>2</sub> excited energy levels are fragmentary.<sup>523</sup> Several theoretical studies of UO<sub>2</sub> were carried out at relativistic post-Hartree-Fock levels, reporting electronic energy levels.<sup>503,506,516–519,524</sup> From these diverse studies we adopted the results by<sup>518</sup> obtained using the Dirac-Coulomb intermediate Hamiltonian multireference coupled cluster approach (DC-IHFSCC). This method is superior to the ones applied in the other papers by (i) using the coupled cluster methods to account for electron correlation; (ii) including the

6d orbitals, that have so far been left out of the active space; (iii) including spin-orbit coupling effects by utilizing the 4-component Dirac-Coulomb Hamiltonian. From this study we took the excited electronic states with the energy up to 19491 cm<sup>-1</sup>. The energy levels above 19491 cm<sup>-1</sup> were united in combined levels with summed up statistical weights (see Table 73).

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (277.027 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 44.35744 + 37.63585 \times 10^{-3}(\text{T/K}) \\ &\quad - 23.15563 \times 10^{-6}(\text{T/K})^2 + 5.50268 \\ &\quad \times 10^{-9}(\text{T/K})^3 - 0.7485093 \\ &\quad \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1500 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 59.57586 + 5.392403 \times 10^{-3}(\text{T/K}) \\ &\quad + 0.09463181 \times 10^{-6}(\text{T/K})^2 \\ &\quad - 0.1028723 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 6.319451 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 1500–4000 K range.

## 6.7.2. Enthalpy of formation

The enthalpy of formation for UO<sub>2</sub>(g) is calculated from the enthalpy of sublimation of UO<sub>2</sub>(cr). The results of the determination of the enthalpy of sublimation of UO<sub>2</sub> are listed in Table 74. Mass-spectrometric investigations (see e.g., Pattoret *et al.*<sup>515</sup>) have shown that the vapor over congruently vaporizing uranium dioxide of slightly substoichiometric composition mainly consists of UO<sub>2</sub> molecules; UO and UO<sub>3</sub> molecules are present in the vapor with the total pressure of several % of UO<sub>2</sub>(g). To derive the selected value of the enthalpy of sublimation, the UO(g) and UO<sub>3</sub>(g) pressures were subtracted from the total pressure of uranium-bearing species.



TABLE 74. The enthalpy of sublimation of  $\text{UO}_2(\text{g})$ , in  $\text{kJ mol}^{-1}$ 

Authors	Method <sup>a</sup>	T/K	$\Delta_{\text{sub}}H^\circ(298.15 \text{ K})$
Ackermann <i>et al.</i> <sup>531</sup>	K	1600–2200	628.2 ± 10
Ivanov <i>et al.</i> <sup>532</sup>	b	1930–2160	623.4 ± 10
Voronov <i>et al.</i> <sup>525</sup>	L	1723–2573	641.9 ± 10/-20
Ohse <sup>533</sup>	K	2278–2768	622.5 ± 10
Gorban' <i>et al.</i> <sup>534</sup>	K	1850–2600	618.9 ± 12
Pattoret <i>et al.</i> <sup>515</sup>	K+M	1890–2420	621.5 ± 8
Tetenbaum and Hunt <sup>535</sup>	T	2080–2705	629.3 ± 12
Reedy and Chasanov <sup>536</sup>	T	2615–3391	614.8 ± 18
Ackermann and Tetenbaum <sup>526</sup>	M	1540–2315	626.2 ± 15
Selected value:			622.9 ± 12

<sup>a</sup>K = Knudsen effusion; L = Langmuir; M = mass spectrometry; T = transpiration.

<sup>b</sup>Vaporization from a cylindrical crucible; results of weight-loss measurements are close to those of Knudsen effusion.

The values of  $p(\text{UO}) + p(\text{UO}_3)$  were calculated using thermodynamic data from Gurvich *et al.*<sup>180</sup>; they amounted from  $\approx 0.01$  p(total) at 1600 K to  $\approx 0.24$  p(total) at 3400 K.

The selected enthalpy of sublimation of  $\text{UO}_2(\text{g})$  is obtained from the data presented in Table 74 as a weighted average:  $\Delta_{\text{sub}}H^\circ(\text{UO}_2, \text{cr}, 298.15) = (622.9 \pm 12) \text{ kJ mol}^{-1}$ . The data of Voronov *et al.*<sup>525</sup> and of Ackermann and Tetenbaum<sup>526</sup> were not taken into account. In the former work, free evaporation of a uranium dioxide rod heated by electric current might lead to serious errors due to nonuniformity of the rod temperature and to a non unity evaporation coefficient of the substance. The result obtained by Ackermann and Tetenbaum<sup>526</sup> is formally close to the selected value. However, independent sensitivity calibration of the mass-spectrometric equipment was not carried out. Instead,  $p(\text{UO}_2)$  at the temperature 2050 K was taken equal to an averaged value of all published data.

Combination of the enthalpy of sublimation with the selected enthalpy of formation of  $\text{UO}_2(\text{cr})$  gives the selected value:

$$\Delta_f H^\circ(\text{UO}_2, \text{g}, 298.15) = -(462.1 \pm 12) \text{ kJ mol}^{-1}.$$

This value corresponds to an atomization energy  $D_0(\text{UO}_2, \text{g}) = (1486.8 \pm 15) \text{ kJ mol}^{-1}$ .

In the above analysis the data of laser heating experiments on uranium dioxide and other techniques (see, e.g., Ohse *et al.*,<sup>527</sup> Breitung and Reil,<sup>528</sup> Pflieger *et al.*<sup>529</sup>) were not discussed. We have preferred the data which allow unambiguous application of thermodynamics of ideal gases, and to avoid difficulties in extracting the  $\text{UO}_2(\text{g})$  partial pressure from the total pressure values at very high temperatures (up to 8000 K), complicated by the possibility of substantial deviation of the vapor from the ideal gas behavior. Nevertheless, extrapolation of our results into the region of extremely high temperatures shows satisfactory agreement with experimental data and results derived from the equation of state of uranium dioxide summarized by Fink.<sup>530</sup>

## 6.8. $\text{UO}(\text{g})$

### 6.8.1. Heat capacity and entropy

The thermal functions of  $\text{UO}(\text{g})$  in the standard state have been calculated using the data given in Table 75.

Electronic spectra of the uranium monoxide molecule were studied by Kaledin *et al.*<sup>537</sup> in absorption and emission, by Heaven *et al.*<sup>538</sup> at the laser excitation of jet cooled  $\text{UO}(\text{U}^{16}\text{O}$  and  $\text{U}^{18}\text{O})$ , by Kaledin *et al.*<sup>539</sup> Gurvich *et al.*<sup>540</sup> and Kaledin *et al.*<sup>258</sup> ( $\text{U}^{16}\text{O}$  and  $\text{U}^{18}\text{O}$ ) in fluorescence, and by Kaledin and Heaven<sup>541</sup> by means of resonantly enhanced two photon ionization with mass selected ion detection ( $\text{U}^{16}\text{O}$  and  $\text{U}^{18}\text{O}$ ). The studies gave information about the ground  $X\Omega = 4$  ( $v \leq 6$ ) state, 8 low-lying and about 20 higher-lying excited electronic states. The ground state  $\Delta G_{1/2}$  value was found to be  $882.351 \text{ cm}^{-1}$ . In addition, the bond distance was determined to be  $183.83 \text{ pm}$  by Kaledin *et al.*<sup>258</sup>

Infrared spectra of matrix-isolated uranium oxide species were investigated by Carstens *et al.*,<sup>542</sup> Gabelnick *et al.*,<sup>501</sup> Abramowitz and Acquista,<sup>543</sup> Hunt and Andrews,<sup>503</sup> and Zhou *et al.*<sup>506</sup> The bands observed near  $820 \text{ cm}^{-1}$  in the Ar and Kr matrices were assigned to the  $\text{UO}$  molecule. Gabelnick *et al.*<sup>501</sup> determined also the anharmonicity value. In the IR spectrum of uranium oxides isolated in Ne matrix the band assigned to  $\text{UO}$  was that at  $882.4 \text{ cm}^{-1}$ .<sup>506</sup> Most quantum chemical calculations<sup>257,460,506,544,545</sup> gave harmonic vibrational frequency values between 845–858 taking into account the anharmonicity and matrix-shift effects in good agreement with the Ar-matrix value. Compared to these values, however, the frequency measured in the Ne matrix is too high.

TABLE 75. Molecular constants of  $^{238}\text{U}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$					$r_e$ pm	$p_i$	
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$			$D_e 10^7$
0 <sup>a</sup>	X(1)4	0	888.5	3.1	0.3346	3.2	1.9 <sup>c</sup>	183.3	2
1 <sup>a</sup>	(2)4	294							2
2 <sup>a</sup>	(1)3	652							2
3 <sup>a</sup>	(1)2	958							2
4 <sup>a</sup>	(1)5	1043							2
5 <sup>b</sup>		1200							3
6 <sup>b</sup>		1500							2
7 <sup>a</sup>	(3)4	1574							2
8 <sup>a</sup>	(3)3	1941							2
9 <sup>b</sup>		2225							7 <sup>d</sup>
10 <sup>b</sup>		2450							3
11 <sup>b</sup>		4500							7 <sup>e</sup>
12 <sup>b</sup>		5200							9
13 <sup>b</sup>		5700							15
14 <sup>b</sup>		7500							38
15 <sup>b</sup>		10 000							116
16 <sup>b</sup>		12 500							238
17 <sup>b</sup>		15 000							375
18 <sup>b</sup>		20 000							615
19 <sup>b</sup>		25 000							1060
20 <sup>b</sup>		30 000							1125
21 <sup>b</sup>		35 000							1215
22 <sup>b</sup>		40 000							1610
23 <sup>b</sup>		45 000							1615

<sup>a</sup>Experimental state.

<sup>b</sup>Estimated state.

The reason of these discrepancies is that the ground state vibrational levels of gaseous UO are irregular. Kaledin and Kulikov,<sup>546</sup> Kaledin *et al.*<sup>547</sup> found that three lowest  $\Omega = 4$  states were mutually perturbed. Kaledin *et al.*<sup>258</sup> performed deperturbation of these levels to determine adiabatic molecular constants. The deperturbed constant  $\omega_e = 846.5 \text{ cm}^{-1}$  agreed well with the bands observed in IR spectra of matrix-isolated uranium oxides (taking into account the well-known red shift in matrices) and with theoretical predictions.

The information on the excited states is not sufficient for the calculation of the thermal functions. To estimate the electronic partition function we use the results of the Ligand field theory calculations by Kaledin *et al.*<sup>258</sup> All but one low-lying states were assigned to  $f^3s$  and the  $\Omega = 4$  state with energy  $294 \text{ cm}^{-1}$  to  $f^2s^2$  configurations. In the present work, the detailed experimental or calculated data on low-lying electronic states up to  $7500 \text{ cm}^{-1}$  are taken into account; the calculated data on the  $f^3s$ ,  $f^2s^2$ ,  $d^2s^2$ ,  $f^2ds$ ,  $f^3d$ ,  $f^2d^2$ ,  $f^2p^2$ ,  $f^2ps$ ,  $f^4$ ,  $fds^2$ , and  $f^2dp$  electronic states in the range  $7500\text{--}45\,000 \text{ cm}^{-1}$  are given in Table 75 as united terms with the fixed energies and the corresponding statistical weights.

The ground state molecular constants of UO selected for the thermal functions calculations (see Table 75) are estimated from the experimental value  $\Delta G_{1/2} = 882.351 \text{ cm}^{-1}$  and the adopted value of the dissociation energy; these constants describe correctly the  $v = 0$  and 1 levels. The simple presentation of the vibrational-rotational energy levels used in our program does not permit to recalculate levels from the deperturbed values of constants. Use of the deperturbed constants ( $\omega_e = 846.5 \text{ cm}^{-1}$  and  $\omega_e x_e = 2.3 \text{ cm}^{-1}$ ) would lead to large underestimation of the  $v = 1$  level energy. Moreover the vibrational constants assumed are close to the mean of those for three lowest  $\Omega = 4$  states ( $846$ ,  $935$ , and  $843 \text{ cm}^{-1}$ ), and with simplification used in our program that  $Q_{\text{vib,rot}}^{(i)} = (p_i/p_X)Q_{\text{vib,rot}}^{(X)}$  our choice should give better results.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (252.137 \pm 1.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 38.48092 + 33.0187 \times 10^{-3}(\text{T/K}) \\ &\quad - 40.4519 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 14.7496 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 5.15534 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1300 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 50.04939 - 18.1106 \times 10^{-3}(\text{T/K}) \\ &\quad + 12.3772 \times 10^{-6}(\text{T/K})^2 \\ &\quad - 1.71438 \times 10^{-9}(\text{T/K})^3 \\ &\quad + 2.50947 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1300–4000 K range.

## 6.8.2. Enthalpy of formation

The results of determination of the enthalpy of formation of UO(g) are listed in Table 76. From many works published, the most reliable mass spectrometric data including reference species with well established thermodynamic properties were chosen for the calculation of the enthalpy of formation. The selected enthalpy of formation value

$$\Delta_f H^\circ(\text{UO}, \text{g}, 298.15 \text{ K}) = (21.4 \pm 10) \text{ kJ mol}^{-1}$$

was obtained as a weighted average of results in Table 76 with reduced weights for the most divergent values from works of Ackermann *et al.*<sup>548</sup> and Steiger and Cater.<sup>549</sup> To the selected enthalpy of formation corresponds dissociation energy of UO (g) molecule  $D_0(\text{UO}) = (758.9 \pm 10) \text{ kJ mol}^{-1}$ .

## 6.9. NpO<sub>2</sub>(g)

### 6.9.1. Heat capacity and entropy

The thermodynamic functions of NpO<sub>2</sub>(g) in the standard state in the temperature range 298.15–4000 K have been calculated using the data given in Table 77.

Experimental data on molecular structure and spectra of NpO<sub>2</sub> are unknown. Neptunium dioxide was studied by relativistic DFT calculations by Liao *et al.*<sup>550</sup> A linear structure of  $D_{\infty h}$  symmetry was found for the  $X^4\Sigma_g$  ground state and the first excited state  $^4H_g$  lies only  $0.01 \text{ eV}$  ( $\approx 80 \text{ cm}^{-1}$ ) above  $X^4\Sigma_g$  state. According to Liao *et al.*<sup>550</sup> this small difference leaves the identity of the ground state in some doubt. The recent study of Infante *et al.*<sup>462</sup> at the SO-CASPT2 level clarified the electronic ground state to be  $^4H_{3,5g}$ . This state has a linear structure too and an equilibrium bond distance of  $r_e(\text{Np-O}) = 176.1 \text{ pm}$ , which was adopted for computing the principal moment of inertia of NpO<sub>2</sub>.

TABLE 76. The enthalpy of formation of UO(g), in  $\text{kJ mol}^{-1}$

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_r H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Drowart <i>et al.</i> <sup>514</sup>	M	2130–2530	$\text{UO}(\text{g}) + \text{Si}(\text{g}) = \text{U}(\text{g}) + \text{SiO}(\text{g})$	−36.6	20.7
	M	1700–2150	$\frac{1}{2}\text{U}(\text{l}) + \frac{1}{2}\text{UO}_2(\text{cr}) = \text{UO}(\text{g})$	569.9	27.4
Coppens <i>et al.</i> <sup>208</sup>	M	2238–2315	$\text{UO}(\text{g}) + \text{B}(\text{g}) = \text{U}(\text{g}) + \text{BO}(\text{g})$	−40.2	18.1
Ackermann <i>et al.</i> <sup>548</sup>	M	1540–2150	$\frac{1}{2}\text{U}(\text{l}) + \frac{1}{2}\text{UO}_2(\text{g}) = \text{UO}(\text{g})$	582.0	39.5
Steiger and Cater <sup>549</sup>	M	2148–2369	$\text{U}(\text{g}) + \text{YO}(\text{g}) = \text{UO}(\text{g}) + \text{Y}(\text{g})$	−51.0	12.4
Younés <i>et al.</i> <sup>221</sup>	M	1930–2280	$\text{UO}(\text{g}) + \text{La}(\text{g}) = \text{U}(\text{g}) + \text{LaO}(\text{g})$	−32.8	14.8
Selected value:			$D_0(\text{UO}) = 758.9 \pm 10$		$21.4 \pm 10$

<sup>a</sup>M = mass spectrometry.

TABLE 77. The molecular parameters for  $\text{NpO}_2(\text{g})$ 

Parameter	Value
Ground electronic state	$^4H_{3.5g}$
Symmetry group	$D_{\infty h}$
Symmetry number, $\sigma$	2
$I$ ( $\text{g cm}^2$ ) <sup>a</sup>	$16.5 \times 10^{-39}$
Vibrational frequencies ( $\text{cm}^{-1}$ ) <sup>b</sup>	939, 185(2), 880
Electronic states ( $\text{cm}^{-1}$ ) <sup>c</sup>	0(2), 470(2), 783(2), 1126(2), 1284(2), 5875(2), 5902(2), 5992(2), 6084(2), 6461(2), 6676(2), 8760(2), 9050(2), 10386(2), 10386(2), 10481(2), 10846(2), 10989(2), 11264(2), 11428(2), 12000(16), 14000(20), 16000(20), 18000(12), 20000(34), 22000(22), 24000(30), 26000(24)

<sup>a</sup>Moment of inertia.<sup>b</sup>Numbers in parentheses represent the degeneracies.<sup>c</sup>Numbers in parentheses represent the statistical weights.

Liao *et al.*<sup>550</sup> reported two vibrational frequencies for the  $X^4\Sigma_g$  state, but in Table 77 we adopted the more reliable frequencies evaluated by Kovács and Konings<sup>460</sup> for the ground state. Liao *et al.*<sup>550</sup> published also a few spin-free relativistic excitation energies to low-lying electronic states of  $\text{NpO}_2$ . In the view of the failed electronic ground state at their SO-VWN-B-P level these data have a limited reliability. Instead, we used the recent SO-CASPT2 vertical excitation energies of Kovács<sup>456</sup> for the evaluation of the thermodynamic functions. We have supplemented the set of  $\text{NpO}_2$  excited electron energy levels using the experimental data on the  $\text{Np}^{4+}$  energy levels in crystals. The information on the splitting of the  $\text{Np}^{4+}$  multiplets in the crystal fields of different symmetry is available from optical spectroscopy (e.g. Lahalle *et al.*<sup>551</sup>), and inelastic neutron scattering experiments (e.g. Kern *et al.*<sup>552</sup>; Fournier *et al.*<sup>553</sup>). We preferred to use the energy levels derived from the low-temperature absorption spectra of low-symmetry monoclinic  $\text{NpF}_4(\text{cr})$ , interpreted and supplemented with the results of crystal-field calculations by Carnall *et al.*<sup>554</sup> In the crystal field of monoclinic neptunium tetrafluoride with the  $C_2$  site symmetry the  $\text{Np}^{4+}$  multiplets split into doublets, constituting a set of levels appropriate for modeling  $\text{NpO}_2(\text{g})$  electron energy levels (see Table 77; the energy levels above  $11428 \text{ cm}^{-1}$  represent the results of combining of several close-lying levels).

The derived standard entropy at room temperature is:

$$S^\circ(298.15 \text{ K}) = (269.892 \pm 6.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 56.33269 + 40.03943 \times 10^{-3}(\text{T/K}) \\ &\quad - 54.9771 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 23.02883 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 0.732805 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1000 K range, and

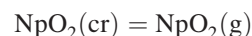
$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 68.29804 - 9.034032 \times 10^{-3}(\text{T/K}) \\ &\quad + 5.459315 \times 10^{-6}(\text{T/K})^2 \\ &\quad - 0.6628476 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 3.700337 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 1000–4000 K range.

## 6.9.2. Enthalpy of formation

The evaporation behavior of neptunium dioxide has been studied by Ackermann *et al.*<sup>438</sup> and Gotcu-Freis *et al.*<sup>555</sup> by mass-spectrometric measurements, showing that the gaseous oxides  $\text{NpO}_2$  and  $\text{NpO}$  are the only neptunium-bearing species in the vapor,  $\text{NpO}_2$  being the major constituent. In both studies it was argued that  $\text{NpO}_2$  becomes substoichiometric at the high temperatures in vacuum. The extent of substoichiometry, unknown in the former and about  $\text{O/Np} = 1.94$  at 2260 K in the latter grossly affects the partial pressure of  $\text{NpO}(\text{g})$  but has an insignificant influence on the partial pressure of  $\text{NpO}_2(\text{g})$ . This was also confirmed by Gotcu-Freis *et al.*<sup>555</sup> who studied the vaporisation of  $\text{NpO}_2$  in a slight flow of oxygen gas, showing no significant difference with the measurement in vacuum. Bartscher and Sari<sup>556</sup> have shown that the  $\text{O/Np}$  ratio may reach 1.97 at 1900 K. However, no thermodynamic data for the substoichiometric compositions are available in the temperature range of the vaporization experiments and, hence, the condensed phase can only be treated as stoichiometric.

Ackermann *et al.*<sup>438</sup> also measured the weight loss of neptunium dioxide from 1852 to 2474 K by Knudsen effusion technique. They obtained the equation for the temperature dependence of the total vapor pressure  $\log(P/\text{atm}) = -(31\,100 \pm 300)/T + (8.39 \pm 0.13)$ . Considering the equilibrium to be



we obtain from this equation the values  $\Delta_{\text{sub}}H^\circ(\text{NpO}_2, \text{cr}, 298.15) = (619.5 \pm 10) \text{ kJ mol}^{-1}$  by third-law and  $(623.2 \pm 12) \text{ kJ mol}^{-1}$  by second-law analysis. The mean second-law enthalpy for  $\text{NpO}_2$  in six mass-spectrometric experiments by Ackermann *et al.* (2100–2450 K), derived from the  $\text{NpO}_2^+$  ion currents is  $\Delta_{\text{sub}}H^\circ(\text{NpO}_2, \text{cr}, 298.15) = (622 \pm 5) \text{ kJ mol}^{-1}$ . The second-law values derived from the measurements in oxygen and vacuum by Gotcu-Freis *et al.*<sup>555</sup> are  $(650 \pm 4)$  and  $(627 \pm 7) \text{ kJ mol}^{-1}$ , in fair agreement.

We select the mean value of the work of Ackermann *et al.*<sup>438</sup> which is considered more precise,  $\Delta_{\text{sub}}H^\circ(\text{NpO}_2, \text{cr}, 298.15) = (621 \pm 20) \text{ kJ mol}^{-1}$ , which yields, when combined with the enthalpy of formation of  $\text{NpO}_2(\text{cr})$ :

$$\Delta_f H^\circ(\text{NpO}_2, \text{g}, 298.15) = -(457 \pm 20) \text{ kJ mol}^{-1}.$$

TABLE 78. Molecular constants of  $^{237}\text{Np}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$						$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$		
0	X0.5	0	840	2.92	0.334	1.6	2.1	183.5	2
1		200							2
2		700							4
3		1200							6
4		2600							18
5		4000							24
6		5300							16
7		6600							32
8		8100							20
9		10 000							40
10		15 000							100
11		20 000							200
12		25 000							600
13		30 000							900
14		35 000							1000
15		40 000							1200

## 6.10. NpO(g)

### 6.10.1. Heat capacity and entropy

The thermal functions of NpO(g) in the standard state in the temperature range 298.15–4000 K have been calculated using the data given in Table 78. As there were no spectral data on the NpO molecule at the time of writing of this review, the molecular constants were estimated using the trends revealed in the spectral data of the lanthanide monoxides. The electronic structure of the NpO molecule is assumed to be analogous to that of PmO. The energies of the ground state  $4f^4 6s$  super-configuration rounded to 2 significant digits were taken from the Ligand field calculation by Dulick *et al.*,<sup>247</sup> the energies of the other states were estimated as for PmO with small correction due to a little difference between relative positions the atomic states of the  $5f^4$  core. These values fairly well agree with the results of recent quantum chemical studies of Infante *et al.*<sup>462</sup> and Kovács and Konings.<sup>460</sup>

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (253.060 \pm 4.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 40.73102 + 5.06903 \times 10^{-3} (\text{T/K}) \\ &+ 5.58835 \times 10^{-6} (\text{T/K})^2 \\ &- 3.26062 \times 10^{-9} (\text{T/K})^3 \\ &- 3.95105 \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

TABLE 79. The enthalpy of formation of NpO(g), in  $\text{kJ mol}^{-1}$ 

Authors	Method <sup>a</sup>	T/K	Reaction	$\Delta_f H^\circ(298.15 \text{ K})$	$\Delta_f H^\circ(298.15 \text{ K})$
Ackermann <i>et al.</i> <sup>557</sup>	M	1665–1972	$\text{NpO}(\text{g}) + \text{La}(\text{g}) = \text{Np}(\text{g}) + \text{LaO}(\text{g})$	−63.6	−15.2
	M	1173–1972	$\text{Np}(\text{g}) + \text{YO}(\text{g}) = \text{NpO}(\text{g}) + \text{Y}(\text{g})$	−22.0	−19.6
Selected value:			$D_0(\text{NpO}) = 734.8 \pm 10$		−16.6 ± 10

<sup>a</sup>M = mass spectrometry.

for the 298.15–1400 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) &= 53.06105 + 1.44612 \times 10^{-3} (\text{T/K}) \\ &- 1.95684 \times 10^{-6} (\text{T/K})^2 \\ &+ 0.362588 \times 10^{-9} (\text{T/K})^3 \\ &- 5.12159 \times 10^6 (\text{T/K})^{-2} \end{aligned}$$

for the 1400–4000 K range.

### 6.10.2. Enthalpy of formation

In their study of the vaporisation of neptunium dioxide, Ackermann *et al.*<sup>557</sup> have revealed nonstoichiometric and not single-mode NpO<sub>2</sub> evaporation, excluding the determination of thermodynamic properties of NpO(g) directly from the effusion measurements. The thermodynamic properties of the NpO(g) molecule were determined by Ackermann and Rauh<sup>558</sup> from mass-spectrometric measurements of equilibrium constants for two oxygen-exchange reactions,  $\text{NpO}(\text{g}) + \text{La}(\text{g}) = \text{Np}(\text{g}) + \text{LaO}(\text{g})$  and  $\text{Np}(\text{g}) + \text{YO}(\text{g}) = \text{NpO}(\text{g}) + \text{Y}(\text{g})$ . Results of the calculation of the enthalpy of formation from both reactions are given in Table 79. For both reactions, the results are in good agreement.

The selected value

$$\Delta_f H^\circ(\text{NpO}, \text{g}, 298.15 \text{ K}) = -(16.6 \pm 10.0) \text{ kJ mol}^{-1}$$

is taken as a weighted mean of two values obtained, with the weight of NpO(g) + La(g) equilibrium twice as that of Np(g) + YO(g), due to the higher statistical errors of the latter. To the selected enthalpy of formation corresponds the value of dissociation energy,  $D_0(\text{NpO}) = (734.8 \pm 10) \text{ kJ mol}^{-1}$ . This is in good agreement with the value  $(744 \pm 21) \text{ kJ mol}^{-1}$  estimated by Marçalo and Gibson<sup>500</sup> on the basis of gas phase oxidation reactions involving the ionised species and known ionisation energies.

## 6.11. PuO<sub>3</sub>(g)

### 6.11.1. Heat capacity and entropy

The thermodynamic functions of PuO<sub>3</sub>(g) in the standard state in the temperature range 298.15–4000 K have been calculated using the data given in Table 80. There are no experimental data on molecular structure and spectra of PuO<sub>3</sub>. Gao *et al.*<sup>559</sup> and Zaitsevskii *et al.*<sup>560</sup> have calculated the molecular structure for PuO<sub>3</sub> by Hartree-Fock and DFT methods. The results of the calculations show that the PuO<sub>3</sub> molecule has a planar structure of C<sub>2v</sub> symmetry similar to



TABLE 80. The molecular parameters for PuO<sub>3</sub>(g)

Parameter	Value
Ground electronic state	<sup>7</sup> B <sub>1</sub>
Symmetry group	C <sub>2v</sub>
Symmetry number, σ	2
I <sub>A</sub> I <sub>B</sub> I <sub>C</sub> (g <sup>3</sup> cm <sup>6</sup> ) <sup>a</sup>	4810 × 10 <sup>-117</sup>
Vibrational frequencies (cm <sup>-1</sup> )	840, 700, 170, 800, 190, 145
Electronic states (cm <sup>-1</sup> ) <sup>b</sup>	0(2), 2530(1), 4870(2), 6700(2), 10334(1), 10983(2), 11225(1), 11651(2), 12326(1), 16713(1), 17737(2), 18565(2), 20029(1), 22703(2), 22889(2), 23022(2), 29710(2), 32198(2), 32759(1), 34080(2), 34702(2), 34982(2)

<sup>a</sup>Product of moments of inertia.<sup>b</sup>Numbers in parentheses represent the statistical weights.

that of UO<sub>3</sub>. The molecular constants of PuO<sub>3</sub> obtained by Gao *et al.*<sup>559</sup> seem unreliable being significantly different from the corresponding molecular constants of UO<sub>3</sub>. The product of the principal moments of inertia of PuO<sub>3</sub> (see Table 80) is calculated using  $r_e(\text{Pu-O}) = (185 \pm 5)$  pm (two equal bonds),  $r_e(\text{Pu-O}') = (190 \pm 5)$  pm, and  $\angle \text{O-Pu-O} = (160 \pm 10)^\circ$ . The bond distances are estimated by comparison with molecular constants of UO, UO<sub>2</sub>, UO<sub>3</sub>, PuO, and PuO<sub>2</sub>. The value of the O-Pu-O bond angle is accepted to be equal to that in UO<sub>3</sub> molecule (see UO<sub>3</sub>).

The vibrational frequencies calculated by Gao *et al.*,<sup>559</sup> which have too low values, were not used for estimation of vibrational contribution to the PuO<sub>3</sub> thermodynamic functions. The values of vibrational frequencies (see Table 80) are estimated by comparison with corresponding frequencies adopted for UO, UO<sub>2</sub>, UO<sub>3</sub>, PuO, and PuO<sub>2</sub> molecules.

According to the calculations of Gao *et al.*,<sup>559</sup> the PuO<sub>3</sub> molecule has a <sup>7</sup>B<sub>1</sub> ground state. Experimental and theoretical investigations of PuO<sub>3</sub> excited electron energy levels are unknown. The energy levels of PuO<sub>3</sub> are assumed to be approximately the same as for the isoelectronic ion PuO<sub>2</sub><sup>2+</sup> (configuration [Rn]5f<sup>2</sup>). These energy levels have been obtained by Infante *et al.*<sup>561</sup> using the novel relativistic intermediate Hamiltonian Fock-space coupled-cluster method.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (319.450 \pm 4) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 52.89593 + 75.46988 \times 10^{-3} (\text{T/K}) \\ & - 71.42497 \times 10^{-6} (\text{T/K})^2 \\ & + 25.87474 \times 10^{-9} (\text{T/K})^3 \\ & - 3.753825 \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

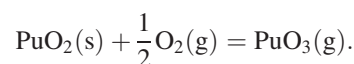
for the 298.15–900 K range, and

$$\begin{aligned} C_p^\circ / (\text{J K}^{-1} \text{ mol}^{-1}) = & 76.50587 + 8.903191 \times 10^{-3} (\text{T/K}) \\ & - 1.504422 \times 10^{-6} (\text{T/K})^2 \\ & + 0.07709618 \times 10^{-9} (\text{T/K})^3 \\ & - 16.13944 \times 10^5 (\text{T/K})^{-2} \end{aligned}$$

for the 900 K–4000 K range.

## 6.11.2. Enthalpy of formation

Krikorian *et al.*<sup>562</sup> have carried out transpiration experiments to study the volatility of PuO<sub>2</sub>(cr) in the presence of oxygen and steam. In experiments in presence of oxygen, it was expected, in analogy to uranium vaporisation behavior, that volatilization can occur to PuO<sub>3</sub>(g) species, with insignificant amounts of the known plutonium oxide species, PuO(g) and PuO<sub>2</sub>(g), according to known thermodynamic data. The existence of the PuO<sub>3</sub> vapor species was later confirmed by Ronchi *et al.*<sup>563</sup> who detected the molecule in mass-spectrometric study of vapors over plutonium dioxide. Krikorian *et al.*<sup>562</sup> used their data for 1 atm oxygen gas present (3 points, 1482 and 1483 K) to obtain the pressures of PuO<sub>3</sub>(g) and to calculate equilibrium constants  $K_p = p(\text{PuO}_3)/a(\text{PuO}_2)p(\text{O}_2)^{1/2}$ , for the reaction



Taking the activity of PuO<sub>2</sub>(cr) as unity, Krikorian *et al.*<sup>562</sup> obtained an average third-law value  $\Delta_f H^\circ(298.15 \text{ K}) = (493.0 \pm 4.9) \text{ kJ mol}^{-1}$  (uncertainty is standard deviation). We have recalculated the equilibrium constants of the above reaction with the thermodynamic functions for PuO<sub>2</sub>(cr) and PuO<sub>3</sub>(g) adopted in this assessment:  $\Delta_f H^\circ(298.15 \text{ K}) = (488.2 \pm 15) \text{ kJ mol}^{-1}$ , with an estimate of total uncertainty. Combining this value with the enthalpy of formation of PuO<sub>2</sub>(cr) gives the adopted value

$$\Delta_f H^\circ(\text{PuO}_3, \text{g}, 298.15) = -(567.6 \pm 15) \text{ kJ mol}^{-1}.$$

## 6.12. PuO<sub>2</sub>(g)

### 6.12.1. Heat capacity and entropy

The thermodynamic functions of PuO<sub>2</sub>(g) in the standard state have been calculated using the data given in Table 81.

The only experimental study on the molecular properties of PuO<sub>2</sub> is the infrared spectroscopic measurement of PuO<sub>2</sub> isotopomers in Ar and Kr matrices by Green and Reedy.<sup>564</sup> The O-Pu-O bond angle of 180° was calculated from the measured isotope shift. This is consistent with the absence of peaks attributable to  $\nu_1$  for the case of Pu<sup>16</sup>O<sub>2</sub> and Pu<sup>18</sup>O<sub>2</sub>. There are no experimental data on Pu–O bond distance in

TABLE 81. The molecular parameters for PuO<sub>2</sub>(g)

Parameter	Value
Ground electronic state	<sup>5</sup> Σ <sub>g</sub>
Symmetry group	D <sub>∞h</sub>
Symmetry number, σ	2
I (g cm <sup>2</sup> ) <sup>a</sup>	18.0 × 10 <sup>-39</sup>
Vibrational frequencies (cm <sup>-1</sup> ) <sup>b</sup>	792, 109(2), 828
Electronic states (cm <sup>-1</sup> ) <sup>c</sup>	0(1), 420(1), 513(1), 516(1), 525(1), 1077(1), 1079(1), 1358(1), 1383(1), 4955(1), 5094(1), 5149(1), 5149(1), 5208(1), 5277(1), 5390(1), 5571(1), 5571(1), 5698(1), 5698(1), 8700(3), 9400(12), 10400(3), 11700(5), 12500(11)

<sup>a</sup>Moment of inertia.<sup>b</sup>Numbers in parentheses represent the degeneracies.<sup>c</sup>Numbers in parentheses represent the statistical weights.

PuO<sub>2</sub>. The principal moment of inertia of PuO<sub>2</sub> (see Table 81) is calculated using  $r_e(\text{Pu-O}) = (184 \pm 5)$  pm. This value is an average of two bond distances (181.2 and 187.0 pm) obtained in *ab initio* calculations of Liao *et al.*<sup>550</sup> and Archibong and Ray.<sup>565</sup> The recent computational study by Zaitsevskii *et al.*<sup>560</sup> is in fair agreement.

The adopted values of PuO<sub>2</sub> vibrational frequencies (see Table 81) are taken from the *ab initio* calculation of Archibong and Ray<sup>565</sup> at the CCSD(T) level. The adopted values  $\nu_1$  and  $\nu_3$  correlate with the results of density-functional method including spin-orbit effects.

There are no experimental data on electronic spectra of PuO<sub>2</sub>. Only results obtained in theoretical calculations are available. According to the calculations made by Liao *et al.*<sup>550</sup> and Archibong and Ray<sup>565</sup> the ground state of PuO<sub>2</sub> is X<sup>5</sup>Σ<sub>g</sub>. Both calculations do not take into consideration the effects of spin-orbit interactions and present limited numbers of energy levels. Therefore, these data were not used for the evaluation of electronic contribution to the PuO<sub>2</sub> thermodynamic functions. We preferred to use the energy levels derived from the low-temperature absorption spectra of low-symmetry monoclinic PuF<sub>4</sub>(cr), interpreted and supplemented with the results of crystal-field calculations by Carnall *et al.*<sup>554</sup> In the crystal field of monoclinic plutonium tetrafluoride with the C<sub>2</sub> site symmetry the Pu<sup>4+</sup> multiplets split into singlets. These singlet levels were used for modeling PuO<sub>2</sub>(g) levels (see Table 81; the energy levels above 5698 cm<sup>-1</sup> represent the results of combining of several close-lying levels).

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (278.741 \pm 5) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & 70.29021 - 13.46001 \times 10^{-3}(\text{T/K}) \\ & + 9.065878 \times 10^{-6}(\text{T/K})^2 \\ & - 1.403447 \times 10^{-9}(\text{T/K})^3 \\ & - 4.810746 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1500 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) = & 49.85911 + 11.46596 \times 10^{-3}(\text{T/K}) \\ & - 1.472281 \times 10^{-6}(\text{T/K})^2 \\ & - 8.555747 \times 10^{-12}(\text{T/K})^3 \\ & + 41.20745 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 1500–4000 K range.

### 6.12.2. Enthalpy of formation

Total pressure over substoichiometric plutonium dioxide was measured in several investigations. Phipps *et al.*<sup>566</sup> employed the Knudsen effusion method to measure total vapor pressures between 1593 and 2063 K; evaporation was performed from a tantalum effusion cell. The authors came to conclusion that “reduction of plutonium dioxide occurs when it is heated in vacuum in a tantalum oven.” Mulford and Lamar<sup>567</sup> reported the vapor pressure data of plutonium dioxide measured at temperatures between 2000 and 2400 K using tungsten Knudsen effusion cells. The effusion rates were considerably less than those observed by Phipps. Pardue and Keller<sup>568</sup> measured the vapor pressure of plutonium dioxide in air, argon, and oxygen at temperatures between 1723 and 2048 K, using the transpiration technique. Total vapor pressures of PuO<sub>2-x</sub> compositions were measured by Messier<sup>569</sup> gravimetrically with tungsten Knudsen cells from 2070 to 2380 K, and the vapor pressure equation for the composition approximating the congruent one, PuO<sub>1.82</sub>, was obtained. According to mass-spectrometric measurements of Battles *et al.*,<sup>570</sup> PuO(g) is the major species in the vapor phase of the Pu<sub>2</sub>O<sub>3</sub> + PuO<sub>1.61</sub> system. In nonreducing atmosphere PuO pressure significantly diminishes. Nevertheless, its part in the total vapor pressure must be taken into account in deriving PuO<sub>2</sub> pressure over PuO<sub>2-x</sub> phase.

Extensive experimental study and detailed thermodynamic analysis of the vaporization of the substoichiometric plutonium dioxide has been carried out by Ackermann *et al.*<sup>438</sup> The rates of evaporation of plutonium-bearing species from the plutonium dioxide phase in the range 1646–2104 K were measured by the effusion method. The effusion cells were made from tungsten, rhenium, and tantalum. The results obtained with tungsten and rhenium cells were found to be in good agreement. The total pressure calculated under assumption of PuO<sub>2</sub>(g) as the only vapor species was

approximated by the equation

$$\log(p/\text{atm}) = -(29620 \pm 280)/(T/\text{K}) + (7.50 \pm 0.15).$$

The above equation corresponds to evaporation of the  $\text{PuO}_{1.92}$  composition, which did not significantly change during the time of measurements. Results obtained with the tantalum effusion cell, i.e. under reducing conditions, correspond to the univariant evaporation of the  $\text{Pu}_2\text{O}_3 + \text{PuO}_{1.61}$  system. Using data on evaporation in reducing conditions, free energy of formation of  $\text{PuO}(\text{g})$  was derived and the  $\text{PuO}(\text{g})$  pressure over  $\text{PuO}_{1.92}$  phase was calculated. Values of  $\text{PuO}_2(\text{g})$  pressure evaluated in the 1600–2150 K range were fitted to the equation:

$$\log(p/\text{atm}) = -29640/(T/\text{K}) + 7.67.$$

In combination with auxiliary data from literature, this equation was used by Ackermann *et al.*<sup>438</sup> for deriving the linear equation for the standard free energy of formation of  $\text{PuO}_2(\text{g})$ , in cal/mol:

$$\Delta_f G^\circ(\text{PuO}_2, \text{g}) = -113100 + 4.35(T/\text{K}).$$

This equation is combined with Gibbs energy of formation of  $\text{PuO}_2(\text{cr})$  for calculation of the  $\text{PuO}_2(\text{g})$  pressures over stoichiometric  $\text{PuO}_2(\text{cr})$  in the 1650–2150 K temperature range. With the pressure values so obtained we find the third-law value for the enthalpy of sublimation:  $\Delta_{\text{sub}} H^\circ(\text{PuO}_2, \text{s}, 298.15) = (643.9 \pm 15) \text{ kJ mol}^{-1}$ , yielding  $\Delta_f H^\circ = -(411.9 \pm 15) \text{ kJ mol}^{-1}$ .

Gotcu-Freis *et al.*<sup>571</sup> studied the evaporation of  $\text{PuO}_2$  by mass spectrometry in vacuum and in a flow of low pressure oxygen. Their results in vacuum are in excellent agreement with most of the earlier studies, including Ackermann *et al.*<sup>438</sup> They observed, however, a significant difference between the measurement in vacuum and oxygen, suggesting that the change in O/Pu ratio has an influence on the vaporisation equilibria. From the analysis of the measurements in oxygen considering the reaction  $\text{PuO}_2(\text{cr}) = \text{PuO}_2(\text{g})$  they obtained  $\Delta_{\text{sub}} H^\circ(298.15 \text{ K}) = (627 \pm 7) \text{ kJ mol}^{-1}$  by second law analysis and  $\Delta_{\text{sub}} H^\circ(298.15 \text{ K}) = (616 \pm 6) \text{ kJ mol}^{-1}$  by third law analysis, corresponding to  $\Delta_f H^\circ = -(428 \pm 7) \text{ kJ mol}^{-1}$  and  $\Delta_f H^\circ = -(440 \pm 6) \text{ kJ mol}^{-1}$ , respectively.

The selected value of  $\text{PuO}_2(\text{g})$  enthalpy of formation is

$$\Delta_f H^\circ(\text{PuO}_2, \text{g}, 298.15) = -(428 \pm 20) \text{ kJ mol}^{-1}.$$

## 6.13. PuO(g)

### 6.13.1. Heat capacity and entropy

The thermal functions of  $\text{PuO}(\text{g})$  in the standard state have been calculated using the data given in Table 82.

The electronic spectrum of  $\text{PuO}$  is unknown. The  $\text{Pu}^{16}\text{O}$  and  $\text{Pu}^{18}\text{O}$  molecules were identified in Ar and Kr matrices by Green and Reedy.<sup>564</sup> The  $\omega_e$  and  $\omega_e x_e$  values given in Table 82 were calculated using these data taking into account the matrix shift ( $\sim 15 \text{ cm}^{-1}$ ) from the condition of converging the vibrational levels to the adopted dissociation limit. The internuclear distance  $r_e = (184 \pm 2) \text{ pm}$  is estimated from the comparison of

TABLE 82. Molecular constants of  $^{239}\text{Pu}^{16}\text{O}(\text{g})$

No.	State	$\text{cm}^{-1}$						$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$		
0 <sup>a</sup>	$X0^{-c}$	0	830	3.1	0.33	1.6	2.13	184.0	1
1 <sup>b</sup>		150							2
2 <sup>b</sup>		600							3
3 <sup>b</sup>		900							2
4 <sup>b</sup>		1300							2
5 <sup>b</sup>		1600							6
6 <sup>b</sup>		2000							2
7 <sup>b</sup>		2400							8
8 <sup>b</sup>		3100							22
9 <sup>b</sup>		4200							20
10 <sup>b</sup>		5000							4
11 <sup>b</sup>		10 000							20
12 <sup>b</sup>		15 700							120
13 <sup>b</sup>		20 000							320
14 <sup>b</sup>		25 000							340
15 <sup>b</sup>		30 000							470
16 <sup>b</sup>		35 000							1640
17 <sup>b</sup>		40 000							2400

<sup>a</sup>Assumed to be analogous to  $\text{SmO}(\text{g})$ .

<sup>b</sup>Estimated state.

the  $r_e$  values in the ground states for the  $\text{CeO}$  and  $\text{NdO}$  with the same values for the  $\text{ThO}$  and  $\text{UO}$  molecules.

*Ab initio* studies by Gao *et al.*<sup>572</sup> and by Li and Xu<sup>573</sup> gave the ground state  $X^5\Sigma^-$ ,  $r_e = 183 \text{ pm}$ ,  $\omega_e = 781.15 \text{ cm}^{-1}$ ; the latter value disagreed with the matrix data. The electronic structure of  $\text{PuO}$  should be analogous to that of  $\text{SmO}$ , which has the  $X0^-$  ground state and near lying  $\Omega = 1$  and  $\Omega = 2$  states. The ground state configuration of  $\text{SmO}$  is  $4f^5 6s$ , which does not generate a  $^5\Sigma^-$  state. Details of the *Ab initio* studies by Gao *et al.*<sup>572</sup> and by Li and Xu<sup>573</sup> are not available. The information was taken from abstracts in Chemical Abstract, where the symmetry of the  $X^5\Sigma^-$  may be indicated erroneously. The  $\omega_e = (830 \pm 10) \text{ cm}^{-1}$  value, derived from the matrix spectra, is characteristic for the  $5f^{N-1}7s$  configuration: see  $\omega_e(\text{UO}) \sim 846$  (deperturbed value). In the present work for  $\text{PuO}$  we assume the electronic structure analogous to that of  $\text{SmO}$ . The more dense structure (total statistical weight of excited states is equal to 5382) is due to the more close arrangement of the atomic states of the  $5f^6$  core  $\text{PuIV}$ .

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (252.254 \pm 3.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 25.39529 + 54.2560 \times 10^{-3}(T/\text{K}) \\ &\quad - 36.2549 \times 10^{-6}(T/\text{K})^2 \\ &\quad + 6.47005 \times 10^{-9}(T/\text{K})^3 \\ &\quad + 3.72127 \times 10^4(T/\text{K})^{-2} \end{aligned}$$

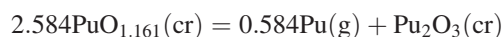
for the 298.15–1200 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 78.23663 - 27.3687 \times 10^{-3}(T/\text{K}) \\ &\quad + 6.77906 \times 10^{-6}(T/\text{K})^2 \\ &\quad - 0.440726 \times 10^{-9}(T/\text{K})^3 \\ &\quad - 7.04572 \times 10^6(T/\text{K})^{-2} \end{aligned}$$

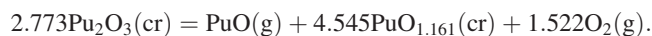
for the 1200–4000 K range.

### 6.13.2. Enthalpy of formation

The total pressure of plutonium-bearing species in the process of evaporation of plutonium oxides in tantalum crucibles is well established and is interpreted to correspond to the equilibrium state in the  $\text{Pu}_2\text{O}_3 + \text{PuO}_{1.61}$  two-phase system (see Cordfunke and Konings<sup>8</sup>). According to mass-spectrometric measurements by Battles *et al.*,<sup>570</sup>  $\text{PuO}(\text{g})$  is the major species in the vapor phase of the  $\text{Pu}_2\text{O}_3 + \text{PuO}_{1.61}$  system. For calculation of the enthalpy of formation of  $\text{PuO}(\text{g})$  the data obtained by Ackermann *et al.*<sup>438</sup> were used. The data of this paper are in good agreement with the work of Phipps *et al.*,<sup>566</sup> as well as with data of Ohse,<sup>574</sup> Messier<sup>569</sup> and Gotcu-Freis *et al.*<sup>571</sup> The pressure of  $\text{PuO}_2(\text{g})$  in the system is small and can be neglected. A small amount of  $\text{Pu}(\text{g})$  was calculated from the equilibrium constant for the reaction



After correcting the total pressure for  $\text{Pu}(\text{g})$ , the enthalpy of formation for  $\text{PuO}(\text{g})$  was calculated from the enthalpy of reaction



The third-law value obtained,  $\Delta_f H^\circ(298.15 \text{ K}) = 560.4 \text{ kJ mol}^{-1}$  leads to the enthalpy of formation  $\text{PuO}(\text{g})$ , which is selected:

$$\Delta_f^\circ(\text{PuO}, \text{g}, 298.15 \text{ K}) = -(51.7 \pm 15) \text{ kJ mol}^{-1}$$

This value corresponds to the dissociation energy  $D_0(\text{PuO}) = (648.7 \pm 30) \text{ kJ mol}^{-1}$ . This is in good agreement with the value  $(658 \pm 10) \text{ kJ mol}^{-1}$  estimated by Marçalo and Gibson<sup>500</sup> on the basis of gas phase oxidation reactions involving the ionised species and known ionisation energies.

## 6.14. $\text{AmO}_2(\text{g})$

### 6.14.1. Heat capacity and entropy

The thermodynamic functions of  $\text{AmO}_2(\text{g})$  in the standard state have been calculated using the data given in Table 83. Experimental data on the molecular structure and spectra of  $\text{AmO}_2$  are not available. Theoretical calculations of the structure

TABLE 83. The molecular parameters for  $\text{AmO}_2(\text{g})$

Parameter	Value
Ground electronic state	$X^6\Pi_{2,5u}$
Symmetry group	$D_{\infty h}$
Symmetry number, $\sigma$	2
$I$ ( $\text{g cm}^2$ ) <sup>a</sup>	$17.35 \times 10^{-39}$
Vibrational frequencies ( $\text{cm}^{-1}$ ) <sup>b</sup>	815, 105(2), 757
Electronic states ( $\text{cm}^{-1}$ ) <sup>c</sup>	0(2), 447(2), 5515(2), 6565(2), 12622(2), 15418(2), 15668(2), 16664(2), 18676(2), 21580(2), 21925(2), 22230(2), 22469(2), 23882(2), 25844(2), 28909(2)

<sup>a</sup>Moment of inertia.

<sup>b</sup>Numbers in parentheses represent degeneracy.

<sup>c</sup>Numbers in parentheses represent the statistical weights.

of the  $\text{AmO}_2$  molecule have been made by Kovács *et al.*,<sup>575</sup> which showed that the molecule has a linear  $D_{\infty h}$  structure with  $r(\text{Am-O}) = 180.7 \text{ pm}$ . The principal moment of inertia of  $\text{AmO}_2$  is calculated from this value. Subsequent calculations by Infante *et al.*<sup>462</sup> provided details on the ground-state electronic structure of the molecule. The harmonic and anharmonic vibrational frequencies were calculated by Kovács and Konings<sup>460</sup> using DFT methods, while the most important low-lying electronic states were reported in Ref. 575.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (279.464 \pm 6) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 52.7906 + 26.0768 \times 10^{-3}(\text{T/K}) \\ &\quad - 27.1648 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 10.2293 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 3.3759 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1000 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 54.4906 + 7.17462 \times 10^{-3}(\text{T/K}) \\ &\quad - 1.46280 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 0.13147 \times 10^{-9}(\text{T/K})^3 \\ &\quad + 1.2604 \times 10^6(\text{T/K})^{-2} \end{aligned}$$

for the 1000–4000 K range.

### 6.14.2. Enthalpy of formation

Gotcu-Freis *et al.*<sup>576</sup> measured the vapor pressure over  $\text{AmO}_{2-x}$  identifying  $\text{AmO}(\text{g})$ ,  $\text{AmO}_2(\text{g})$  and  $\text{Am}(\text{g})$  as vapor species. However, due to the strong change in the O/Am ratio of the samples during the measurements, it is not possible to derive the enthalpy of formation of  $\text{AmO}_2(\text{g})$  from these results. Kovács *et al.*<sup>575</sup> calculated the dissociation energy of the reaction  $\text{AmO}_2(\text{g}) = \text{AmO}(\text{g}) + \text{O}(\text{g})$  as 6.03 eV (581.8  $\text{kJ mol}^{-1}$ ) using multiconfigurational relativistic quantum chemical methods. From these data we obtain

$$\Delta_f H^\circ(\text{AmO}_2, \text{g}, 298.15) = -(514.0 \pm 30) \text{ kJ mol}^{-1}.$$

## 6.15. $\text{AmO}(\text{g})$

### 6.15.1. Heat capacity and entropy

Thermal functions of  $\text{AmO}(\text{g})$  in the standard state have been calculated using the data given in Table 84.

As there is no spectral data on the  $\text{AmO}$  molecule, the estimates of the molecular constants given in that table are based on the trends revealed in the spectral data of the lanthanide monoxides. After the completion of this assessment, a few quantum chemical calculations on the molecular geometry, ro-vibrational properties and low-lying electronic states were published.<sup>460,462,575</sup> Infante *et al.*<sup>462</sup> confirmed the electronic ground state of  $\text{AmO}$  to be  $X^8\Sigma_{1/2}$ . But significant



TABLE 84. Molecular constants of  $^{243}\text{Am}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$						$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$		
0	$X^8\Sigma$	0	700	2.95	0.311	1.7	2.461	190	8
1		5000							8
2		6000							16
3		7000							18
4		8000							28
5		12 000							50
6		15 000							30
7		20 000							30
8		25 000							430
9		30 000							2200
10		35 000							2800
11		40 000							3000

differences were found for the other properties ( $\omega_e = 872 \text{ cm}^{-1}$ ,  $r_e = 180.1 \text{ pm}$ ).

The electronic structure of AmO is assumed to be analogous to that of EuO. Small difference in the recommended statistical weights of the united terms is due mainly to a little smaller accepted energy of  $A^8\Sigma(f^6s)$  state.

Uncertainties of the calculated thermal functions are due mainly to the unknown energy of the first excited state  $A^8\Sigma(f^6s)$ . The minimum on the curve presented the variation of the actinide oxides dissociation energy with the charge on the actinide atoms implies that the ground state of the AmO molecule is  $X^8\Sigma(f^7)$ . However, as in case of EuO, the position of  $A^8\Sigma(f^6s)$  is quite unclear.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (259.105 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 34.31966 + 6.13074 \times 10^{-3}(\text{T/K}) \\ &\quad - 6.40603 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 4.00456 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 2.68922 \times 10^5(\text{T/K})^{-2} \end{aligned}$$

for the 298.15–1500 K range, and

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= -85.66885 + 111.574 \times 10^{-3}(\text{T/K}) \\ &\quad - 30.2584 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 2.82855 \times 10^{-9}(\text{T/K})^3 \\ &\quad + 4.35180 \times 10^7(\text{T/K})^{-2} \end{aligned}$$

for the 1500–4000 K range.

### 6.15.2. Enthalpy of formation

No experimental measurement of the enthalpy of formation AmO(g) is known. Estimates of this value can be made using estimates of the AmO dissociation energy. The dissociation energy of AmO(g) was estimated by Haire<sup>577</sup> as  $550 \text{ kJ mol}^{-1}$  using correlation between dissociation energy and promotion energy, needed for excitation of the free actinide atom in the  $ds^2$

configuration. The estimation was confirmed by ICR mass spectrometric study of reactivity of americium oxide ions by Santos *et al.*<sup>578</sup> From the results of experiments, Santos *et al.* estimated  $D_0(\text{Am}^+-\text{O}) = (580 \pm 50) \text{ kJ mol}^{-1}$  and found the ionization energy  $\text{IE}(\text{AmO}) = (5.9 \pm 0.2) \text{ eV}$ , or  $(569 \pm 20) \text{ kJ mol}^{-1}$ . Later, Marçalo and Gibson<sup>500</sup> revised the former value to  $(560 \pm 28) \text{ kJ mol}^{-1}$ . The thermochemical relation

$$\text{IE}(\text{AmO}) + D_0(\text{Am}^+ - \text{O}) - \text{IE}(\text{Am}) = D_0(\text{AmO})$$

resulted in the value  $D_0(\text{AmO}) = 553 \text{ kJ mol}^{-1}$ , in agreement with an previous estimate.<sup>577</sup> The rounded value  $D_0(\text{AmO}) = (550 \pm 50) \text{ kJ mol}^{-1}$  is selected in this work. It corresponds to

$$\Delta_f H^\circ(\text{AmO}, \text{g}, 298.15 \text{ K}) = -(15 \pm 50) \text{ kJ mol}^{-1}.$$

## 6.16. CmO(g)

### 6.16.1. Heat capacity and entropy

Thermal functions of CmO(g) in the standard state have been calculated using the data given in Table 85. Since there is no molecular data on the CmO molecule, the molecular constants given in this table were estimated using the trends revealed in the spectral data of the lanthanide monoxides. The electronic structure is assumed to be analogous to that of GdO. The more dense structure (total statistical weight of excited states is estimated to be equal to 3209, that of GdO to 1834) is due to the more dense arrangement of the atomic states of the  $5f^5$  core CmIV. After the completion of this assessment, a few quantum chemical calculations on the molecular geometry, rovibrational properties and low-lying electronic states were published.<sup>460,462,575</sup> Infante *et al.*<sup>462</sup> confirmed the electronic ground state of CmO to be  $X^9\Sigma_4$ . The other properties agree well with the estimated values.

The derived standard entropy at room temperature is

$$S^\circ(298.15 \text{ K}) = (259.071 \pm 5.0) \text{ J K}^{-1} \text{ mol}^{-1}$$

and the coefficients of the equations for the heat capacity are

$$\begin{aligned} C_p^\circ/(\text{J K}^{-1} \text{ mol}^{-1}) &= 26.15923 + 23.0665 \times 10^{-3}(\text{T/K}) \\ &\quad - 11.8949 \times 10^{-6}(\text{T/K})^2 \\ &\quad + 1.83564 \times 10^{-9}(\text{T/K})^3 \\ &\quad - 3.50044 \times 10^4(\text{T/K})^{-2} \end{aligned}$$

TABLE 85. Molecular constants of  $^{247}\text{Cm}^{16}\text{O}(\text{g})$ 

No.	State	$\text{cm}^{-1}$						$r_e$ pm	$p_i$
		$T_e$	$\omega_e$	$\omega_e x_e$	$B_e$	$\alpha_e 10^3$	$D_e 10^7$		
0	$X^9\Sigma_4$	0	840	2.88	0.333	1.5	2.1	183.5	9
1		2300							7
2		10 000							18
3		17 000							50
4		20 000							65
5		25 000							120
6		30 000							400
7		35 000							1040
8		40 000							1500

TABLE 86. Molecular constants and enthalpies of formation for AnO(g) (An = Bk–Lr)

An	$\omega_e$	$\omega_e x_e^a$	$B_e$	$\alpha_e 10^{3a}$	$r_e$	$\Delta_r H(0 \text{ K})^b$		$\Delta_r H(298.15 \text{ K})$
						Calculated	Estimated <sup>c</sup>	
			$\text{cm}^{-1}$		$\text{pm}$	$\text{kJ mol}^{-1}$		$\text{kJ mol}^{-1}$
BkO	833	3	0.333	1	183.5	537	598	543
CfO	833	3	0.338	1	182.2	479	498	485
EsO	825	3	0.338	1	182.2	403	460	409
FmO	735	3	0.327	1	185.0	369	443	373
MdO	673	3	0.311	1	189.8	341	418	349
NoO	650	3	0.303	1	192.3	290	268	295
LrO	756	3	0.319	1	187.1	583	665	588

<sup>a</sup>Estimated on the basis of the data of light actinide monoxides.

<sup>b</sup>Bond dissociation  $\text{AnO} \rightarrow \text{An} + \text{O}$ .

<sup>c</sup>Haire.

for the 298.15–1200 K range, and

$$C_p^\circ / (\text{JK}^{-1} \text{mol}^{-1}) = 46.16751 - 3.81910 \times 10^{-3} (\text{T/K}) \\ + 0.473867 \times 10^{-6} (\text{T/K})^2 + 0.160603 \\ \times 10^{-9} (\text{T/K})^3 - 3.86840 \\ \times 10^6 (\text{T/K})^{-2}$$

for the 1200–4000 K range.

### 6.16.2. Enthalpy of formation

Smith and Peterson<sup>579</sup> have performed Knudsen-effusion measurements of  $\text{Cm}_2\text{O}_3$  evaporation in the temperature range 1800–2600 K. Curium oxide has been shown to vaporize congruently. Thermodynamic analysis has led to conclusion that the predominant vaporization process is the formation of  $\text{CmO}(\text{g})$  and  $\text{O}(\text{g})$ . From the enthalpy of reaction



an approximate value  $D_0(\text{CmO}) = 728 \text{ kJ mol}^{-1}$  was obtained, using for the curium oxide the enthalpy of formation equal to that of  $\text{Pu}_2\text{O}_3$ . The mechanism of curium oxide vaporization proposed by Smith and Peterson obtained unambiguous confirmation in the mass spectrometric work on curium vaporization from a mixed curium-plutonium oxide by Hiernaut and Ronchi.<sup>580</sup> It was shown that  $\text{CmO}$  is the only effective vapor species.

With the thermodynamic functions of  $\text{Cm}_2\text{O}_3(\text{cr})$  from this assessment, the enthalpy of the above reaction was calculated from the equilibrium constant for the temperature

1843 K:  $\Delta_r H^\circ(298.15 \text{ K}) = 1782.4 \text{ kJ mol}^{-1}$ . This value yields

$$\Delta_r H^\circ(\text{CmO}, \text{g}, 298.15 \text{ K}) = -(75.4 \pm 20) \text{ kJ mol}^{-1}$$

which is selected. The corresponding dissociation energy is  $D_0(\text{CmO}) = 708.3 \text{ kJ mol}^{-1}$ . It should be mentioned that Gibson<sup>261</sup> estimated the value  $D_0(\text{CmO}) = 709 \text{ kJ mol}^{-1}$  using the concept of a Ln or An atom into a  $d^2s$  configuration thought to be needed for formation of the Ln-O or An-O bond.

### 6.17. Computed data for AnO(g) and AnO<sub>2</sub>(g) (An = Bk–Lr)

Presently, there is no experimental information available in the literature on the molecular parameters on the monoxides and dioxides of the late actinides. The bond distance and harmonic vibrational frequency of  $\text{LrO}$  has been computed by Cao and Dolg.<sup>581</sup> The DFT study by Kovács *et al.*<sup>498</sup> using relativistic effective core potentials on the actinides provided the first data on the molecular geometries and harmonic vibrational frequencies of the other mono- and dioxides. That study reported also the computed dissociation enthalpies and Gibbs free energies. The reliability of the latter data can be assumed to be comparable to that for the monoxides and dioxides of early actinides, where average deviations of ca.  $20 \text{ kJ mol}^{-1}$  for the dioxides and ca.  $50 \text{ kJ mol}^{-1}$  for the monoxides were obtained as compared with available experimental dissociation enthalpies at 298 K. The computed dissociation enthalpies are in good agreement with estimated data of Haire and Eyring<sup>582</sup> using the promotion model. The molecular constants and thermochemical properties are summarized in Tables 86 and 87.

 TABLE 87. Molecular constants and enthalpies of formation for AnO<sub>2</sub>(g) (An = Bk–Lr)

An	$\nu_i$	Symmetry	$r_e$	$\angle(\text{O}-\text{An}-\text{O})$	$I_A$	$\Delta_r H(0 \text{ K})^a$	
						$\text{cm}^{-1}$	$\text{kg m}^2$
BkO <sub>2</sub>	791, 152(2), 725	2	182.0	180.0	$1.75952 \times 10^{-45}$	420	425
CfO <sub>2</sub>	795, 183(2), 716	2	181.7	180.0	$1.75303 \times 10^{-45}$	434	440
EsO <sub>2</sub>	816, 213(2), 738	2	179.5	180.0	$1.71066 \times 10^{-45}$	427	433
FmO <sub>2</sub>	816, 221(2), 730	2	179.1	180.0	$1.70399 \times 10^{-45}$	413	419
MdO <sub>2</sub>	789, 182(2), 703	2	181.2	180.0	$1.74408 \times 10^{-45}$	354	360
NoO <sub>2</sub>	753, 170(2), 668	2	184.3	180.0	$1.80437 \times 10^{-45}$	291	296
LrO <sub>2</sub>	686, 114, 271	2	194.0	101.5	$1.6373 \times 10^{-135c}$	330	334

<sup>a</sup>Bond dissociation  $\text{AnO}_2 \rightarrow \text{AnO} + \text{O}$ .

<sup>b</sup>Neglecting electronic contributions.

<sup>c</sup> $I_A I_B I_C$  for  $\text{LrO}_2$  in  $\text{kg}^3 \text{m}^6$ .

## 7. Discussion and Conclusions

### 7.1. Comparison to existing reviews

The current review has resulted in a consistent set of thermodynamic data for the lanthanide and actinide oxides as summarized in Tables 88 and 89. As mentioned in the Introduction, a number of critical reviews are existing but they are generally restricted to the condensed state and do not include the gaseous oxides, which were treated in detail in the present work. In 1968 Holley Jr. *et al.*<sup>583</sup> published a rather complete review of the crystalline lanthanide oxides based on the emerging literature on these intriguing compounds. The current work has extended that review by providing data in a wider temperature range, and including many more recent literature sources. Although the selected values are generally in good agreement with the recommendations of Holley Jr. *et al.*,<sup>583</sup> differences may exist, particularly for the high temperature properties. During the course of our work Zinkevich<sup>584</sup> published a comprehensive review of the thermodynamics of the rare earth sesquioxides. He also gave rather complete overview of the existing literature, and complemented the known values with estimated data. A major difference between that work and the current results is obvious for the (estimated) properties of the phase transitions. Whereas we have based the estimation of the entropies of transition and fusion on the high pressure work by Hoekstra,<sup>116</sup> the experimental study of  $Gd_2O_3$  by Barkhatov *et al.*,<sup>148</sup> the recent DTA work on  $La_2O_3$  by Ushakov and Navrotsky,<sup>14</sup> and the observation for the lanthanide trihalides that the sum of the transition and melting entropies is fairly constant throughout those series,<sup>160</sup> Zinkevich<sup>584</sup> based his estimations on the measured entropies of fusion for  $Y_2O_3$  and  $Sc_2O_3$  and a comparison to the relation between volume change upon melting and entropy of fusion for the lanthanide metals. These are two different approaches, resulting in somewhat different values, but in absence of experimental data it is difficult to validate them. We think that our approach based on comparison to the (ionic) halides is physically more justified than the comparison to metals.

A comprehensive review for the actinide oxides that treats the condensed and the gaseous phases in a consistent manner does not exist. In the frame of the thermodynamic database project OECD/NEA<sup>430,585–587</sup> the solid oxides of Th, U, Np, Pu and U were reviewed, but with emphasis on the room temperature values. No significant differences exist between our and that work, since the same sources of information have been used, with the exception of the high temperature heat capacity of the oxides of Np and Am for which new measurements were reported recently. The molecular species of some of the major actinides (Th, U, Pu) have been dealt with in general thermochemical assessments<sup>395,588</sup> but the information was very limited at the time those works were completed. Currently quantum chemical computational methods are a main source of molecular data for most actinide oxide species, and are considered to be more reliable than the assumed analogy with the better known

lanthanide compounds. There are four comprehensive quantum chemical studies<sup>460,518,575,589</sup> used in this assessment which covered the mono- and dioxides of early actinides (Th, Pa, U, Np, Pu, Am, Cm), although in some cases the information appeared too late to be included. Quantum chemical studies are also used to calculate dissociation energies of early actinide oxides species and their cations (e.g., Averkiev *et al.*<sup>589</sup>). Although the errors of several experimental values are quite large, the comparison indicates that the quantum chemical computations are sufficiently reliable to predict trends among various actinides but also that the reliability of the computed values still does not reach that of the advanced gas-phase experimental methods.

### 7.2. Trends

The thermodynamic data from the present review allow evaluating the trends in the lanthanide and actinide series, as we have also done for the elements.<sup>590</sup> As is obvious from the sections for the individual compounds, these trends have been used extensively for checking consistency and estimating unknown properties. However, unlike the metals the comparison of the trends in the lanthanide and actinide oxides is less obvious because the electronic configurations are different. Whereas in the lanthanide series the compounds are predominantly trivalent due to the fact that the 4f electrons do not participate in the bonding (localised), this is not the general case for the actinide series. In the light actinides (Th-Am) the 5f electron are delocalised (itinerant) and as a result these elements show a wide range of valence states in compounds. The crystalline sesquioxides are formed in both series for a large number of elements and can be compared very well. The crystalline dioxides are typical for the actinides, and are stable only for a few lanthanides. In the gas phase the LnO and AnO molecules are stable for all elements, the  $AnO_2$  and  $AnO_3$  molecules are again typical for the light actinides (Th-Am). We will shortly discuss the trends in the crystalline sesquioxides, the crystalline dioxides and the gaseous monoxides.

#### 7.2.1. The crystalline sesquioxides

A comparison of the polymorphism of the actinide with the lanthanide sesquioxides is made in Fig. 23, which is a remake of Fig. 4 showing the stability domain of the crystallographic modification not as a function of the position in the lanthanide series, but as a function of the ionic radius. It is evident that the actinide sesquioxides fall in the mid range where the polymorphism is most complex. It can be seen that the polymorphism of  $Cm_2O_3$  fits reasonably well in the  $Ln_2O_3$  series, but the other  $An_2O_3$  compounds show discrepancies: the transition temperatures, particularly the melting points, are significantly lower.

Figure 24 shows the trend in the standard entropy of the sesquioxides, which can be well described as the sum of a lattice component, arising mainly from the vibrations of the

TABLE 88. Selected thermodynamic data of the solid and liquid phases of the lanthanide and actinide oxides

Phase	$\Delta_f H^\circ(298.15)$ kJ mol <sup>-1</sup>	$S^\circ(298.15)$ J K <sup>-1</sup> mol <sup>-1</sup>	$C_p/J\ K^{-1}\ mol^{-1} = A + B \cdot T + C \cdot T^2 + D \cdot T^3 + E \cdot T^{-2}$					Temperature range (K)	$T_{trs}/K$	$\Delta_{trs} H$ kJ mol <sup>-1</sup>
			A	B	C	D	E			
La <sub>2</sub> O <sub>3</sub> (A)	-1791.6 ± 2.0	127.32 ± 0.84	120.6805	13.42414 × 10 <sup>-3</sup>	-	-	-14.13668 × 10 <sup>5</sup>	298-1800	2313 ± 30 (A → H)	23 ± 5
La <sub>2</sub> O <sub>3</sub> (H)	-	-	150	-	-	-	-	2313-2386	2386 ± 30 (H → X)	17 ± 5
La <sub>2</sub> O <sub>3</sub> (X)	-	-	150	-	-	-	-	2386-2577	2577 ± 15 (X → liq)	78 ± 10
La <sub>2</sub> O <sub>3</sub> (liq)	-	-	162	-	-	-	-	>2577	-	-
CeO <sub>2</sub> (cr)	-1090.4 ± 1.0	62.29 ± 0.07	74.4814	5.83682 × 10 <sup>-3</sup>	-	-	-1.29710 × 10 <sup>6</sup>	298-3083	3083 ± 50 (cr → liq)	69 ± 5
CeO <sub>2</sub> (liq)	-	-	120	-	-	-	-	>3083	-	-
Ce <sub>2</sub> O <sub>3</sub> (A)	-1799.8 ± 1.8	148.1 ± 0.4	113.736	28.4344 × 10 <sup>-3</sup>	-	-	-6.41205 × 10 <sup>5</sup>	298-2392	2392 ± 30 (A → H)	28 ± 8
Ce <sub>2</sub> O <sub>3</sub> (H)	-	-	145	-	-	-	-	2392-2406	2406 ± 30 (H → X)	19 ± 5
Ce <sub>2</sub> O <sub>3</sub> (X)	-	-	145	-	-	-	-	2406-2512	2512 ± 50 (X → liq)	85 ± 10
Ce <sub>2</sub> O <sub>3</sub> (liq)	-	-	157	-	-	-	-	>2512	-	-
PrO <sub>2</sub> (cr)	-959.1 ± 2.3	80.8 ± 2.0	72.9881	16.628 × 10 <sup>-3</sup>	-	-	-0.9990 × 10 <sup>5</sup>	298-663	-	-
PrO <sub>1.833</sub> (cr)	-944.6 ± 2.5	79.2 ± 2.0	68.4932	15.9207 × 10 <sup>-3</sup>	-	-	-8.0968 × 10 <sup>5</sup>	298-750	-	-
Pr <sub>2</sub> O <sub>3</sub> (A)	-1809.9 ± 3.0	152.7 ± 0.3	121.6594	25.5611 × 10 <sup>-3</sup>	-	-	-9.8942 × 10 <sup>5</sup>	298-2310	2310 ± 30 (A → H)	28 ± 8
Pr <sub>2</sub> O <sub>3</sub> (H)	-	-	145	-	-	-	-	2310-2397	2397 ± 30 (H → X)	19 ± 5
Pr <sub>2</sub> O <sub>3</sub> (X)	-	-	145	-	-	-	-	2397-2583	2583 ± 25 (X → liq)	88 ± 10
Pr <sub>2</sub> O <sub>3</sub> (liq)	-	-	157	-	-	-	-	>2583	-	-
Nd <sub>2</sub> O <sub>3</sub> (A)	-1806.9 ± 3.0	158.7 ± 1.0	117.1079	28.13655 × 10 <sup>-3</sup>	-	-	-1.25845 × 10 <sup>6</sup>	298-2379	2379 ± 30 (A → H)	29 ± 8
Nd <sub>2</sub> O <sub>3</sub> (H)	-	-	145	-	-	-	-	2379-2477	2477 ± 30 (H → X)	20 ± 5
Nd <sub>2</sub> O <sub>3</sub> (X)	-	-	145	-	-	-	-	2477-2577	2577 ± 15 (X → liq)	88 ± 10
Nd <sub>2</sub> O <sub>3</sub> (liq)	-	-	160	-	-	-	-	>2577	-	-
Pm <sub>2</sub> O <sub>3</sub> (B)	-1811 ± 21	158 ± 5.0	122.9493	30.0141 × 10 <sup>-3</sup>	-	-	-1.85217 × 10 <sup>6</sup>	298-2013	2013 ± 20 (B → A)	6 ± 3
Pm <sub>2</sub> O <sub>3</sub> (A)	-	-	129.454	19.960 × 10 <sup>-3</sup>	-	-	-	2013-2407	2407 ± 20 (A → H)	31 ± 8
Pm <sub>2</sub> O <sub>3</sub> (H)	-	-	165	-	-	-	-	2407-2497	2497 ± 20 (H → X)	20 ± 5
Pm <sub>2</sub> O <sub>3</sub> (X)	-	-	165	-	-	-	-	2497-2592	2592 ± 30 (X → liq)	88 ± 10
Pm <sub>2</sub> O <sub>3</sub> (liq)	-	-	179	-	-	-	-	>2597	-	-
Sm <sub>2</sub> O <sub>3</sub> (C)	-1826.8 ± 4.8	-	132.4358	18.7799 × 10 <sup>-3</sup>	-	-	-2.40860 × 10 <sup>6</sup>	298-900	~ 900 (C → B)	6 ± 3
Sm <sub>2</sub> O <sub>3</sub> (B)	-1823.0 ± 4.0	150.6 ± 0.3	129.7953	19.03114 × 10 <sup>-3</sup>	-	-	-1.86227 × 10 <sup>6</sup>	298-2190	2190 ± 20 (B → A)	7 ± 3
Sm <sub>2</sub> O <sub>3</sub> (A)	-	-	140	-	-	-	-	2190-2395	2395 ± 20 (A → H)	32 ± 8
Sm <sub>2</sub> O <sub>3</sub> (H)	-	-	165	-	-	-	-	2395-2533	2533 ± 30 (H → X)	20 ± 5
Sm <sub>2</sub> O <sub>3</sub> (X)	-	-	165	-	-	-	-	2533-2613	2613 ± 15 (X → liq)	89 ± 10
Sm <sub>2</sub> O <sub>3</sub> (liq)	-	-	179	-	-	-	-	>2613	-	-
Eu <sub>2</sub> O <sub>3</sub> (C)	-1662.5 ± 6.0	135.4 ± 2.0	136.2978	14.9877 × 10 <sup>-3</sup>	-	-	-1.4993 × 10 <sup>6</sup>	298-1350	1350 ± 15 (C → B)	9 ± 3
Eu <sub>2</sub> O <sub>3</sub> (B)	-	-	133.3906	16.6443 × 10 <sup>-3</sup>	-	-	-1.42435 × 10 <sup>6</sup>	298-2327	2327 ± 30 (B → A)	7 ± 2
Eu <sub>2</sub> O <sub>3</sub> (A)	-	-	141	-	-	-	-	2327-2427	2427 ± 30 (A → H)	33 ± 8
Eu <sub>2</sub> O <sub>3</sub> (H)	-	-	144	-	-	-	-	2427-2557	2557 ± 30 (H → X)	21 ± 5
Eu <sub>2</sub> O <sub>3</sub> (X)	-	-	144	-	-	-	-	2557-2622	2622 ± 20 (X → liq)	89 ± 10
Eu <sub>2</sub> O <sub>3</sub> (liq)	-	-	156	-	-	-	-	>2622	-	-
Eu <sub>3</sub> O <sub>4</sub> (cr)	-2276 ± 10	315 ± 4	182.464	26.108 × 10 <sup>-3</sup>	-	-	-	298-2000	-	-
EuO(cr)	-593.2 ± 5.0	83.6 ± 0.8	46.5453	7.360 × 10 <sup>-3</sup>	-	-	1.42047 × 10 <sup>4</sup>	419-1724	-	-
Gd <sub>2</sub> O <sub>3</sub> (C)	-1819.7 ± 3.6	150.6 ± 0.2	114.8086	17.2911 × 10 <sup>-3</sup>	-	-	-1.28397 × 10 <sup>6</sup>	298-2000	1473 (C → B)	9 ± 2
Gd <sub>2</sub> O <sub>3</sub> (B)	-	-	114.6104	15.2344 × 10 <sup>-3</sup>	-	-	-1.24917 × 10 <sup>6</sup>	298-2430	2430 ± 30 (B → A)	6.3 ± 3.3
Gd <sub>2</sub> O <sub>3</sub> (A)	-	-	142	-	-	-	-	2430-2470	2470 ± 30 (A → H)	34.7 ± 3.3
Gd <sub>2</sub> O <sub>3</sub> (H)	-	-	130	-	-	-	-	2470-2538	2538 ± 20 (H → X)	20 ± 5
Gd <sub>2</sub> O <sub>3</sub> (X)	-	-	130	-	-	-	-	2538-2693	2693 ± 15 (X → liq)	92 ± 10
Gd <sub>2</sub> O <sub>3</sub> (liq)	-	-	140	-	-	-	-	>2693	-	-
TbO <sub>2</sub> (cr)	-972.2 ± 5.0	86.9 ± 3.0	73.259	13.2023 × 10 <sup>-3</sup>	-	-	1.0424 × 10 <sup>6</sup>	298-1400	Decomposition	-
Tb <sub>2</sub> O <sub>3</sub> (C)	-1865.2 ± 6.0	159.2 ± 3.0	120.6682	22.17194 × 10 <sup>-3</sup>	-	-	-1.00261 × 10 <sup>6</sup>	298-1823	1823 ± 30 (C → B)	12 ± 4
Tb <sub>2</sub> O <sub>3</sub> (B)	-	-	152	-	-	-	-	1823-2488	2488 ± 30 (B → H)	55 ± 8
Tb <sub>2</sub> O <sub>3</sub> (H)	-	-	170	-	-	-	-	2488-2682	2682 ± 15 (H → liq)	83 ± 8
Tb <sub>2</sub> O <sub>3</sub> (liq)	-	-	182	-	-	-	-	>2682	-	-
Dy <sub>2</sub> O <sub>3</sub> (C)	-1863.4 ± 5.0	149.8 ± 0.15	121.2302	15.27609 × 10 <sup>-3</sup>	-	-	-8.4580 × 10 <sup>5</sup>	298-2223	2223 ± 30 (C → B)	14 ± 5
Dy <sub>2</sub> O <sub>3</sub> (B)	-	-	155	-	-	-	-	2223-2488	2488 ± 30 (B → H)	55 ± 8
Dy <sub>2</sub> O <sub>3</sub> (H)	-	-	173	-	-	-	-	2488-2680	2680 ± 15 (H → liq)	83 ± 8
Dy <sub>2</sub> O <sub>3</sub> (liq)	-	-	188	-	-	-	-	>2680	-	-
Ho <sub>2</sub> O <sub>3</sub> (C)	-1883.3 ± 8.2	156.38 ± 0.15	121.9340	10.11623 × 10 <sup>-3</sup>	-	-	-8.8628 × 10 <sup>5</sup>	298-2538	2538 ± 30 (C → B)	16 ± 5
Ho <sub>2</sub> O <sub>3</sub> (B)	-	-	135	-	-	-	-	2538-2588	2588 ± 30 (B → H)	57 ± 8
Ho <sub>2</sub> O <sub>3</sub> (H)	-	-	149	-	-	-	-	2588-2686	2686 ± 15 (H → liq)	83 ± 8
Ho <sub>2</sub> O <sub>3</sub> (liq)	-	-	162	-	-	-	-	>2686	-	-
Er <sub>2</sub> O <sub>3</sub> (C)	-1900.1 ± 6.5	153.13 ± 0.15	123.2921	8.62245 × 10 <sup>-3</sup>	-	-	-1.54433 × 10 <sup>6</sup>	298-2538	2538 ± 30 (C → H)	25 ± 5
Er <sub>2</sub> O <sub>3</sub> (H)	-	-	162	-	-	-	-	2538-2690	2690 ± 15 (H → liq)	83 ± 5
Er <sub>2</sub> O <sub>3</sub> (liq)	-	-	176	-	-	-	-	>2690	-	-
Tm <sub>2</sub> O <sub>3</sub> (C)	-1889.3 ± 5.7	139.7 ± 0.4	128.4322	5.23209 × 10 <sup>-3</sup>	-	-	-1.17891 × 10 <sup>6</sup>	298-2588	2588 ± 30 (C → H)	26 ± 5
Tm <sub>2</sub> O <sub>3</sub> (H)	-	-	155	-	-	-	-	2588-2682	2682 ± 30 (H → liq)	83 ± 8
Tm <sub>2</sub> O <sub>3</sub> (liq)	-	-	168	-	-	-	-	>2682	-	-
Yb <sub>2</sub> O <sub>3</sub> (C)	-1814.5 ± 6.0	133.1 ± 0.3	130.6438	3.34628 × 10 <sup>-3</sup>	-	-	-1.44820 × 10 <sup>6</sup>	298-2687	2687 ± 20 (C → H)	27 ± 5
Yb <sub>2</sub> O <sub>3</sub> (H)	-	-	134	-	-	-	-	2687-2707	2707 ± 15 (H → liq)	84 ± 8
Yb <sub>2</sub> O <sub>3</sub> (liq)	-	-	146	-	-	-	-	>2707	-	-
Lu <sub>2</sub> O <sub>3</sub> (cr)	-1877.0 ± 7.7	109.96 ± 0.13	122.4593	7.29001 × 10 <sup>-3</sup>	-	-	-2.03414 × 10 <sup>6</sup>	298-2762	2762 ± 15 (C → liq)	113 ± 10
Lu <sub>2</sub> O <sub>3</sub> (liq)	-	-	152	-	-	-	-	>2762	-	-
ThO <sub>2</sub> (cr)	-1226.4 ± 3.5	65.23 ± 0.20	55.9620	5.12579 × 10 <sup>-2</sup>	-3.68022 × 10 <sup>-5</sup>	9.2245 × 10 <sup>-9</sup>	-5.740310 × 10 <sup>5</sup>	298-3500	3651 ± 17 (cr → liq)	88 ± 6
ThO <sub>2</sub> (liq)	-	-	61.8	-	-	-	-	>3651	-	-
γ-UO <sub>3</sub> (cr)	-1223.8 ± 2.0	96.11 ± 0.40	90.2284	1.385332 × 10 <sup>-2</sup>	-	-	-1.12795 × 10 <sup>6</sup>	416-886	-	-



TABLE 88. Selected thermodynamic data of the solid and liquid phases of the lanthanide and actinide oxides—Continued

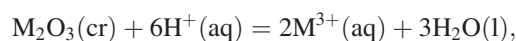
Phase	$\Delta_f H^\circ(298.15)$ kJ mol <sup>-1</sup>	$S^\circ(298.15)$ J K <sup>-1</sup> mol <sup>-1</sup>	$C_p/J\text{ K}^{-1}\text{ mol}^{-1} = A + B \cdot T + C \cdot T^2 + D \cdot T^3 + E \cdot T^{-2}$					Temperature range (K)	$T_{trs}/\text{K}$	$\Delta_{trs} H$ kJ mol <sup>-1</sup>
			A	B	C	D	E			
$\alpha\text{-U}_3\text{O}_8(\text{cr})$	$-3574.8 \pm 2.5$	$282.55 \pm 0.50$	279.267	$2.7480 \times 10^{-2}$	—	—	$-4.3116 \times 10^6$	298–483	483	0.171
			279.267	$2.7480 \times 10^{-2}$	—	—	$-4.3116 \times 10^6$	483–658	658	0.444
			279.267	$2.7480 \times 10^{-2}$	—	—	$-4.3116 \times 10^6$	658–850	850	0.942
			279.267	$2.7480 \times 10^{-2}$	—	—	$-4.3116 \times 10^6$	>850	—	—
$\text{U}_4\text{O}_9(\text{cr})$	$-4512 \pm 7$	$334.1 \pm 0.7$	319.163	$4.9691 \times 10^{-2}$	—	—	$-3.9602 \times 10^6$	298–1600	—	—
$\text{UO}_2(\text{cr})$	$-1085.0 \pm 1.0$	$77.03 \pm 0.20$	66.7437	$4.31393 \times 10^{-2}$	$-3.5640 \times 10^{-5}$	$1.1655 \times 10^{-8}$	$-1.16863 \times 10^6$	298–3000	$3120 \pm 20$ (cr → liq)	$75 \pm 3$
$\text{UO}_2(\text{liq})$	—	—	1365.4956	-0.85866	$1.91305 \times 10^{-4}$	$-1.41608 \times 10^{-8}$	—	3120–5000	—	—
$\text{Np}_2\text{O}_5(\text{cr})$	$-2162.7 \pm 9.3$	$186 \pm 15$	99.2	$9.86 \times 10^{-2}$	—	—	—	298–750	~ 700 (decomposition)	—
$\text{NpO}_2(\text{cr})$	$-1078.5 \pm 2.7$	$80.3 \pm 0.4$	64.7712	$43.8574 \times 10^{-3}$	$-35.0695 \times 10^{-6}$	$13.1917 \times 10^{-9}$	$-0.78932 \times 10^6$	298–3072	$3072 \pm 66 \pm 50$ (cr → liq)	$70 \pm 6$
$\text{NpO}_2(\text{liq})$	—	—	66	—	—	—	—	>3072	—	—
$\text{PuO}_2(\text{cr})$	$-1055.8 \pm 1.0$	$66.13 \pm 0.30$	35.2952	$1.5225 \times 10^{-1}$	$-1.27255 \times 10^{-4}$	$3.6289 \times 10^{-8}$	$-3.47593 \times 10^5$	298–3017	$3017 \pm 28$ (cr → liq)	$64 \pm 6$
$\text{PuO}_2(\text{liq})$	—	—	70	—	—	—	—	>3017	—	—
$\text{Pu}_2\text{O}_3(\text{A})$	$-1647 \pm 10 \pm$	$163.02 \pm 0.65$	130.6670	$1.84357 \times 10^{-2}$	—	—	$-1.70530 \times 10^6$	298–2300	$2300 \pm 50$ (A → H)	$32 \pm 10$
$\text{Pu}_2\text{O}_3(\text{H})$	—	—	165	—	—	—	—	2300–2352	$2352 \pm 10$ (H → liq)	$71 \pm 10$
$\text{Pu}_2\text{O}_3(\text{liq})$	—	—	179	—	—	—	—	>2352	—	—
$\text{AmO}_2(\text{cr})$	$-932.2 \pm 3.0$	$75.5 \pm 3.0$	78.9718	$3.8365 \times 10^{-3}$	—	—	$-1.40591 \times 10^6$	298–1200	—	—
$\text{Am}_2\text{O}_3(\text{A})$	$-1690.4 \pm 7.9$	$134.2 \pm 5.0$	126.0084	$8.0097 \times 10^{-3}$	—	—	$-1.05752 \times 10^6$	298–2350	$2350 \pm 50$ (A → H)	$33 \pm 10$
$\text{Am}_2\text{O}_3(\text{H})$	—	—	141	—	—	—	—	2350–2410	$2410 \pm 50$ (H → X)	$10 \pm 10$
$\text{Am}_2\text{O}_3(\text{X})$	—	—	141	—	—	—	—	2410–2481	$2481 \pm 15$ (X → liq)	$74 \pm 10$
$\text{Am}_2\text{O}_3(\text{liq})$	—	—	156	—	—	—	—	>2481	—	—
$\text{Cm}_2\text{O}_3(\text{B})$	$-1684 \pm 14$	$167 \pm 5$	123.532	$1.4550 \times 10^{-2}$	—	—	$-1.3489 \times 10^6$	298–1888	$1888 \pm 15$ (B → A)	$6 \pm 3$
$\text{Cm}_2\text{O}_3(\text{A})$	—	—	142	—	—	—	—	1888–2273	$2273 \pm 20$ (A → H)	$32 \pm 6$
$\text{Cm}_2\text{O}_3(\text{H})$	—	—	130	—	—	—	—	2273–2383	$2383 \pm 20$ (H → X)	$10 \pm 5$
$\text{Cm}_2\text{O}_3(\text{X})$	—	—	130	—	—	—	—	2283–2543	$2543 \pm 15$ (H → liq)	$66 \pm 10$
$\text{Cm}_2\text{O}_3(\text{liq})$	—	—	140	—	—	—	—	>2543	—	—

ions in the crystal, and an excess component, which is of electronic origin:<sup>127,445</sup>

$$S^\circ = S_{\text{lat}} + S_{\text{exs}} \quad (9)$$

$S_{\text{exs}}$  is zero for the lanthanide ions with empty (La,  $4f^0$ ), half-filled (Gd,  $4f^7$ ) and completely filled (Lu,  $4f^{14}$ )  $f$  sub-shell and thus the lattice contribution can be derived from the experimental data for the sesquioxides of those elements. From the results it is clear that the lattice entropy for the lanthanide sesquioxides falls in two groups, the hexagonal and monoclinic compounds in which the Ln coordination is sixfold, and the cubic compounds in which it is eightfold. This trend has been used in this work to estimate the values for those compounds for which experimental data are missing. Figure 25 shows how the trend for the lanthanides has been extrapolated to the actinides, based on the single experimental value for  $\text{Pu}_2\text{O}_3$ . A similar relation hold for the heat capacity.

The trends in the enthalpies of formation are shown in Fig. 26. We have plotted the hypothetical solution enthalpy of the reaction ( $\Delta_{\text{sln}} H^\circ(298.15\text{ K})$ ),



which represents in part the difference between the lattice enthalpy of the crystalline dioxide and the enthalpy of

hydration of its ionic components. As it can be seen, the results for the lanthanide sesquioxides fall into three groups of isostructural compounds. The few results for the actinides parallel the trend for the lanthanide sesquioxides very well, irrespectively of the crystal structure, suggesting that the reaction enthalpy is about 20–30 kJ mol<sup>-1</sup> less negative than the corresponding lanthanide sesquioxides. Morss *et al.*<sup>454</sup> suggested a relationship between the solution enthalpy of the above reaction and the molar volume (Fig. 27). That correlation captured well the three crystallographic groups of the lanthanide sesquioxides, but it did not show a correlation between the lanthanides and actinides, except for the B-type structure.

## 7.2.2. The crystalline dioxides

The melting point of the actinide dioxides have been revisited by Manara *et al.*<sup>371,404,413</sup> recently. These authors have shown that, due to the high oxygen potential of these compounds at temperatures close to melting, their observed melting/freezing behavior is not independent of the atmosphere in which fusion and solidification occur. Taking into account this important aspect, the trend established with these new data is shown in Fig. 28. It shows a strong decrease from  $\text{ThO}_2$  to  $\text{UO}_2$ , after which the melting temperature decreases further in a moderate way. The absence of data for the melting of  $\text{PaO}_2$  make it difficult the establish the trend in a more reliable way. A similar trend is observed

**THERMODYNAMIC PROPERTIES OF LANTHANIDE AND ACTINIDE OXIDE COMPOUNDS 013101-85**

TABLE 89. Selected thermodynamic data of the gaseous lanthanide and actinide oxides

Phase	$\Delta_f H^\circ(298.15)$ kJ mol <sup>-1</sup>	$S^\circ(298.15)$ J K <sup>-1</sup> mol <sup>-1</sup>	$C_p/J\ K^{-1}\ mol^{-1} = A + B \cdot T + C \cdot T^2 + D \cdot T^3 + E \cdot T^{-2}$					Temperature range (K)
			A	B	C	D	E	
LaO(g)	-119 ± 8	239.594 ± 0.03	28.0550 41.7593	21.9688 × 10 <sup>-3</sup> -6.82858 × 10 <sup>-3</sup>	-1.925691 × 10 <sup>-5</sup> 3.529957 × 10 <sup>-6</sup>	6.083418 × 10 <sup>-9</sup> -2.81197 × 10 <sup>-10</sup>	-1.145313 × 10 <sup>5</sup> -1.500459 × 10 <sup>6</sup>	298-1200 1200-4000
CeO(g)	-132 ± 8	246.099 ± 0.10	22.01944 62.79967	55.5050 × 10 <sup>-3</sup> -1.753953 × 10 <sup>-2</sup>	-4.314095 × 10 <sup>-5</sup> 5.431417 × 10 <sup>-6</sup>	1.089494 × 10 <sup>-8</sup> -4.87448 × 10 <sup>-10</sup>	-4.23189 × 10 <sup>4</sup> -4.947446 × 10 <sup>6</sup>	298-1300 1300-4000
CeO <sub>2</sub> (g)	-538 ± 20	274.417 ± 3.0	37.7646 55.9864	55.1209 × 10 <sup>-3</sup> 2.90431 × 10 <sup>-3</sup>	-57.1816 × 10 <sup>-6</sup> -1.4361 × 10 <sup>-6</sup>	21.2692 × 10 <sup>-9</sup> 0.25254 × 10 <sup>-9</sup>	-2.6077 × 10 <sup>5</sup> -1.11908 × 10 <sup>6</sup>	298-900 900-4000
PrO(g)	-145.5 ± 8	244.367 ± 0.05	22.98903 84.67666	20.3103 × 10 <sup>-3</sup> -2.85857 × 10 <sup>-2</sup>	2.02961 × 10 <sup>-5</sup> 8.73036 × 10 <sup>-6</sup>	-1.46500 × 10 <sup>-8</sup> -8.4566 × 10 <sup>-10</sup>	2.72239 × 10 <sup>5</sup> -1.44324 × 10 <sup>7</sup>	298-1200 1200-4000
NdO(g)	-120 ± 8	242.817 ± 0.05	-9.80368 57.65812	1.88146 × 10 <sup>-1</sup> -9.784651 × 10 <sup>-3</sup>	-1.90285 × 10 <sup>-4</sup> 2.76479 × 10 <sup>-6</sup>	6.23079 × 10 <sup>-8</sup> -2.18778 × 10 <sup>-10</sup>	5.70945 × 10 <sup>5</sup> 4.42849 × 10 <sup>5</sup>	298-1100 1100-4000
PmO(g)	-145 ± 50	246.519 ± 2.0	31.6417 50.9578	3.86653 × 10 <sup>-2</sup> 1.31663 × 10 <sup>-3</sup>	-3.03021 × 10 <sup>-5</sup> -1.57622 × 10 <sup>-6</sup>	8.45852 × 10 <sup>-9</sup> 2.82026 × 10 <sup>-10</sup>	-2.08747 × 10 <sup>5</sup> -2.75930 × 10 <sup>6</sup>	298-1100 1100-4000
SmO(g)	-105.3 ± 8	246.592 ± 0.10	28.65158 70.5862	4.58171 × 10 <sup>-2</sup> -2.02158 × 10 <sup>-2</sup>	-3.44081 × 10 <sup>-5</sup> 4.6805 × 10 <sup>-6</sup>	7.99213 × 10 <sup>-9</sup> -2.43006 × 10 <sup>-10</sup>	-4.99673 × 10 <sup>4</sup> -6.91999 × 10 <sup>6</sup>	298-1400 1400-4000
EuO(g)	-56.5 ± 8	253.419 ± 0.10	33.8838 -79.4911	7.70507 × 10 <sup>-3</sup> 9.34647 × 10 <sup>-2</sup>	-7.38219 × 10 <sup>-6</sup> -2.23194 × 10 <sup>-5</sup>	3.41476 × 10 <sup>-9</sup> 1.8699 × 10 <sup>-9</sup>	-2.52934 × 10 <sup>5</sup> 5.2756 × 10 <sup>7</sup>	298-1700 1700-4000
GdO(g)	-68.0 ± 8	253.495 ± 0.03	21.26451 51.77714	4.09137 × 10 <sup>-2</sup> -1.00325 × 10 <sup>-2</sup>	-3.01655 × 10 <sup>-5</sup> 2.57810 × 10 <sup>-6</sup>	7.50616 × 10 <sup>-9</sup> -1.11668 × 10 <sup>-10</sup>	6.83276 × 10 <sup>4</sup> -4.8038 × 10 <sup>6</sup>	298-1300 1300-4000
TbO(g)	-84.7 ± 10	245.758 ± 0.10	30.97785 64.48191	3.25225 × 10 <sup>-2</sup> -8.31957 × 10 <sup>-3</sup>	-1.96568 × 10 <sup>-5</sup> 8.83659 × 10 <sup>-7</sup>	4.14576 × 10 <sup>-9</sup> 2.70508 × 10 <sup>-11</sup>	-1.18135 × 10 <sup>4</sup> -1.04723 × 10 <sup>7</sup>	298-1400 1400-4000
DyO(g)	-71 ± 20	242.208 ± 0.10	19.08366 68.49712	6.17856 × 10 <sup>-2</sup> -1.13191 × 10 <sup>-2</sup>	-3.7479 × 10 <sup>-5</sup> 1.99999 × 10 <sup>-6</sup>	7.29509 × 10 <sup>-9</sup> -4.18797 × 10 <sup>-11</sup>	2.19805 × 10 <sup>4</sup> -8.39005 × 10 <sup>6</sup>	298-1300 1300-4000
HoO(g)	-57.8 ± 10	244.590 ± 0.10	48.31232 43.11154	7.68332 × 10 <sup>-2</sup> -5.49748 × 10 <sup>-3</sup>	-1.55715 × 10 <sup>-4</sup> 3.07490 × 10 <sup>-6</sup>	7.69567 × 10 <sup>-8</sup> -3.50566 × 10 <sup>-10</sup>	-1.57926 × 10 <sup>6</sup> 4.11955 × 10 <sup>6</sup>	298-900 900-4000
ErO(g)	-32.9 ± 8	256.473 ± 0.10	43.23815 33.60868	-4.71231 × 10 <sup>-3</sup> 3.85434 × 10 <sup>-3</sup>	5.71735 × 10 <sup>-7</sup> -3.82306 × 10 <sup>-7</sup>	7.47515 × 10 <sup>-10</sup> 4.64068 × 10 <sup>-11</sup>	-2.55285 × 10 <sup>5</sup> 2.52559 × 10 <sup>6</sup>	298-1300 1300-4000
TmO(g)	-13.6 ± 8	255.041 ± 0.15	37.63499 39.61651	4.06981 × 10 <sup>-3</sup> -5.40323 × 10 <sup>-4</sup>	-2.78859 × 10 <sup>-6</sup> 1.26385 × 10 <sup>-6</sup>	9.67411 × 10 <sup>-10</sup> -1.43823 × 10 <sup>-10</sup>	-6.69507 × 10 <sup>4</sup> -1.21233 × 10 <sup>6</sup>	298-1700 1700-4000
YbO(g)	-16 ± 10	238.521 ± 0.10	37.70801 32.77497	4.85496 × 10 <sup>-2</sup> 1.73649 × 10 <sup>-2</sup>	-6.84668 × 10 <sup>-5</sup> -5.08804 × 10 <sup>-6</sup>	3.05227 × 10 <sup>-8</sup> 5.03717 × 10 <sup>-10</sup>	-7.73176 × 10 <sup>5</sup> 1.98479 × 10 <sup>6</sup>	298-1000 1000-4000
LuO(g)	-3.4 ± 10	242.089 ± 0.03	28.20132 40.19679	2.04184 × 10 <sup>-2</sup> -7.58989 × 10 <sup>-4</sup>	-1.65159 × 10 <sup>-5</sup> -6.23484 × 10 <sup>-7</sup>	4.71909 × 10 <sup>-9</sup> 2.73798 × 10 <sup>-10</sup>	-1.1965 × 10 <sup>5</sup> -2.75040 × 10 <sup>6</sup>	298-1300 1300-4000
AcO(g)	-193 ± 20	245.545 ± 5.0	26.7681 47.1310	26.00447 × 10 <sup>-3</sup> -13.09869 × 10 <sup>-3</sup>	-24.23966 × 10 <sup>-6</sup> 6.02184 × 10 <sup>-6</sup>	8.10964 × 10 <sup>-9</sup> -0.628330 × 10 <sup>-9</sup>	-0.83961 × 10 <sup>5</sup> -2.91011 × 10 <sup>6</sup>	298-1100 1100-4000
ThO <sub>2</sub> (g)	-435.6 ± 12.6	285.233 ± 2.0	36.83878 56.72155	55.95391 × 10 <sup>-3</sup> 2.163537 × 10 <sup>-3</sup>	-56.33462 × 10 <sup>-6</sup> -1.189606 × 10 <sup>-6</sup>	20.31019 × 10 <sup>-9</sup> 0.227498 × 10 <sup>-9</sup>	-1.94049 × 10 <sup>5</sup> -1.407937 × 10 <sup>6</sup>	298-900 900-4000
ThO(g)	-21.5 ± 5.0	240.071 ± 0.03	30.10768 -13.55069	1.39388 × 10 <sup>-2</sup> 4.45401 × 10 <sup>-2</sup>	-1.15407 × 10 <sup>-5</sup> -8.99062 × 10 <sup>-6</sup>	4.92964 × 10 <sup>-9</sup> 5.83589 × 10 <sup>-10</sup>	-1.86435 × 10 <sup>5</sup> 1.48587 × 10 <sup>7</sup>	298-1500 1500-4000
PaO <sub>2</sub> (g)	-514.0 ± 30	279.711 ± 5	35.4855 57.8335	69.7567 × 10 <sup>-3</sup> 4.88325 × 10 <sup>-3</sup>	-71.6190 × 10 <sup>-6</sup> 6.76950 × 10 <sup>-8</sup>	27.8213 × 10 <sup>-9</sup> -1.07986 × 10 <sup>-10</sup>	-0.58791 × 10 <sup>5</sup> -1.40951 × 10 <sup>6</sup>	298-900 900-4000
PaO(g)	8 ± 30	250.778 ± 10	22.73634 84.42095	1.80345 × 10 <sup>-2</sup> -2.90423 × 10 <sup>-2</sup>	2.8825 × 10 <sup>-5</sup> 8.96146 × 10 <sup>-6</sup>	-2.01289 × 10 <sup>-8</sup> -8.76655 × 10 <sup>-10</sup>	2.78485 × 10 <sup>5</sup> -1.36244 × 10 <sup>7</sup>	298-1100 1100-4000
UO <sub>3</sub> (g)	-795.0 ± 10	310.648 ± 3	46.69199 81.70962	94.6942 × 10 <sup>-3</sup> 2.009478 × 10 <sup>-3</sup>	-94.91701 × 10 <sup>-6</sup> -1.110505 × 10 <sup>-6</sup>	34.1585 × 10 <sup>-9</sup> 0.2162739 × 10 <sup>-9</sup>	-2.793843 × 10 <sup>5</sup> -2.580355 × 10 <sup>6</sup>	298-900 900-4000
UO <sub>2</sub> (g)	-462.1 ± 12	277.027 ± 3	44.35744 59.57586	37.6358 × 10 <sup>-3</sup> 5.392403 × 10 <sup>-3</sup>	-23.15563 × 10 <sup>-6</sup> 0.09463181 × 10 <sup>-6</sup>	5.50268 × 10 <sup>-9</sup> -0.1028723 × 10 <sup>-9</sup>	-0.7485093 × 10 <sup>5</sup> -6.319451 × 10 <sup>5</sup>	298-900 900-4000
UO(g)	21.4 ± 10	252.137 ± 1.0	38.48092 50.04939	3.30187 × 10 <sup>-2</sup> -1.81106 × 10 <sup>-2</sup>	-4.04519 × 10 <sup>-5</sup> 1.23772 × 10 <sup>-5</sup>	1.47496 × 10 <sup>-8</sup> -1.71438 × 10 <sup>-9</sup>	-5.15534 × 10 <sup>5</sup> 2.50947 × 10 <sup>6</sup>	298-1300 1300-4000
NpO <sub>2</sub> (g)	-457 ± 20	269.892 ± 6	56.33269 68.29804	40.03943 × 10 <sup>-3</sup> -9.034032 × 10 <sup>-3</sup>	-54.97771 × 10 <sup>-6</sup> 5.459315 × 10 <sup>-6</sup>	23.02883 × 10 <sup>-9</sup> -0.6628476 × 10 <sup>-9</sup>	-7.32805 × 10 <sup>5</sup> -3.700337 × 10 <sup>5</sup>	298-1000 1000-4000
NpO(g)	-16.6 ± 10	253.060 ± 4	40.73102 53.06105	5.06903 × 10 <sup>-3</sup> 1.44612 × 10 <sup>-3</sup>	5.58835 × 10 <sup>-6</sup> -1.95684 × 10 <sup>-6</sup>	-3.26062 × 10 <sup>-9</sup> 3.62588 × 10 <sup>-10</sup>	-3.95105 × 10 <sup>5</sup> -5.12159 × 10 <sup>6</sup>	298-1400 1400-4000
PuO <sub>3</sub> (g)	-567.6 ± 15	319.450 ± 4	52.89593 76.50587	75.46988 × 10 <sup>-3</sup> 8.903191 × 10 <sup>-3</sup>	-71.42497 × 10 <sup>-6</sup> -1.504422 × 10 <sup>-6</sup>	25.87474 × 10 <sup>-9</sup> 0.07709618 × 10 <sup>-9</sup>	-3.753825 × 10 <sup>5</sup> -16.13944 × 10 <sup>5</sup>	298-900 900-4000
PuO <sub>2</sub> (g)	-428 ± 20	278.741 ± 5	70.29021 49.85911	-13.46001 × 10 <sup>-3</sup> 11.46596 × 10 <sup>-3</sup>	9.065878 × 10 <sup>-6</sup> -1.472281 × 10 <sup>-6</sup>	-1.403447 × 10 <sup>-9</sup> -8.55574 × 10 <sup>-9</sup>	-4.810746 × 10 <sup>5</sup> 41.20745 × 10 <sup>5</sup>	298-1500 1500-4000
PuO(g)	-51.7 ± 15	252.254 ± 3	25.39529 78.23663	5.42560 × 10 <sup>-2</sup> -2.73687 × 10 <sup>-2</sup>	-3.62549 × 10 <sup>-5</sup> 6.77906 × 10 <sup>-6</sup>	6.47005 × 10 <sup>-12</sup> -4.40726 × 10 <sup>-10</sup>	3.72127 × 10 <sup>4</sup> -7.04572 × 10 <sup>6</sup>	298-1200 1200-4000
AmO <sub>2</sub> (g)	-514.0 ± 30	279.464 ± 6	52.7906 54.4906	26.0768 × 10 <sup>-3</sup> 7.17462 × 10 <sup>-3</sup>	-27.1648 × 10 <sup>-6</sup> -1.46280 × 10 <sup>-6</sup>	10.2293 × 10 <sup>-9</sup> +0.13147 × 10 <sup>-9</sup>	-3.3759 × 10 <sup>5</sup> 1.2604 × 10 <sup>6</sup>	298-1000 1000-4000
AmO(g)	-15 ± 50	259.105 ± 5	34.31966 -85.66885	6.13074 × 10 <sup>-3</sup> 1.11574 × 10 <sup>-1</sup>	-6.40603 × 10 <sup>-6</sup> -3.02584 × 10 <sup>-5</sup>	4.00456 × 10 <sup>-9</sup> 2.82855 × 10 <sup>-9</sup>	-2.68922 × 10 <sup>5</sup> 4.35180 × 10 <sup>7</sup>	298-1500 1500-4000
CmO(g)	-75.4 ± 20	259.071 ± 5	26.15923 46.16751	2.30665 × 10 <sup>-2</sup> -3.81910 × 10 <sup>-3</sup>	-1.18949 × 10 <sup>-5</sup> 4.73867 × 10 <sup>-7</sup>	1.83564 × 10 <sup>-9</sup> 1.60603 × 10 <sup>-10</sup>	-3.50044 × 10 <sup>4</sup> -3.86840 × 10 <sup>6</sup>	298-1200 1200-4000

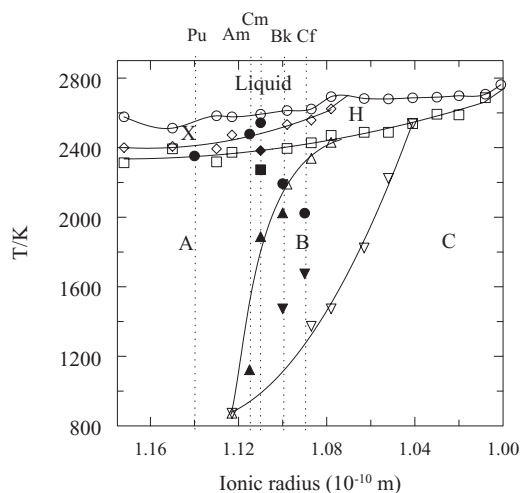


FIG. 23. The polymorphism of  $\text{Ln}_2\text{O}_3$  (open symbols) and  $\text{An}_2\text{O}_3$  (closed symbols) compounds expressed as ionic radius versus temperature. The lines are based on the transition temperatures in the lanthanide series (see Fig. 4).

in other properties of the actinide dioxides, for example the enthalpy of sublimation, which is a good measure for the cohesion energy in the crystalline lattice, shown also in the figure. It suggests a notable influence of the  $5f$  electrons on the bonding. In the lanthanide series, only the melting point of  $\text{CeO}_2$  is known, which is substantially lower than the value of  $\text{ThO}_2$ . Although it is somewhat speculative, this is consistent with the much lower enthalpy of sublimation compared to the actinide dioxides.

The trend in the standard entropies of the actinide dioxides is shown in Fig. 29, which has served as a basis for estimating the values for actinide and lanthanide dioxides for which no experimental data are existing. As is the case for the sesquioxides, the trend in the dioxides can be well described by Eq. (9), estimating the lattice from the measured compounds, in absence of experimental data for the  $5f^7$  compound ( $\text{CmO}_2$ ). Again, in the lanthanide series only the entropy of  $\text{CeO}_2$  has been measured.

The trend in the enthalpy of formation of the solid dioxides has been evaluated from the hypothetical solution

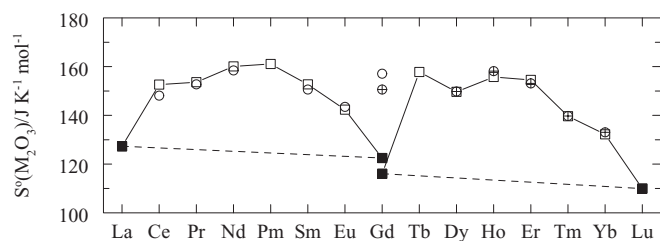


FIG. 24. The standard entropy  $S^\circ(298.15 \text{ K})$  of the lanthanide sesquioxides; ■ the lattice entropies derived from experimental studies; □ values calculated from the lattice, represented by the dashed lines, and excess entropy as explained in the text; ○ and ⊕ the experimental values from the hexagonal/monoclinic and cubic compounds, respectively.

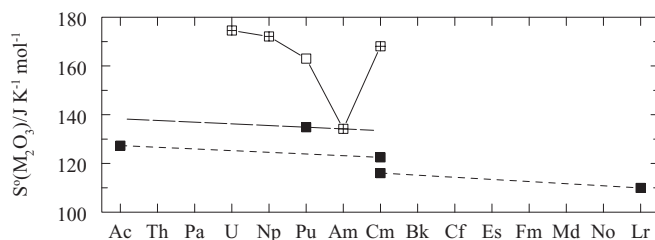
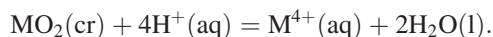


FIG. 25. The standard entropy  $S^\circ(298.15 \text{ K})$  of the actinide sesquioxides; ■ the lattice entropies derived from experimental studies; □ experimental value for  $\text{Pu}_2\text{O}_3$ ; ⊕ estimated values from the lattice and excess entropy as explained in the text.

enthalpy of the reaction



As shown in Fig. 30, the trend for this quantity in the actinide series is quite regular, showing a slightly deviating value for  $\text{NpO}_2$ . The values for the three lanthanide dioxides indicate also a negative slope, but less than the actinides. Morss and Fuger<sup>441</sup> used a different approach by correlating the hypothetical solution enthalpy to the molar volume, as shown in Fig. 31. In this case the lanthanide and actinide dioxides plot almost on a straight line, with the exception of  $\text{TbO}_2$  that deviates significantly. Morss and Fuger<sup>441</sup> suggested this may be due to an erroneous value of the enthalpy of formation of  $\text{Tb}^{4+}(\text{aq})$ , but this remains speculative.

### 7.2.3. The gaseous monoxides

The bond distances of  $\text{LnO}$  and  $\text{AnO}$  molecules are compared in Fig. 32. Those of the  $\text{LnO}$  molecules are mostly experimental data derived from rotationally resolved electronic spectra. In the  $\text{AnO}$  series only the bond distances of  $\text{ThO}$  and  $\text{UO}$  have been determined experimentally (derived similarly from electronic spectra), while the others were taken from the theoretical study of Kovács *et al.*,<sup>498</sup> except  $\text{AcO}$ , which was calculated at the same level of theory in the present study. The theoretical level used in the latter study (B3LYP/small-core relativistic pseudopotentials for the

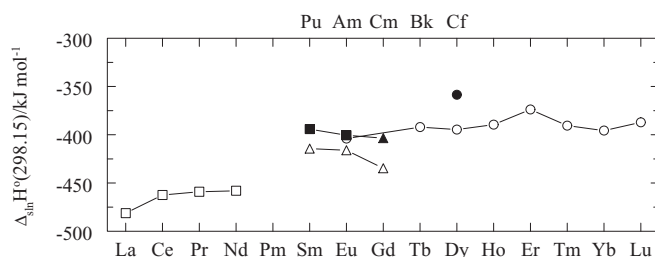


FIG. 26. The enthalpy of the hypothetical solution reaction for the lanthanide (open symbols) and actinide (closed symbols) sesquioxides, indicating the different structures (A-type, □; B-type, △; C-type, ○).

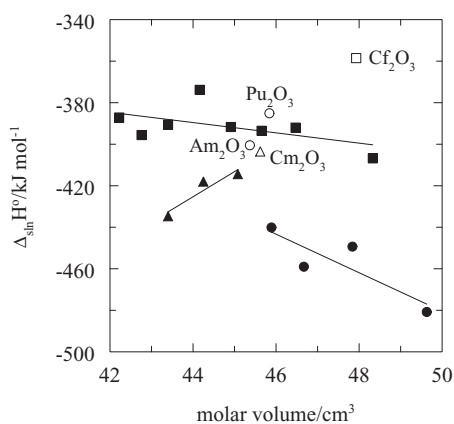


FIG. 27. The enthalpy of the hypothetical solution reaction for the lanthanide (open symbols) and actinide (closed symbols) sesquioxides as a function of the molar volume, indicating the different structures (A-type,  $\square$ ; B-type,  $\blacksquare$ ; C-type,  $\triangle$ ; D-type,  $\circ$ ).

actinides) gave bond distances in excellent agreement (within 0.7 pm) with the experimental data of ThO and UO. A similar reliability may be expected for the other actinide monoxides.

Both the LnO and AnO curves show a double-well shape with some distinct differences. These differences can be attributed to some special features in the electronic structures of the concerned molecules. The very long bond distance in AcO is in agreement with the covalent radii of Ac larger by about 1 pm with respect to that of Th.<sup>591</sup> According to our present population analysis the explanation of the longer bond of AcO may mainly be related to the unpaired 7s electron requiring larger space than the closed 7s<sup>2</sup> subshell in ThO and PaO. On the other hand, in test computations of LaO, CeO, and PrO we observed an open 7s<sup>1</sup> subshell.

The considerably longer bonds toward the end of the actinide row in FmO, MdO, and NoO have been ascribed to substantial population of antibonding orbitals in these molecules.<sup>498</sup> These antibonding orbitals consist of 5f An and 2p O atomic orbitals. Such low-energy antibonding orbitals are not present in the LnO molecules due to the core-like nature of 4f electrons. Therefore the partly filled 4f orbitals have no substantial influence on the bond distance of LnO molecules.

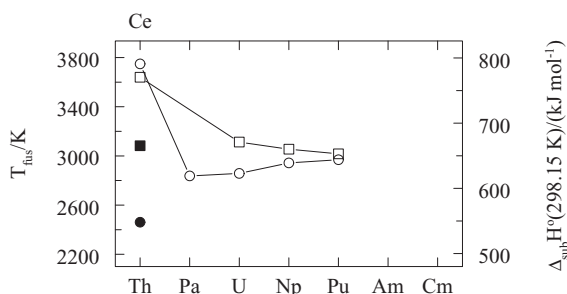


FIG. 28. The melting temperature ( $\square$ ,  $\blacksquare$ ) and the enthalpies of sublimation ( $\circ$ ,  $\bullet$ ) of the actinide (open symbols) and lanthanide (closed symbols) dioxides.

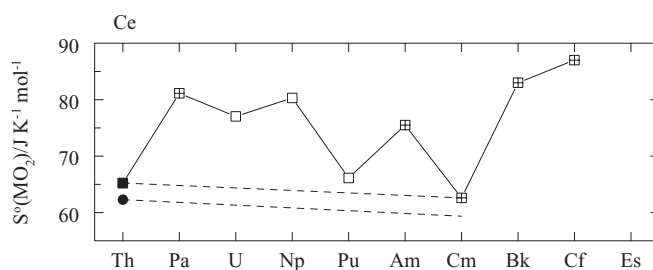


FIG. 29. The standard entropy  $S^\circ(298.15\text{ K})$  of the actinide dioxides;  $\blacksquare$  the lattice entropies derived from experimental studies. The experimental value of  $\text{CeO}_2$  is also shown ( $\bullet$ ).

Population analysis shows that the bonding in EuO is significantly different from the other LnO molecules. While in those other LnO molecules the 4f orbitals have a core-like character (being lower-energy, not mixing with O orbitals and with each other), in EuO the 4f orbitals participate in the bonding orbitals with 2p orbitals of O, and even the non-bonding 4f orbitals are also high-energy (higher than the bonding orbitals). Hence they do not have a core-character in Eu.

The dissociation energy of the lanthanide and actinide monoxides are plotted in Fig. 33. The trends for the two series are very similar. It is generally accepted that this trend can be described by a base energy  $D_0$  and an excess energy  $\Delta E$ ,<sup>260,261,582</sup> arising from the promotion energy from the ground state of the metal to the bonding state in the molecule.

### 7.3. Recommendations for further research

Our review has shown that the thermodynamic properties of the lanthanide sesquioxides are well established in the low to medium temperature range (up to about 2000 K). At high temperature, still large uncertainty exists about the properties of the H, X, and liquid phases. This is still *terra incognita* and is a challenging topic for further research. Additionally, the thermodynamic properties of the intermediate oxides of general formula  $\text{Ln}_n\text{O}_m$  that occur in several of the Ln-O systems, are still poorly known.

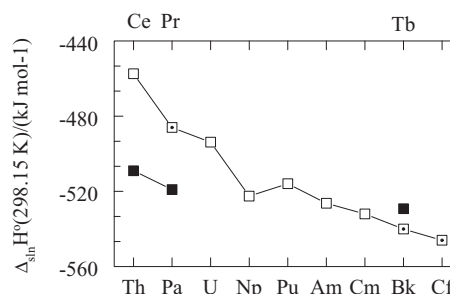


FIG. 30. The enthalpy of formation of the lanthanide ( $\blacksquare$ ) and actinide ( $\square$ ) dioxides.



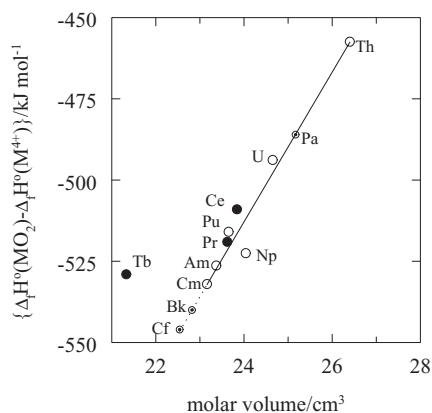


FIG. 31. The enthalpy of formation of the actinide dioxides as a function of molar volume.

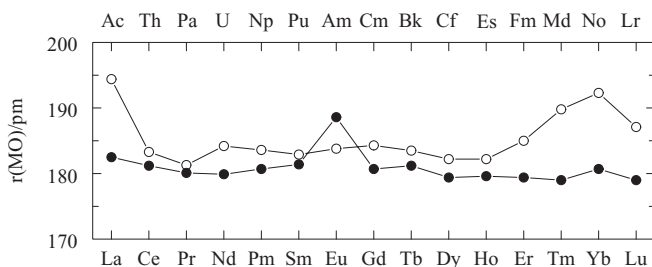


FIG. 32. The interatomic bond distance of the lanthanide (●) and actinide (○) gaseous monoxides.

For the actinide oxides the situation is somewhat different, also because the issue of radioactivity comes into play. The thermodynamic properties of the crystalline dioxides of the major actinides ( $\text{ThO}_2$ ,  $\text{UO}_2$ , and  $\text{PuO}_2$ ) are also well established, even at very high temperatures, though further studies of the liquid phase would be welcomed. The properties of the crystalline oxides of the minor actinides (Np, Am, Cm) are still poorly known, which is in part due to their highly radioactive nature. Some progress has been made in recent years, making use of measuring techniques suitable for small quantities, but also for these compounds high temperature studies are highly needed. Their properties are mainly based on estimation methods that need further validation. The other actinide oxides are hardly studied, and this will probably not change. Improvement for these compounds must be the result of better understanding of the trends in the actinide series.

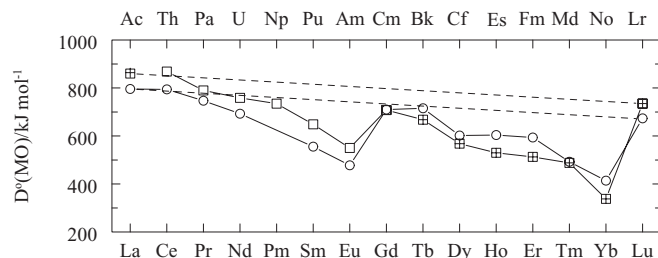


FIG. 33. The dissociation enthalpy of the lanthanide (○) and actinide (□) gaseous monoxides.

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