Temperature effects on the universal equation of state of solids

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Recently it has been argued, based on theoretical calculations and experimental data, that there is a universal form for the equation of state of solids. This observation was restricted to the range of temperatures and pressures such that there are no phase transitions. The use of this universal relation to estimate pressure-volume relations (i.e., isotherms) required three input parameters at each fixed temperature. In this paper we show that for many solids the input data needed to predict high-temperature thermodynamical properties can be dramatically reduced. In particular, only four numbers are needed: (1) the zero pressure (P=0) isothermal bulk modulus, (2) its P=0 pressure derivative, (3) the P=0 volume, and (4) the P=0 thermal expansion; all evaluated at a *single* (reference) temperature. Explicit predictions are made for the high-temperature isotherms, the thermal expansion as a function of temperature, and the temperature variation of the isothermal bulk modulus and its pressure derivative. These predictions are tested using experimental data for three representative solids: gold, sodium chloride, and xenon. Good agreement between theory and experiment is found.

I. INTRODUCTION

Knowledge of the equation of state [the pressure, volume, temperature (P-V-T) relation], based on either calculation or measurement, is of primary importance in both basic and applied sciences. It provides insight into the nature of solid-state theories, and determines the values of fundamental thermodynamic parameters.

Many theoretical and semiempirical calculations have been carried out to describe the isothermal equation of state (EOS). Analytic semiempirical expressions for the EOS for each class of solid abound in a variety of forms,¹ with care often taken to present different forms for different classes of solids.¹⁻³ We have found, however, that there is a universal EOS for all classes of solids in compression and in the absence of phase transitions.⁴ Further, we found that this universal EOS can be predicted from a knowledge of three equilibrium quantities at each fixed temperature T: the equilibrium volume V_0 , isothermal bulk modulus B_0 , and isothermal $(\partial B/\partial P)_0$. Here, we use the notation $(\partial B/\partial P)_0$ for the zero-pressure value of the isothermal pressure derivative of the isothermal bulk modulus, $(\partial B/\partial P)_{T,P=0}$.

In the present study we will show how to predict hightemperature properties of the EOS based on an even smaller amount of input data: the thermal-expansion coefficient α_0, V_0, B_0 and $(\partial B / \partial P)_0$ at zero pressure and at a single (reference) temperature. Using this approach, involving input data at a single temperature, we assumed that the thermal pressure is independent of the volume and linear with temperature above the Debye temperature. This property of near volume independence and linearity with temperature has been found for a number of solids (see, e.g., Refs. 5-8). Based on this assumption the new approach not only predicts the pressure-volume relation at high temperatures, but also the temperature dependence of the thermal expansion, B_0 , and $(\partial B / \partial P)_0$. We will see that rather good accuracy can be obtained with these predictions in comparison with experimental data for some typical solids (a metal, an alkali-halide and a rare-gas solid). Even nonlinear effects in the thermal expansion, which have been difficult to obtain,⁹ are accurately predicted in this approach.

In the following we will first briefly review the observation of a universal EOS for all classes of solids in compression.⁴ Next we will formulate a method to predict high-temperature properties of the universal EOS based only on zero pressure data at a reference temperature. We then present a simple analytical approximation to the universal EOS which is then used as the reference isotherm. The final section will show a set of predictions for the thermodynamic properties of some typical solids (a metal, an alkali-halide crystal, and a rare-gas solid). These predictions will include high-temperature pressure-

<u>35</u> 1945

volume curves, thermal expansion curves, and the temperature dependence of the isothermal bulk modulus and its pressure derivative.

II. A UNIVERSAL EQUATION OF STATE

A universal energy relation between the total energy E and a characteristic distance has been discovered for bimetallic adhesion,¹⁰ chemisorption on metals¹¹ and metallic cohesion¹² (and even then hypothesized for the EOS of nuclear matter¹³). This relation can be written as

$$E(a) = \Delta E E^*(a^*) , \qquad (2.1)$$

where

$$a^* = \frac{(r_{\rm WS} - r_{\rm WSe})}{l}$$
 (2.2)

Here ΔE is the equilibrium binding energy for metallic cohesion, $r_{\rm WS}$ is the Wigner-Seitz radius with a P=0 value of $r_{\rm WSe}$, and *l* is a scaling length, determined by ΔE and the zero-pressure value of d^2E/da^2 (see Ref. 4).

Consider now the classes of solids other than metals. There is some evidence^{12,14} that covalent bonds satisfy Eq. (2.1). However, rare-gas and ionic bonds do not. For energy relations in either compression or expansion, ionic and rare-gas interactions are of inherently different form than the wave-function-overlap interactions¹⁵ characteristic of metallic and covalent bonds. It is these differences that have led to the proposal of different forms for the EOS of different classes of solids.^{1,2}

However, we have found that in compression the form of the pressure-volume relation is dominated by overlap interactions for all classes of solids. That is, at T=0 K, the pressure P(V) has the form

$$P(V) = -dE/dV \tag{2.3}$$

for all classes of solids, where E is the universal form for the energy [Eq. (2.1)] and $V=4\pi r_{\rm WS}^3/3$ is the volume. Thus many different forms are not needed to describe the EOS. Rather, a single universal expression suffices.

In this paper we are primarily interested in temperature effects. For T > 0 Eq. (2.3) becomes

$$P(T,V) = -[\partial F(T,V)/\partial V]_T$$

= $-dE(V)/dV + P^*(T,V)$, (2.4)

where F(T,V) is the Helmholtz free energy, which includes a temperature dependent internal energy and entropy, and $P^*(T,V)$ is the thermal pressure. Now by definition, $P^*(T,V) \rightarrow 0$ as $T \rightarrow 0$. Thus for low T, -dE/dV dominates and one might expect the form of P(T,V) to be similar to that of -dE(V)/dV with the temperature dependence being contained in the scaling parameters. For higher temperatures, $P^*(T,V)$ is not negligible, but in general it is known^{5-8,16,17} to be a weak function of V. In this case, again one might expect that the form of -dE(V)/dV and P(T,V) would be similar with the scaling parameters containing the temperature information. This is because a weakly volume-dependent $P^*(T,V)$ would be expected to shift the equilibrium point, but not effect a substantial change of the shape of the isotherm.

To make this more concrete, we first define the function H as

$$H(V) = \frac{(V/V_0)^{2/3}}{3[1 - (V/V_0)^{1/3}]} P(V) .$$
 (2.5)

This definition is prompted by the form of -dE(V)/dV obtained from the universal energy relation, Eq. (2.1), for metals.¹⁸ Our results for metals¹⁸ suggest that plots of ln*H* versus $1-(V/V_0)^{1/3}$ should be approximately linear, of slope $=\frac{3}{2}[(\partial B/\partial P)_o - 1]$ and intercept $=B_0$, where the subscript 0 refers to equilibrium values. We found⁴ that this is in fact an accurate description of EOS data for many materials from all classes of solids over a broad range of temperatures. This is an experimental confirmation of the expectations expressed above based on the weak volume dependence of $P^*(T, V)$.

As further confirmation, we added a volumeindependent $P^*(T)$ to pressure-volume data⁶ for cesium of strength sufficient to increase or decrease $V_0(T)$ by up to 4%. This thermal volume change and corresponding change in thermal pressure is larger than one could actually encounter in solid cesium. Nevertheless, we found that the linearity of plots of $\ln H(V,T)$ versus $1 - [V/V_0(T)]^{1/3}$ was virtually unchanged by the addition of $P^*(T)$. Values of the correlation coefficient indicating the quality of least-mean-square linear fit, varied from 0.999 95 for the original experimental data to 0.999 92 with the largest added $P^*(T)$. However, the effect of the added $P^*(T)$ was found in changes of corresponding V_0 , B_0 , and $(\partial B/\partial P)_0$ with temperature.

Thus we have a universal form for the EOS of all classes of solids including the effects of temperature. In addition, there is the capability⁴ to predict EOS, requiring only this single form and input equilibrium data: V_0 , B_0 , and $(\partial B / \partial P)_0$ at each temperature. While this is a relatively small amount of information necessary to predict the EOS, in the following section we will derive a predictive scheme which requires zero-pressure data only at a single (reference) temperature.

III. HIGH-TEMPERATURE PREDICTIONS FROM REFERENCE TEMPERATURE ZERO-PRESSURE DATA

The form of the universal EOS allows us to make predictions given V_0 , B_0 , and $(\partial B / \partial P)_0$. Here we would like to show that for some solids, high-temperature thermodynamical properties can be predicted via the universal EOS given zero-pressure data at a single (reference) temperature. [Note that for metals, the universal binding energy relation, Eq. (2.1), provided a means of predicting the thermal expansion coefficient at a single temperature and zero-pressure melting temperatures (Ref. 19)].

The universal form [see discussion following Eq. (2.5) as well as Eq. (4.1)], depends at high temperatures on $P^*(T, V)$ having a much weaker dependence on V than -dE(V)/dV has. This characteristic seems to be generally true (see, e.g., Refs. 5–8, and 17). The new assumptions in this paper are that $P^*(T, V)$ is independent of V and linearly dependent on T for $T \geq \Theta_D$ (where Θ_D is the Debye temperature).

Experimentally, $P^*(T, V)$ can be very often assumed to be independent of V. Examples are the alkali metals (Ref. 6), solid xenon (Refs. 7 and 17), NaCl, LiF, and MgO (see Ref. 8 for the last three). However, for solid argon and krypton the volume dependence of $P^*(T, V)$ was found⁷ to be not negligible, but still much weaker than that of -dE(V)/dV. Thus, where appropriate, one might assume that

$$P^*(T,V) \to P^*(T) . \tag{3.1}$$

In addition, it has been observed experimentally that above the Debye temperature, $P^*(T, V)$ is a linear function of T (see, e.g., Refs. 6–8 and 17). By definition,

$$\left[\frac{\partial P(T,V)}{\partial T}\right]_{V} = \left[\frac{\partial P^{*}(T,V)}{\partial T}\right]_{V} = \alpha_{0}(T,V)B_{0}(T,V) ,$$
(3.2)

where $\alpha_0(T, V)$ is the volumic thermal expansion coefficient. Thus (for $T \ge \Theta_D$) linearity with \overline{T} and Eq. (3.1) implies

$$P^{*}(T) = P^{*}(T_{R}) + \alpha_{0}(T_{R})B_{0}(T_{R})(T - T_{R}), \qquad (3.3)$$

where $T_R \ge \Theta_D$ and T_R is a reference temperature. Combining Eqs. (2.4) and (3.3), we have for $T \ge \Theta_D$,

$$P(T, V) = P(T_R, V) + \alpha_0(T_R)B_0(T_R)(T - T_R) . \quad (3.4)$$

By Eq. (3.4), the high-temperature EOS and corresponding thermodynamical properties can be predicted from a knowledge of zero-pressure properties at the reference temperature T_R . Specifically, $P(T_R, V)$ is specified by $B_0(T_R)$, $V_0(T_R)$ and $(\partial B/\partial P)_0(T_R)$ because of the universal form (Ref. 4) of EOS. The only additional input number is $\alpha_0(T_R)$.

From Eq. (3.4) we find that the isothermal bulk modulus as given by

$$B(T,V) = B(V) = -V \frac{dP(T_R,V)}{dV}$$
(3.5)

and

$$\left[\frac{\partial B}{\partial P}\right]_{T}(T,V) = -\frac{V}{B(V)}\frac{dB(V)}{dV}.$$
(3.6)

The zero-pressure thermal expansion can also be predicted from the EOS, Eq. (3.4), as

$$P(T_R, V) + \alpha_0(T_R)B_0(T_R)(T - T_R) = 0$$
(3.7)

We will now test the predictions from Eqs. (3.4)-(3.7), making use⁴ of the universal EOS for the reference isotherm, $P(T_R, V)$.

IV. ANALYTIC APPROXIMATION TO UNIVERSAL EQUATION OF STATE

The universal EOS has been described in Ref. 4 and Sec. II. It was pointed out there that plots of $\ln H(V,T)$ [Eq. (2.5)] versus $\{1 - [V/V_0(T)]^{1/3}\}$ were approximately linear at each temperature, with slope and intercept yielding $B_0(T)$ and $(\partial B / \partial P)_0(T)$. Thus a simple, analytic approximation (Ref. 4) to the universal EOS is

$$P(T,X) = \frac{3B_0(T)}{X^2} (1-X) \exp[\eta_0(T)(1-X)], \quad (4.1)$$

where

$$X \equiv \left[\frac{V}{V_0(T)}\right]^{1/3} \tag{4.2}$$

and

$$\eta_0(T) = \frac{3}{2} \left[\left(\frac{\partial B}{\partial P} \right)_0(T) - 1 \right].$$
(4.3)

This analytic form, Eq. (4.1), is particularly convenient when used as the reference isotherm since it allows one to make analytic predictions of high-temperature thermodynamical properties through Eqs. (3.4)-(3.7). In terms of Eq. (4.1), Eqs. (3.4)—(3.7) become, respectively,

$$P(T,X) = \frac{3B_0(T_R)}{X^2} (1-X) \exp[\eta_0(T_R)(1-X)] + \alpha_0(T_R)B_0(T_R)(T-T_R), \qquad (4.4)$$

$$B(T,X) = \frac{B_0(T_R)}{X^2} \{2 + [\eta_0(T_R) - 1]X - \eta_0(T_R)X^2\} \exp[\eta_0(T_R)(1 - X)], \qquad (4.5)$$

and

$$\left[\frac{\partial B}{\partial P}\right](T,X) = \frac{4 + [3\eta_0(T_R) - 1]X + \eta_0(T_R)[\eta_0(T_R) - 1]X^2 - \eta_0^2(T_R)X^3}{3\{2 + [\eta_0(T_R) - 1]X - \eta_0(T_R)X^2\}},$$
(4.6)

where, for Eqs. (4.4)-(4.6),

$$X = \left[\frac{V}{V_0(T_R)}\right]^{1/3}.$$
(4.7)

It is appropriate now to compare Eqs. (4.1) and (4.4). As discussed in Sec. III, Eq. (4.4) is an approximation to Eq. (4.1) which is reasonable for $T \ge \Theta_D$ and for those solids for which Eq. (3.1) is a good approximation. The numerical test described in the next to last paragraph of Sec. II suggests that Eqs. (4.1) and (4.4) are consistent with each other to a good approximation. As discussed earlier, the advantage of Eq. (4.4) is that values of $B_0(T)$, $\alpha_0(T)$, and $(\partial B / \partial P)_0(T)$ are required only at $T = T_R$, while for Eq. (4.1), $V_0(T)$, $B_0(T)$ and $(\partial B/\partial P)_0(T)$ are required at all T of interest.

		Gold	NaCl	Xenon	
Input data	$\begin{array}{c} B_0(T_R) \; (10^{10} \; \mathrm{Pa}) \\ \alpha_0(T_R) \; (10^{-5} \; \mathrm{K}^{-1}) \\ (\partial B / \partial P)_0 \; (T_R) \end{array}$	16.6 ^a 4.25 ^d 5.5-6.5 ^a	2.35 ^b 12.0 ^e 5.35 ^b	0.302° 60.0 ^f 7.8°	
Temperatures	$ \begin{array}{l} T_R \ (\mathbf{K}) \\ \Theta_D \ (\mathbf{K})^{\mathbf{g}} \end{array} $	300 162	298 320	60 64	
^a Reference 24. ^b Reference 4.		^e Reference 27, data taken at 293 K. ^f Reference 28.			

^gReference 3.

TABLE I. Summary of the three input parameters as well as Debye temperatures Θ_D and reference temperatures T_R .

^bReference 4.

^cReference 25.

^dReference 26, data taken at 293 K.

V. COMPARISON WITH EXPERIMENT

We will now compare predictions of Eqs. (3.4)-(3.7)with experimental data. The analytic approximation to the universal EOS will be used for the reference isotherm, Eqs. (4.4)-(4.6). Comparison will also be made, where appropriate, with use of the Murnaghan²⁰ EOS for the reference isotherm in Eqs. (3.4)-(3.7). The Murnaghan EOS, which is extensively used (see, e.g., Refs. 21 and 22), has decreasing accuracy (Refs. 4 and 23) as compression increases.

As the universal EOS applies to all classes of solids, we carried out this test for representatives from three classes of solids. These are gold, sodium chloride, and xenon, exhibiting respectively metallic, ionic, and rare-gas bonding.

A. Input parameters

Equations (4.4)-(4.6) require only three input parameters for each solid. These are listed in Table I for the three representative solids, along with selected reference and Debye temperatures.

A few comments must be made: For sodium chloride and xenon, the reference temperatures have been chosen at temperatures slightly lower than the Debye temperatures. This was necessary due to the very limited EOS experimental data, and is not expected to introduce important errors in the predictions of the model. For gold and sodium chloride the values of $B_0(T_R)$ and $(\partial B/\partial P)_0(T_R)$ have been chosen as the average of values taken from the references indicated in Table I.

It is difficult to estimate $(\partial B / \partial P)_0(T_R)$ for gold from the literature. Consequently, we will examine the predictions of our theory for two different values: the lowest one represents most of the static pressure experiments $[(\partial B/\partial P)_0(T_R)=5.5]$ and the highest value represents most of the ultrasonic measurements $[(\partial B/\partial P)_0(T_R)]$ =6.5] (Ref. 24). Values for xenon have been taken from the analysis carried out by Birch (Ref. 25) on data obtained by Anderson and Swenson. Finally, thermal expansion data have been taken from the complete compilations in Refs. 26 and 27 for gold and sodium chloride respectively. The value for xenon has been taken from the study of Manzhelii et al. (Ref. 28).

B. Prediction of high-temperature isotherms

In this section we will estimate high-temperature isotherms near the melting point from input data measured for $T \approx \Theta_D$. The results are then compared with hightemperature isotherms taken from the literature. Figure 1 compares the results for gold with the estimates of Heinz and Jeanloz.²⁴ Their estimates are not direct experimental data but extrapolations of the considered isotherms using a Mie-Gruneisen equation. Heinz and Jeanloz estimate that their results contain a 1-2% error in the pressure. A problem arises from uncertainties in the value of $(\partial B/\partial P)_0(T_R)$. In their analysis of experimental data, Heinz and Jeanloz find a value close to 5.5, however, most



FIG. 1. Prediction of a high-temperature P(V) isotherm: gold at 1000 K. Comparison of predictions based on universal $P(T_R, V)$ [Eq. (4.4)] and on the Murnaghan model for $P(T_R, V)$ with experimental data from Heinz and Jeanloz (Ref. 24).

ultrasonic measurements give values around 6.5 (Refs. 24 and 29-31). It is interesting to note that a fitting of the room-temperature EOS proposed in Ref. 24 to our universal EOS [Eq. (4.4)] would give a value of 5.8. In the present analysis, we show the results for the two extreme values of 5.5 and 6.5 for $(\partial B / \partial P)_0(T_R)$ for gold (Fig. 1). The agreement is good over the entire range of compression and is considerably better than that obtained using the Murnaghan (Refs. 20 and 22) EOS for the reference isotherm in Eqs. (3.4)-(3.7).

In Fig. 2 we compare the results for NaCl with the experimental data of Boehler and Kennedy³² and with data representative of the Decker model.³³ Again, the prediction via Eq. (4.4) is relatively accurate. The prediction based on the Murnaghan reference isotherm is again accurate at low pressures but becomes increasingly inaccurate as the pressure increases as is to be expected.

Finally, solid xenon, as shown in Fig. 3, confirms the results of both Figs. 1 and 2. Again the predictions based on Eq. (4.4) give good agreement when compared to experimental data [Anderson and Swenson (Ref. 7)], while those based on the Murnaghan $P(T_R, V)$ deviate significantly at high compression.

For the three different materials investigated so far, we can conclude that the use of the universal description of the reference isothermal EOS combined with the assumptions of Sec. III gives an accurate prediction of the hightemperature EOS. We have also demonstrated that the



FIG. 2. Prediction of a high-temperature P(V) isotherm: NaCl at 773 K. Comparison of predictions based on universal $P(T_R, V)$ [Eq. (4.4)] and on the Murnaghan model for $P(T_R, V)$ with experimental data from Boehler and Kennedy (Ref. 32) and representative points from the Decker EOS (Ref. 33).



FIG. 3. Prediction of a high-temperature P(V) isotherm: xenon at 159 K. Comparison of predictions based on universal $P(T_R, V)$ [Eq. (4.4)] and on the Murnaghan model for $P(T_R, V)$ with experimental data from Anderson and Swenson (Ref. 7).

Murnaghan EOS cannot describe the reference isotherm well enough to allow extrapolations to high temperatures when substantial compression occurs.

C. Prediction of thermal expansion

In Sec. III, a method was described for obtaining the thermal expansion curve given the reference temperature isotherm [Eq. (3.7)]. In this section, thermal expansion predictions will be compared with experiment. For simplicity, we will limit our study to zero-pressure thermal expansion [Eq. (3.7)] for which experimental data are easily available in the literature. However, we note that the same approach predicts α at high-pressure [$P(V,T) \neq 0$ in Eq. (3.4)], where data are generally not available.

We will use the universal EOS for the reference isotherm, so that from Eq. (4.4),

$$\frac{3}{X^2}(1-X)\exp[\eta_0(T_R)(1-X)] + \alpha_0(T_R)(T-T_R) = 0.$$
(5.1)

Here X is defined by Eq. (4.7) and $\eta_0(T_R)$ by Eq. (4.3), evaluated at T_R . Thus the zero-pressure thermal expansion is determined entirely by $\alpha_0(T_R)$, $V_0(T_R)$, and $(\partial B / \partial P)_0(T_R)$.

For cubic solids, this function of volume can be reduced to a function of length. Figures 4-6 are plots of the predicted values of thermal expansion along with experimental data for gold, sodium chloride, and solid xenon. The two input parameters can be found in Table I. Also, for comparison the linear expansion $\alpha_0(T_R)(T-T_R)/3$ is shown.

A common feature of Figs. 4-6 is that Eq. (5.1) gives a good prediction of thermal expansion for temperatures ranging from the Debye temperature up to the melting point. The nonlinear behavior of the thermal expansion is well reproduced. This is due primarily to the universal EOS. Such nonlinear dependence on temperature has been difficult to predict.⁹

A second common result is that Eq. (5.1) does not accurately predict the thermal expansion for temperatures less than the Debye temperature. Recall that Eq. (3.3) is valid only for $T \ge \Theta_D$ and that Eq. (3.3) was assumed in the derivation of (5.1). Equations (3.1) and (3.3) require that $\alpha_0(T,V)B_0(T,V)$ be independent of T and V. Since the thermal expansion coefficients go to zero at zero temperature, $\alpha_0(T,V)B_0(T,V)$ also $\rightarrow 0$ as $T \rightarrow 0$. Thus it is to be expected that predictions from Eq. (5.1) would not be accurate for low temperatures.

In Fig. 4 we investigated the sensitivity to the input parameter, $(\partial B / \partial P)_0(T_R)$. Using the two extreme values for gold of 5.5 and 6.5, the maximum deviation between predicted thermal expansion of gold is only 3% of $\Delta L / L(T_R)$.

However, Eq. (5.1) is slightly more sensitive to the choice of the other parameter, $\alpha_0(T_R)$. As shown in Fig. 5, a change of about 2.5% in $\alpha_0(T_R)$, consistent with experimental accuracy $[\alpha_0(T_R)=1.2\times10^{-4} \text{ K}^{-1}$ in Ref. 27 and $1.17\times10^{-4} \text{ K}^{-1}$ in Ref. 32 for the same temperature of 300 K], brings maximum variations of the predicted values of about 4% of $\Delta L / L(T_R)$.

D. Temperature dependence of the bulk modulus

In Sec. III we assumed that the isothermal bulk modulus has no explicit dependence on the temperature



FIG. 4. Prediction of thermal expansion of gold: plot of prediction from Eq. (5.1) with smoothed experimental data from Ref. 26. In order to examine how well the nonlinear expansion is predicted, the linear result is plotted as $(T - T_R)\alpha_0(T_R)/3$.



FIG. 5. Prediction of thermal expansion of sodium chloride plotted as in Fig. 4 with smoothed experimental data from Ref. 27.

and is a function of the volume only [Eq. (3.5)]. In Sec. IV, we proposed an analytical form of this function B(V), based on the universal EOS [Eq. (4.5)]. We can now use the predictions of the thermal expansions obtained previously [Eq. (5.1)] to compute the temperature induced volume changes and consequently, the corresponding changes of isothermal bulk modulus. This computation has been done for gold, sodium chloride, and solid xenon and the results are plotted in Figs. 7–9, respectively. Again, the only input parameters are the three listed in Table I.



FIG. 6. Prediction of thermal expansion of solid xenon plotted as in Fig. 4 with experimental data from Ref. 34 ($T \le 100$ K) and Ref. 35 (T > 100 K). (See also Ref. 7).



FIG. 7. Prediction of temperature dependence of the isothermal bulk modulus of gold. Plot of the present model [Eqs. (4.5) and (5.1)] with experimental data from Refs. 30 and 36.

The results achieved with our model agree well with available experimental data over the investigated temperature range. In addition to these general comments, Fig. 7 shows that, for gold, the choice of an input value of 6.5 for $(\partial B / \partial P)_0(T_R)$, which corresponds to ultrasonic measurements, gives the best results. This was to be expected since the experimental data used here to test the bulk modulus prediction (from Refs. 30 and 36) are also obtained from ultrasonic measurements.

In the case of xenon, where low-temperature experimental data are available (Ref. 25), Fig. 9 shows a deviation of the predicted values from the experimental values for temperatures close to 0 K. This was also to be expected since the approximations made in Sec. III do not adequately represent the low-temperature behavior of solids. The slope of the curves $B_0(T)$ must be zero at zero tem-



FIG. 8. Prediction of temperature dependence of the isothermal bulk modulus of sodium chloride: plot of the present model [Eqs. (4.5) and (5.1)] with experimental data from Refs. 32 and 37.



FIG. 9. Prediction of temperature dependence of the isothermal bulk modulus of solid xenon. Plot of the present model [Eqs. (4.5) and (5.1)] with experimental data from Ref. 25.

perature on general thermodynamic grounds. This is confirmed by the experimental data for xenon, but is not represented by our model.

E. Temperature dependence of $(\partial B / \partial P)_0$

The same approach used for the isothermal bulk modulus can be used for $(\partial B / \partial P)_0$ since this quantity is also assumed to have no explicit temperature dependence [Eq. (3.6)]. We use Eq. (4.6) and the previously calculated values of thermal expansion [Eq. (5.1)] for sodium chloride and xenon to obtain a prediction of the thermal behavior of $(\partial B / \partial P)_0$.

Figures 10 and 11 present, respectively, the results for



FIG. 10. Prediction of temperature dependence of $(\partial B / \partial P)_0$ for sodium chloride. Plot of the present model [Eqs. (4.6) and (5.1)] with experimental data from Refs. 37 and 38.



FIG. 11. Prediction of temperature dependence of $(\partial B / \partial P)_0$ for solid xenon. Plot of the present model [Eqs. (4.5) and (5.1)] with experimental data extrapolated from high-pressure measurements in Refs. 7 and 25.

sodium chloride and xenon, obtained by using the three input parameters listed in Table I. These results are compared with experimental data from ultrasonic measurements (sodium chloride) or from analysis of high-pressure EOS data (xenon). Unfortunately, to our knowledge, no similar high-temperature set of data exists for gold.

For sodium chloride, the agreement between the prediction of Eq. (4.6) and the experimental data, as presented in Fig. 10, is good. For xenon (Fig. 11), agreement is worse. However, the accuracy of the experimental data which is extrapolated from static high-pressure measurements is questionable. For example, the two sets of data presented in Fig. 11 are extrapolated from the same set of highpressure experiments, using two different methods (Refs. 7 and 25). As a consequence, it is clear that one must add the uncertainties of the high pressure measurements to the uncertainty caused by the extrapolation method used to obtain the values of $(\partial B/\partial P)_0$. Therefore, a check of our predictions of the thermal variations of $(\partial B/\partial P)_0$ will require more complete and accurate sets of experimental data.

VI. SUMMARY

We have discussed a recently proposed "universal" EOS which accurately describes isothermal P-V data for all classes of solids in compression. Phase transitions were excluded from the analysis. Results from Ref. 4 show that the EOS can be predicted from the universal relation if $V_0(T)$, $B_0(T)$, and $(\partial B / \partial P)_0(T)$ are known for the temperatures of interest.

In this paper we have used the fact that an additional assumption is often valid. Namely, it was assumed that the thermal pressure is independent of V and varies linearly with T for $T \ge \Theta_D$. With solids for which this is an accurate approximation we showed that hightemperature thermodynamical properties can be predicted from $\alpha_0(T_R)$, $V_0(T_R)$, $B_0(T_R)$, and $(\partial B/\partial P)_0(T_R)$, where T_R is a single reference temperature. An analytic approximation was introduced for the universal EOS. This led to a number of simple formulas characterizing the hightemperature EOS in the absence of phase transitions. Simple expressions of particular interest were given for (1) the high-temperature isotherms, (2) the temperature dependence of the thermal expansion for a given P, and (3) the temperature dependence of the P=0 bulk modulus and its pressure derivative. These simple expressions were compared with experiment for gold, sodium chloride, and xenon. Good agreement between prediction and experiment was obtained for the high-temperature isotherms, as well as the temperature dependence of the thermal expansion, bulk modulus, and pressure derivative of the bulk modulus for gold, sodium chloride, and xenon.

Finally, we want to emphasize the wide generality of a model which describes the high-temperature thermodynamic behavior of a metal and an alkali halide as well as a rage-gas solid, from P=0 input data at a single (reference) temperature.

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- ¹B. K. Godwal, S. K. Sikka, and R. Chidambaram, Phys. Rep. **102**, 121 (1983). See especially Table V.
- ²V. N. Zharkov and V. A. Kalinin, Equations of State for Solids at High Pressures and Temperatures (Consultants Bureau-Plenum, New York, 1971).
- ³A. L. Ruoff, and L. C. Chhabildas, in *High Pressure Science and Technology*, edited by K. D. Timmerhaus and M. S. Barber (Plenum, New York, 1979), Vol. 1, p. 19.
- ⁴P. Vinet, J. Ferrante, J. R. Smith, and J. H. Rose, J. Physics C 19, L467 (1986).
- ⁵C. A. Swenson, J. Phys. Chem. Solids 29, 1337 (1968).
- ⁶M. S. Anderson and C. A. Swenson, Phys. Rev. B 31, 668 (1985).
- ⁷M. S. Anderson and C. A. Swenson, J. Phys. Chem. Solids **36**, 145 (1975).
- ⁸O. L. Anderson, R. Boehler, and Y. Sumino, in *High Pressure Research in Geophysics*, edited by S. Akimoto and M. H. Manghnani (Center for Academic Publications, Japan, Tokyo, 1982), p. 273.

- ⁹R. A. MacDonald and R. C. Shulka, Phys. Rev. B 32, 4961 (1985).
- ¹⁰J. H. Rose, J. Ferrante, and J. R. Smith, Phys. Rev. Lett. 47, 675 (1981); J. Ferrante and J. R. Smith, Phys. Rev. B 31, 3427 (1985); J. R. Smith and J. Ferrante, Mater. Sci. Forum 4, 21 (1985).
- ¹¹J. R. Smith, J. Ferrante, and J. H. Rose, Phys. Rev. B 25, 1419 (1982).
- ¹²J. R. Ferrante, J. R. Smith, and J. H. Rose, Phys. Rev. Lett.
 50, 1385 (1983); J. R. Ferrante, J. R. Smith, and J. H. Rose, Phys. Rev. B 28, 1835 (1983).
- ¹³J. H. Rose, J. P. Vary, and J. R. Smith, Phys. Rev. Lett. 53, 344 (1984).
- ¹⁴For a review, see J. R. Smith, J. H. Rose, J. Ferrante, and F. Guinea in *Many-Body Phenomena at Surfaces*, edited by D. Langreth and H. Suhl (Academic, New York, 1984), p. 159.
- ¹⁵W. A. Harrison, *Electronic Structure and the Properties of Solids*, (Freeman, San Francisco, 1980), especially Appendix C.
- ¹⁶See, e.g., Refs. 5-8.
- ¹⁷J. R. Packard and C. A. Swenson, J. Phys. Chem. Solids 24, 1405 (1963).
- ¹⁸J. H. Rose, J. R. Smith, F. Guinea, and J. Ferrante, Phys. Rev. B 29, 2963 (1984).
- ¹⁹F. Guinea, J. H. Rose, J. R. Smith, and J. Ferrante, Appl. Phys. Lett, 44, 53 (1984).
- ²⁰F. D. Murnaghan, *Finite Deformation of an Elastic Solid* (Wiley, New York, 1951), Chap. 4.
- ²¹N. Dass and M. Kumari, Phys. Status Solidi B 127, 103 (1985).
- ²²M. T. Yin and Marvin L. Cohen, Phys. Rev. B 26, 5668

(1982).

- ²³O. L. Anderson, J. Phys. Chem. Solids 27, 547 (1966).
- ²⁴D. L. Heinz and R. Jeanloz, J. Appl. Phys. 55, 885 (1984).
- ²⁵F. Birch, J. Phys. Chem. Solids 38, 175 (1977).
- ²⁶Y. S. Touloukian, R. K. Kirby, R. E. Taylor, and P. D. Dasai, in *Thermal Expansion, Metallic Elements and Alloys*, Vol. 12 of *Thermophysical Properties of Matter*. (IFI/Plenum, New York, 1975).
- ²⁷Y. S. Touloukian, R. K. Kirby, R. E. Taylor, and Tyr Lee, in *Thermal Expansion, Non Metallic Solids, Vol. 13 of Thermophysical Properties of Matter (IFI/Plenum, New York, 1977.)*
- ²⁸V. G. Manzhelii, V. G. Gavrilko, and E. I. Voitovich, Phys. Status Solidi 17, K139 (1966).
- ²⁹W. B. Daniels and C. S. Smith, Phys. Rev. 111, 713 (1958).
- ³⁰S. N. Biswas, P. Van't Klooster, and N. J. Trappeniers, Physica 103B, 235 (1981).
- ³¹G. R. Barsch and Z. P. Chang, Phys. Status Solidi 19, 139 (1967).
- ³²R. Boehler and G. C. Kennedy, J. Phys. Chem. Solids **41**, 517 (1980).
- ³³D. L. Decker, J. Appl. Phys., **42**, 3239 (1971).
- ³⁴C. R. Tilford and C. A. Swenson, Phys. Rev. B 5, 719 (1972).
- ³⁵V. G. Gavrilko and V. G. Manzhelii, Fiz. Tverd. Tela (Leningrad) **6**, 2194 (1964) [Sov. Phys.—Solid State **6**, 1734, (1965)].
- ³⁶Y. A. Chang and L. Himmel, J. Appl. Phys. 37, 3567 (1966).
- ³⁷H. Spetzler, C. G. Sammis, and R. J. O'Connell, J. Phys. Chem. Solids 33, 1727 (1972).
- ³⁸R. A. Bartels and D. E. Schuele, J. Phys. Chem. Solids 26, 537 (1965).