Computation of collective modes and acoustic investigations at different temperatures of vitreous silica

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MS received 6 May 1994

Abstract. This paper is mainly concerned with elastic and acoustic properties of vitreous silica besides the computation of phonon frequencies. Thus the phonon frequencies of vitreous silica have been calculated assuming the electronic bulk modulus, $K_e$, as equal to zero. New equations have been derived to relate the pressure derivatives of second order elastic constants to the acoustic Gruneisen's parameters using both Bhatia–Singh's parameters and Schofield's equations. The calculated longitudinal and transverse Gruneisen's parameters and the predicted absorption band spectra from Nagendranath's equation and Bhatia Singh's parameters are in good agreement with experiment. The calculated mean acoustic mode Gruneisen's parameter evaluated from the pressure derivative of Nagendranath's equation is also in good agreement with experiment.

Keywords. Phonon frequencies; pressure derivatives of elastic constants; longitudinal and transverse Gruneisen's constants; absorption band.

PACS No. 62·20

1. Introduction

Recently, various aspects of vitreous silica have been investigated [1–5]. Some of the important properties that have been investigated are the acoustic properties [1], neutron diffraction studies [2–4], velocity measurements [5] of densified $a$-$SiO_2$ etc. Vitreous silica is of great technical importance. It is a pure, single-phase glass and its non-linear acoustic properties are singularly anomalous. The temperature dependence of elastic stiffness constants, as measured by Wang et al [6] are negative up to 70 K and then become positive. Recently the pressure dependence of longitudinal sound wave velocity in vitreous silica has been measured [7] through Brillouin spectroscopy between 50 K and room temperature. It is found that there is a reduction in velocity by the application of pressure. Negative pressure derivatives of second order elastic stiffness tensor components and positive third order elastic constants (TOEC) reveal anomalous effects in the vibrational anharmonicity while the mode Gruneisen's parameter's are negative [6]. Hence a knowledge of acoustic mode Gruneisen constant is important in understanding low temperature thermal expansion. Hence we felt a theoretical investigation of phonons, acoustic and elastic properties of vitreous silica is important. As pointed out earlier $SiO_2$ is remarkable in the sense that the pressure derivatives of second order elastic constants (SOEC) and also that of bulk modulus is negative unlike in normal substances like $TeO_2$ glass [8], amorphous As [9] and $As_2S_3$ [10]. Even though the SOEC pressure derivatives are negative it is remarkably isotropic over a wide temperature range. Vitreous silica has been shown to possess tetra-
hedralk structure [11, 12] and the average number of nearest neighbours has been estimated [11] to be 4.3. Extensive elastic and acoustic work has been done on vitrous silica. We give only the reference to the recent studies of Wang et al [6] in which other references can be found. Wang et al made a detailed experimental study of the SOECs, their pressure derivatives as well as the acoustic Gruneisen's constants as a function of temperature. We also derive the phonon frequencies through Bhatia–Singh's (BS) method [13].

2. Theory

A. Computation of phonon frequencies

Bhatia and Singh [13] derived the following expressions for the longitudinal and transverse phonon frequencies

\[ \rho \omega^2_L(k) = \frac{2n_e}{a^2} (\beta I_0 + \delta I_2) + \frac{K_eK_{TF}^2 k^2 [G(kr_s)]^2}{k^2 + K_{TF}^2 \tilde{g}(k)}. \] (1)

\[ \rho \omega^2_T(k) = \frac{2n_e}{a^2} \left[ \frac{\beta}{2} (I_0 - I_2) \right]. \] (2)

Here \( n_e \) stands for the number of nearest neighbours, \( a \) is the nearest neighbour distance, \( K_{TF} \) is the Thomas–Fermi inverse screening length, \( K_e \) is the electronic bulk modulus, \( r_s \) the Wigner–Seitz radius, \( G(kr_s) \) is the shape factor and \( \tilde{g}(k) \) is the Linhard–Langer–Vasko expression for the \( k \) dependent dielectric constant. Since these expressions have been discussed earlier [14, 15] we do not give any more details here. Since vitrous silica is not a semiconductor and is a dielectric we take \( K_e \) the electronic bulk modulus to be equal to zero and calculate \( \omega_L(k) \) and \( \omega_T(k) \) from (1) and (2). To evaluate \( \beta \) and \( \delta \) we use the experimental values of \( C_{11} \) and \( C_{44} \) as given by Wang et al [6].

The equations are [13]

\[ C_{11} = n_e \left[ \frac{1}{3} \beta + \frac{1}{5} \delta \right], \] (3)

\[ C_{44} = n_e \left[ \frac{1}{3} \beta + \frac{1}{15} \delta \right], \] (4)

where

\[ \beta = \frac{\rho a^2}{2M} \left[ \frac{1}{r} \frac{d\phi}{dr} \right]_{r=a}, \] (5)

\[ \delta = \frac{\rho a^3}{2M} \left[ \frac{1}{r} \frac{d}{dr} \left( \frac{1}{r} \frac{d\phi}{dr} \right) \right]_{r=a}. \] (6)

Here \( \phi \) is the pair potential. We calculate the phonon frequencies at four different temperatures.
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B. Derivation of the pressure derivatives of elastic constants from Bhatia–Singh’s β and δ parameters

\[
\frac{dC_{11}}{dP} = n_c \left[ \frac{1}{3} \frac{d\beta}{dP} + \frac{1}{5} \frac{d\delta}{dP} \right]
\]

\[
\frac{dC_{44}}{dP} = n_c \left[ \frac{1}{3} \frac{d\beta}{dP} + \frac{1}{15} \frac{d\delta}{dP} \right].
\]

From (6) it can be shown that

\[
\frac{d\delta}{dP} = -\frac{d\beta}{dP} + \frac{d^2\psi}{2M dP} \left( \frac{d^2 \phi}{dr^2} \right)_{r=a} + \left( \frac{d^2 \phi}{dr^2} \right)_{r=a} \frac{d}{dP} \left( \frac{\rho a^2}{2M} \right).
\]

\[
\frac{d}{dP} \left( \frac{d^2 \phi}{dr^2} \right)_{r=a} = \left[ \frac{d}{dV} \left( \frac{d^2 \phi}{dr^2} \right) \frac{dV}{dP} \right]_{r=a}.
\]

\[
= \left[ -\chi_r r \frac{d^3 \phi}{3dr^3} \right]_{r=a}.
\]

Here \(\chi_r\) is the isothermal compressibility. Remembering that the Gruneisen’s constant \(\gamma_s\) is given by [16, 17]

\[
\gamma_s = -\frac{a}{6} \left( \frac{d^3 \phi}{dr^3} \right)_{r=a} \left( \frac{d^2 \phi}{dr^2} \right)_{r=a}
\]

one can show that

\[
\frac{d\delta}{dP} = -\frac{d\beta}{dP} + 2\chi_r (\delta + \beta) \left( \gamma_s + \frac{1}{6} \right).
\]

Similarly it can be shown that

\[
\frac{d\beta}{dP} = -\frac{\rho a^2 \chi_r \left( d^2 \phi \right)_{r=a}}{6M} + \frac{\rho a \chi_r \left( \frac{d\phi}{dr} \right)_{r=a}}{3M}.
\]

Thus finally we obtain

\[
\frac{d\delta}{dP} = 2\chi_r \left[ \frac{\delta}{3} + (\beta + \delta) \gamma_s \right],
\]

and

\[
\frac{d\beta}{dP} = -\frac{\chi_r (\beta - \delta)}{3}.
\]

Hence from (8) we get

\[
\frac{dC_{11}}{dP} = n_c \chi_r \left[ \frac{\delta + 5\beta}{45} + \frac{2\gamma_s (\beta + \delta)}{5} \right].
\]
In an exactly similar way we obtain

$$\frac{dC_{44}}{dP} = n_r x_T \left[ \frac{1}{9}(\beta - 0.6\delta) + \frac{2}{15}y_T(\beta + \delta) \right].$$

(18)

Equations (17) and (18) are important and offer a method in the evaluation of the pressure derivatives of the SOECs. Vitreous silica is a highly isotropic substance and we can obtain $dC_{12}/dP$ from the following equation

$$\frac{2dC_{44}}{dP} = \frac{dC_{11}}{dP} - \frac{dC_{12}}{dP}.$$  \hspace{1cm} (19)

In this connection it is important to note that liquids in the high frequency region show rigidity which is a characteristic of solids [18, 19]. Hence it will be interesting and tempting to try the equations of Schofield [20] (Sch) to derive the pressure derivatives of SOECs. From Schofield’s equations [20] we derive the following expressions for the pressure derivatives of SOECs as [21]

$$\frac{dC_{11}}{dP} = C_{11} x_T + 1.8(C_1 - 1).$$  \hspace{1cm} (20)

$$\frac{dC_{44}}{dP} = C_{44} x_T + 0.6(C_1 - 1).$$  \hspace{1cm} (21)

$$\frac{dC_{12}}{dP} = C_{12} x_T + 0.6(C_1 - 1).$$  \hspace{1cm} (22)

Here

$$C_1 = (dB/dP)$$  \hspace{1cm} (23)

and $B$ is the bulk modulus.

It is important to remember that $C_1$ is related to third order elastic constants [22]. Thus (20)–(22), even though can strictly be used for fluids can help us to estimate the pressure derivatives of the SOECs. Calculations for computing the pressure derivatives of SOECs have been made for vitreous silica at different temperatures using (17)–(22).

C. Prediction of band absorption spectra of vitreous silica

It was shown that [23–25]

$$\frac{8A(A + 8C_{11} - 16C_{44})}{(3A - 8C_{11} + 16C_{12})^2} = 1,$$

(24)

where

$$A = \omega^2 M/2a.$$  \hspace{1cm} (25)

From (24) and (25) it is possible to solve for the frequency and hence predict the position of the absorption band.

From (6) we get

$$\delta = -\beta + \frac{\rho a^2 \left( \frac{d^2 \phi}{dr^2} \right)_{r=a}}{2M},$$  \hspace{1cm} (26)
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and we can calculate \( (d^2 \phi / dr^2)_{r=a} = K_1 \). With a three-force constant model it is possible to show that for a diamond type of lattice \([25]\)

\[
\omega^2 = \frac{4K_1}{M},
\]

and we have \(1/\lambda = \omega / 2\pi C\) where \(C\) is the velocity of light. Hence we can predict the position of the absorption band and can make a comparison of these results with experiment. Such calculations have been made for vitreous silica in these investigations.

D. Methods of evaluation of acoustic Gruneisen’s parameters \(\gamma_g^T\) and \(\gamma_g^L\)

From (17) we get \(C_1\) associated with \(\gamma_g^L\) the Gruneisen’s constant of longitudinal wave while from (18) we get \(\gamma_g^T\) that of the transverse wave. Similarly we use (20) and (21).

The mean acoustic mode Gruneisen’s parameter \(\gamma_g^{el}\) can be obtained \([6]\) from the following equation

\[
\gamma_g^{el} \left( \frac{\gamma_g^L}{V_L^3} + \frac{2\gamma_g^T}{V_L^3} \right) \left( \frac{1}{V_L^3} + \frac{2}{V_T^3} \right)^{-1}.
\]

We now give a method of evaluation of \(\gamma_g^{el}\) using (24) and (25). We differentiate (24) with respect to pressure and solve for \((dA/dP)_T\). Thus we obtain

\[
\frac{dA}{dP} = \left\{ \frac{dC_{11}}{dP} \left[ 56A + 128C_{12} - 64C_{11} \right] - 64A \left( \frac{dC_{44}}{dP} \right) + \right. \\
\left. \frac{dC_{12}}{dP} \left[ 128C_{11} - 48A - 256C_{12} \right] \right\} \left[ A - 56C_{11} + 48C_{12} + 64C_{44} \right]^{-1}.
\]

(29)

Now we eliminate \(dC_{12}/dP\) using the isotropy condition (vide (19)) and finally obtain

\[
\frac{dA}{dP} = \left\{ 8 \left( \frac{dC_{11}}{dP} \right) (3A - 8C_{11} + 16C_{12}) + 16 \left( \frac{dC_{12}}{dP} \right) \\
(8C_{11} - 16C_{12} - A) \right\} \left[ A - 24C_{11} + 16C_{12} \right]^{-1}.
\]

(30)

Further we have

\[
A = K_2 \omega^2
\]

(31)

where

\[
K_2 = M/2a
\]

(32)

From (31) we get

\[
\frac{dA}{dP} = \beta_T A \left[ 2\gamma_g^{el} + \frac{1}{3} \right],
\]

(33)

where we assumed that \(v = kr^3\). Here \(v\) is the volume and \(r\) is the radius. Hence from (30)–(33) we can solve the mean acoustic mode Gruneisen parameter \(\gamma_g^{el}\).
3. Results and discussion

As pointed out already an application of the various equations derived above have been applied to vitreous silica. The experimental results used are those of Wang et al [6]. In figure 1 we give the phonon frequencies as obtained from (1) and (2) at two different temperatures namely 273 K and 77 K. As pointed out already vitreous silica is a dielectric and hence it is assumed that $K_e = 0$. The $\beta$ and $\delta$ values calculated from (3) and (4) at the four temperatures are given in table 1. The position and maximum values of $\omega_L(k)$ and $\omega_T(k)$ are given in table 2. It may be noted from table 1 that

![Graph showing Phonon frequencies at 77 and 273 K.](image)

**Figure 1.** Phonon frequencies at 77 and 273 K.

**Table 1.** Parameters obtained for $\beta$ and $\delta$ at different temperatures.

<table>
<thead>
<tr>
<th>Temp. K</th>
<th>$\beta \times 10^{12}$ dyn/cm$^2$</th>
<th>$\delta \times 10^{12}$ dyn/cm$^2$</th>
</tr>
</thead>
<tbody>
<tr>
<td>273</td>
<td>0.0469</td>
<td>0.835</td>
</tr>
<tr>
<td>233</td>
<td>0.0406</td>
<td>0.8383</td>
</tr>
<tr>
<td>123</td>
<td>0.0380</td>
<td>0.8311</td>
</tr>
<tr>
<td>77</td>
<td>0.0352</td>
<td>0.832</td>
</tr>
</tbody>
</table>

$K_e = 0$ and $\rho_w = 2.202$ g/cm$^3$; $n_e$ [11] = 4.3; $a$ [11] = 1.61 Å

**Table 2.** Characteristics of phonon frequencies at different temperatures.

<table>
<thead>
<tr>
<th>Temp. (K)</th>
<th>Position and maximum value $\omega_L(k)$ $10^{13}$s$^{-1}$</th>
<th>Position and maximum values $\omega_T(k)$ $10^{13}$s$^{-1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>273</td>
<td>2.4, 9.03</td>
<td>3.4, 7.3</td>
</tr>
<tr>
<td>233</td>
<td>2.4, 8.98</td>
<td>3.4, 7.24</td>
</tr>
<tr>
<td>123</td>
<td>2.4, 8.92</td>
<td>3.4, 7.19</td>
</tr>
<tr>
<td>77</td>
<td>2.4, 8.89</td>
<td>3.4, 7.1</td>
</tr>
</tbody>
</table>

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\( \beta \neq 0 \) and hence central forces are not alone operative. Evidently the tetrahedral angular forces are contributing and hence \((d\phi/dr)_{r=a} \neq 0\). The position of the maximum phonon frequency does not change with temperature both for \(\omega_\gamma(k)\) and \(\omega_T(k)\). However with increase in temperature the frequency increases slightly. As expected \(\omega_\gamma(k)\) attains a maximum at a small value of \(k\) and oscillates showing collective mode propagation for larger values of \(k\) while \(\omega_T(k)\) attains almost a constant value beyond the maximum. From \(\beta\) and \(\delta\) given in table 1 and at a temperature 273 K we get \(\omega = 11.1 \times 10^3 \) s\(^{-1}\) through the use of (27) and the position of the absorption band should appear at 583 cm\(^{-1}\). The value of \(A\) obtained from (24) is \(31.7 \times 10^{12}\) dyn/cm\(^2\) and from (25) we get the absorption band of vitrous silica to appear at 536 cm\(^{-1}\). The observed bands for vitrous silica as reported by Lippincott et al and Hass [26, 27] appear between 500 cm\(^{-1}\) and 600 cm\(^{-1}\) which are reasonably close to the calculated values obtained from \(A\) and \((d^2\phi/dr^2)_{r=a}\). The average value being 560 cm\(^{-1}\). According to Bell and Dean [28] this vibrational mode arises from bond bending vibration in which the oxygen atoms move along the bisectors of Si–O–Si angles. In this connection the absorption band values of 536 cm\(^{-1}\) and 583 cm\(^{-1}\) correspond to a frequency \(10 \times 10^5\) s\(^{-1}\) and \(11.0 \times 10^5\) s\(^{-1}\). These values can be considered to be close to each other. From the observed pressure derivatives we obtain (vide (20) and (21)) \(C_L^T\) corresponding to \(dC_{11}/dP\) and \(C_T^T\) to \(dC_{44}/dP\). We now use the well-known Slater's equation [29]

\[
C_L^{T} = 2\gamma^L_s + \frac{1}{3}
\]

(34)

We evaluate \(\gamma^L_s\) and \(\gamma^T_s\) at 273 and 77 K using (17) and (18) and also from (20) and (21). The values calculated are given in table 3. It is important to stress that \(\gamma^L_s\) and \(\gamma^T_s\) are related to third order elastic constants [30].

It is known that Bhatia–Singh equation is meant for amorphous solids and gives very good results in agreement with experiment both at low and high temperatures for which results are available. The application of Schofield's equations gives satisfactory results as well. It should be borne in mind that Schofield's equations are meant for liquids and hence the present agreement can be considered as very good.

We can calculate from (30)–(33) the mean acoustic mode Gruneisen's parameter \(\dot{\gamma}^{el}_s\) at 273 K. Using the experimental values of SOECs we obtain a value of \(31.7 \times 10^{12}\) dyn/cm\(^2\) for \(A\) from (24) and finally we get a value for \(\dot{\gamma}^{el}_s = -2.2\) where we used (30)–(33). The experimental value is \(-2.0\). Thus the agreement is excellent.

Table 3. Gruneisen constants evaluated by different methods at different temperatures.

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<tbody>
<tr>
<td>77</td>
<td>-5.4</td>
<td>-5.9</td>
<td>-5.8</td>
<td>-2.4</td>
<td>-3.6</td>
<td>-2.8</td>
</tr>
<tr>
<td>273</td>
<td>-2.4</td>
<td>-3.0</td>
<td>-2.5</td>
<td>-1.8</td>
<td>-2.9</td>
<td>-2.1</td>
</tr>
</tbody>
</table>

4. Conclusions

Using Bhatia–Singh's equation and assuming $K_e$, the electronic bulk modulus, equal to zero the phonon frequencies have been derived. A change of temperature over a wide range does not effect the position of the maximum while its magnitude decreases only slightly with decreasing temperature. Equations for the pressure derivatives of SOECs have been derived using both Bhatia–Singh's and Schofield's equations. Using the experimental values of the pressure derivatives of the SOECs the longitudinal and transverse Gruneisen's constants have been calculated at two extremely different temperatures for which accurate measurements are available. The results calculated from Bhatia–Singh's parameters give excellent results in agreement with experiment while those obtained from Schofield's equation are also in good agreement with experiment. Using (24) the absorption band spectrum of vitrous silica has been calculated and compared with that obtained from Bhatia–Singh's 's' parameter. Both the results are in fair agreement with experiment for SiO$_2$. Finally we calculate the mean acoustic mode $\gamma^s$ from the pressure derivatives of (24). The value obtained is in excellent agreement with experiment.

Acknowledgements

The authors are thankful to the DST, CSIR, Government of India for financial support.

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