

s with the nuclei becomes in-
 e substance may be regarded
 perfect Fermi electron gas.
 of dK/dp in the nonrelativistic
 relativistic approximations is
 tively. It should be noted th
 ply only at extremely high pre
 g to the same source, the valu
 atm $\gg p \gg 5 \times 10^{17}$ atm
 e average atomic number of th
 as the value $4/3$ is for $p \gg 10^{17}$
 a ($Z = 11$) the inequality be
 comes $10^{17} \gg p \gg 1.5 \times 10^{17}$
 is far above the range of any
 a and probably even above th
 trapolations are needed! Th
 pressure found in compiling th
 ons is a shock wave point
 for aluminum oxide. Birch
 ated the pressure at the cen
 re of the order of 3.4 Mb. In
 sidered normal for dK/dp to
 n a monotone fashion as th
 s. Equation 2 provides the
 or, but the leveling off of
 a few per cent of the value
 ssures p of the order of $10aK_0$
 able values of a , is very low
 10^{17} atm. Therefore, in order
 eted behavior over the pres
 he extrapolation is desired, it
 t the best m to use in equ
 substantially larger than $5/3$
 ensitive point, however, sine
 or C) remains undetermined
 ue of the second derivative
 3, can still be adjusted by

if C and K_0' both positive we used

$$m - K_0' = \frac{C}{2[K_0' - (2C)^{1/2}]} \quad (4)$$

This expression was obtained as an approxima-
 tion to the smallest value of m that allowed K
 to fall to zero on $-a < P < 0$, assuming
 $C \ll K_0'^2$. The condition that K drop con-
 tinuously to zero on $P < 0$ may be regarded as
 an 'instability' condition. It is satisfied auto-
 matically whenever $K_0' > 0$ and $C < 0$. It is
 not considered essential for the purpose of extrap-
 olating on $P > 0$, but in the absence of any
 other guidance, it seemed to be a reasonable
 criterion for relating the two adjustable param-
 eters, say m and C , when K_0' and C are both
 positive. The idea that m should be near the
 smallest value that allows this instability fol-
 lows from the feeling that the condition $m >$
 K_0' , needed to avoid a singularity on $P > 0$ in
 this case, is likely to give an m that is already
 too large to be a correct limiting value of
 dK/dp as $p \rightarrow \infty$.

When $K_0' < 0$, as for vitreous silica, many
 formulas including those of Murnaghan and
 Keane necessarily predict an instability ($K \leq$
 0) on $P > 0$. Although this may not be a great
 catastrophe, and could even be represented as
 advantageous (because an actual material with
 $K_0' < 0$ could be presumed to undergo a phase
 transition, through which the extrapolation
 should not be continued analytically), it is in-
 teresting to note that the present formula allows
 such an instability to be avoided by choosing
 a sufficiently high positive value for C . This
 is illustrated in Figure 1, which shows K/K_0
 versus P for three different values of C with
 $K_0' = -6.5$ and $m = 1$. The value $K_0' = -6.5$
 applies to vitreous silica [McSkimin as cited by
 Anderson, 1961].

COMPRESSION EQUATION

The next task is to relate the volume v to the
 pressure, subject to equation 2 and the defini-
 tion of the bulk modulus

$$K = -v dp/dv \quad (5)$$

$$V = \left\{ \left[\frac{a}{mP^2 + (1 + A + am)P + a} \right] \right.$$

$$\left. \cdot \left[\frac{4am + 2mP[(q)^{1/2} + (1 + A + am)]}{4am - 2mP[(q)^{1/2} - (1 + A + am)]} \right]^{(1+A-am)/(q)^{1/2}} \right\}^{1/2m} \quad (9)$$

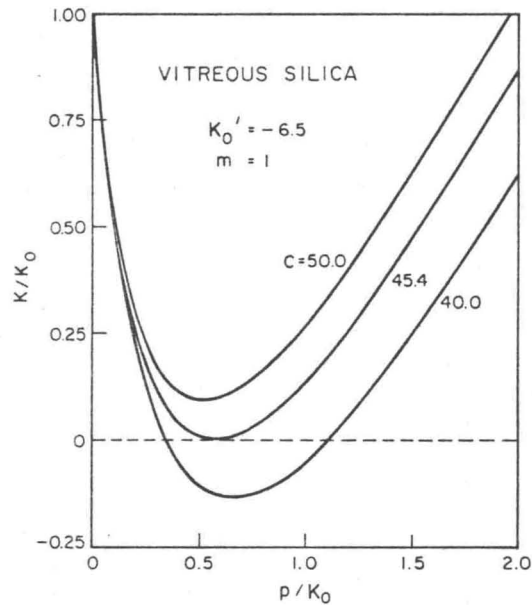


Fig. 1. Determining the value of C that ensures reasonable behavior of K on $p > 0$ for the anomalous case, vitreous silica ($K_0' < 0$) (see text).

Let $V = v/v_0$. Then

$$K/K_0 = -V dP/dV \quad (6)$$

As an abbreviation in equation 2, let $A = a(K_0' - m)$. Then the integral of equation 2 is

$$\frac{K}{K_0} = -V \frac{dP}{dV} = 1 + A + mP - \frac{aA}{P + a} \quad (7)$$

where the constant of integration has been determined to make $K = K_0$ at $P = 0$. From (7)

$$V = \exp \left[- \int \frac{dP}{\left(1 + A + mP - \frac{aA}{P + a} \right)} \right] \quad (8)$$

The evaluation of the integral in the expres-
 sion above, subject to $V = 1$ when $P = 0$
 (Given in Appendix B), gives us