

Fig. 15 Pressure-volume relations in diamond cell

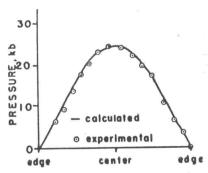


Fig. 16 Calculated pressure gradient -T1Br, 12 kbar

$$V = f (P,T,h,r) = \pi r_0^2 h$$

At constant temperature and pressure

$$(dV/dP)_{P,T,r} = \pi r_0^2 (dh/dP)_r$$

However, this is the same volume change which would occur at a given applied pressure in an identical system of constant h and variable r, Fig.26(b),

$$(dV/dP)_{P,T,h} = 2\pi rh_{o} (dr/dP)$$

This volume change can be related to β and give a relations for dP/dr, the pressure gradient. Combining the definition of β , equation (1) and the foregoing expression, we get

$$- (1/V_0) 2\pi rh_0 (dr/dP) = \beta$$

Multiplying both sides by $V_0/\Delta V_a$, where $\Delta V_a = \pi h_0 (r_0^2 - r_a^2)$, we get

=
$$(2r/r_0^2 - r_a^2)$$
 dr = (βdP) $(V_0/\Delta V_a)$

Integrating between r_0 and r corresponding to P=0 kbar at the edge to P=P at r, we get

$$-\int_{r_{0}}^{r} (2r/r_{0}^{2} - r_{a}^{2}) dr = V_{0}/\Delta V_{a} \int_{0}^{P} \beta dP$$

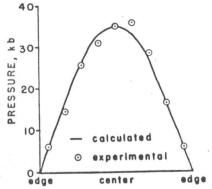


Fig. 17 Calculated pressure gradient - NaC1. 20 kbar

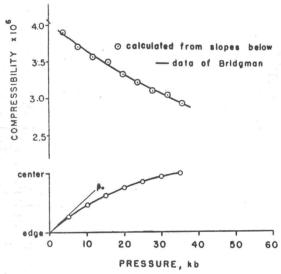


Fig. 18 Compressibility calculation - NaC1

$$\frac{1 - r^2}{1 - r_a^2} = \frac{(aP - bP^2)V_o}{V_a} = \frac{aP - bP^2}{aP_a - bP_a^2}$$
 (2)

Similarly, the pressure, P, and r may be compared to P_m , the maximum pressure at the center of the cell by means of a similar derivation to get relation (3).

$$\frac{1-r^2}{1-r^2_m} = \frac{(aP-bP^2)V_0}{\Delta V_m} = \frac{aP-bP^2}{aP_m-bP_m^2}$$
(3)

Now it is necessary to introduce another relation to evaluate the constant ra; viz.

$$\vec{P} = \sum_{i} A_{i} P_{i} / \sum_{i} A_{i}$$