

Chapter 10

These are answers to the exercises in Chapter 10 of:

Understanding the Properties of Matter
by Michael de Podesta.

If you find an error in these answers, or think they could be clarified in any way, please feel free to contact me.

Thanks

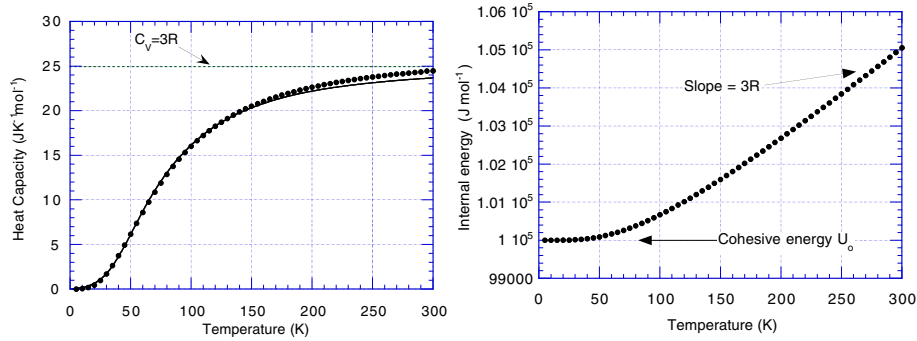
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P1. In general, $U(T) = \int_0^T C(T) dT$ so $U(T) = \frac{\gamma T^2}{2} + \frac{\alpha T^4}{4}$

At 1 K $U(T) \approx 5.25 \times 10^{-5} \text{ J mol}^{-1}$ and at 20 K $U(T) \approx 0.42 \text{ J mol}^{-1}$ and

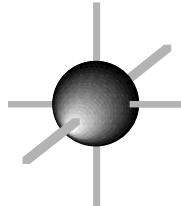
These are both very small compared to $U_0 \approx 10^5 \text{ J mol}^{-1}$

P2.

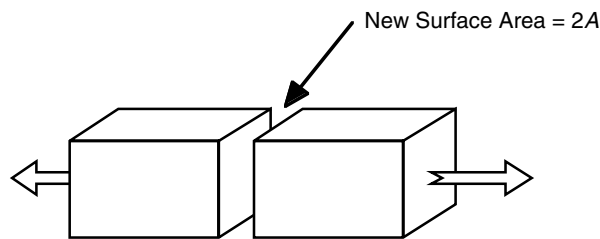


P3. Each atom has 6 bonds, each of which it ‘half owns’ i.e. associated with each atom are $6 \times 0.5 = 3$ bonds. So in 1 mole of substance there are $3N_A$ bonds so the energy per bond is $U_0 / 3N_A$ joules

$\approx 0.346 \text{ eV}$



The number of bonds broken when a bar of cross sectional area A is broken is $\approx A/a^2$ and each bond represents the loss of 0.346 eV of energy. The total area of surface formed is $2A$ (area A on each broken part) and so the energy per unit area of surface formed is $0.346 \times (A/a^2)/2A$ in eV m^{-2} evaluates to 0.31 Jm^{-2} . In practice the surface structure changes to minimise this energy.



P4.

0.7 K below T_M the difference in free energies is $0.7 \times (400-395) \approx 0.35 \text{ JK}^{-1}\text{mol}^{-1}$

1 mole has a volume...

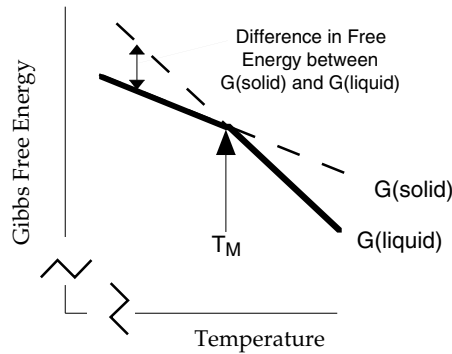
$$N_A a^3 = 1.626 \times 10^{-5} \text{ m}^3$$

So ΔG_{LS} per cubic metre is

$$\Delta G_{LS} \approx 0.215 \times 10^6 \text{ Jm}^{-3}$$

Critical embryo size is $a >$

$$\frac{2 \times 0.6}{0.215 \times 10^6} \approx 5.6 \text{ } \mu\text{m}$$



This seems unfeasible large: it involves VAST numbers of atoms all fluctuating in the same way. In practice the surface energy *difference* between the liquid and solid states is much less than the surface energy of the solid state, perhaps only \approx few % of the 0.6 (or 0.3) Jm^{-2} we estimated here. This reduces the embryo size estimate to \approx few nm, which is much more plausible.

P5. Following Example 10.7,

$$\left. \frac{dP}{dT} \right|_{\text{Water}} = -15.6 \times 10^6 \text{ Pa K}^{-1} \quad \left. \frac{dP}{dT} \right|_{\text{ethanol}} = +5.53 \times 10^6 \text{ Pa K}^{-1}$$

$$\text{So } T_M(\text{water}) = 273.16 - P/15.6 \times 10^6 \text{ and } T_M(\text{ethanol}) = 155.85 + P/5.53 \times 10^6$$

Equating these and solving for P yields $P = 0.478 \text{ GPa}$ or $\approx 4.8 \text{ kBar}$. This is quite easily achievable, but the ice would probably have formed a new structure at around 2 kBar and the slope of the melting curve changed to a positive value (See Chapter 12 Q28). The prediction is that the melting temperatures would be the same at $T_M \approx 242.3 \text{ K} \approx -30.9 \text{ }^\circ\text{C}$. In fact the minimum melting temperature for water is around $-20 \text{ }^\circ\text{C}$.

P6. I don't know for sure whether the slipperiness is connected with the temperature of the sliding object, but I would expect it be, and I find the premelting hypothesis very convincing.

P7.

	T	P	
(a)	200	10^5	Solid
(b)	200	10^8	Liquid according to diagram, but actuality solid (See Chapter 12 Q28)
(c)	400	10^8	Liquid
(d)	400	10^5	Gas
(e)	700	10^5	Gas
(f)	700	10^8	Liquid/Gas near critical point where the distinction is blurred.
(g)	273.15	611	Could be solid, liquid or gas: This is the triple point