

EXAMINATION SOLUTION FORM, Semester 1, 2014-15

Examiner
(initials)

JMA

Module Code

PHY 2063.

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Number

1.

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of

<ul style="list-style-type: none"> • Use BLACK ink only • Legible hand-written model answers are preferred • Indicate whether "bookwork" or not 	MARK S
<p>a) i) Heat energy required to raise temperature of (a specified e.g. 1mol or 1kg) of substance by 1K, under specified conditions (e.g. fixed volume)</p> $C_v = \left. \frac{dQ}{dT} \right _v = T \left. \frac{dS}{dT} \right _v$ <p>\therefore Integrating gives $\Delta S = \int_{T_1}^{T_2} \frac{C_v}{T} dT$</p>	<p>1</p> <p>1</p> <p>1</p> <p>[BW]</p>
<p>ii) Thermodynamic potentials are used to determine equilibrium in open systems (systems in contact with an external reservoir).</p> <p>For example Helmholtz free energy $F = U - TS$ is minimized for systems in contact with a heat bath at fixed temperature T.</p> <p>Equilibrium occurs when: $dF = 0$ (F is a minimum).</p>	<p>1</p> <p>1</p> <p>1</p> <p>[BW]</p>
<p>b) i) $W = \frac{N!}{(N-n)! n!}$</p> <p>N positions for a domain wall. n domain walls</p> <p>ii) $F = \epsilon n - T S$</p> $S = k_B \ln W \approx k_B (N \ln N - n \ln n - (N-n) \ln (N-n))$ <p>using Stirling's approx. $\ln N! \approx N \ln N - N$</p> <p>Combining these gives stated result</p>	<p>2.</p> <p>1</p> <p>2</p> <p>1.</p>

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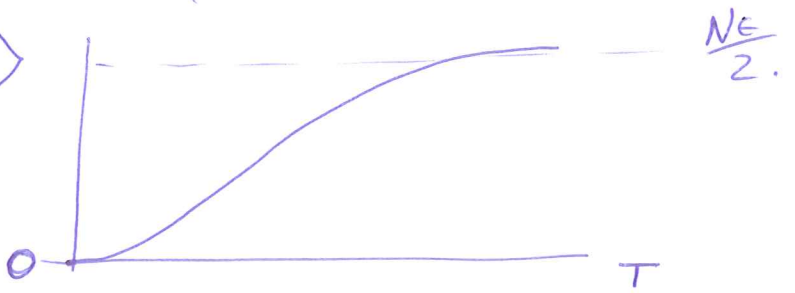

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<p>iii) $\frac{dF}{dn} = 0$ in equilibrium</p> $\Rightarrow 0 = \epsilon - k_B T (-\ln n + \ln(N-n))$ $\Rightarrow \frac{\epsilon}{k_B T} = \ln \left(\frac{N-n}{n} \right)$ $\Rightarrow \frac{N}{n} - 1 = e^{\epsilon/k_B T} \therefore n = \frac{N}{1 + e^{\epsilon/k_B T}}$ $\approx N e^{-\epsilon/k_B T}$ $\therefore \langle E \rangle \approx N \epsilon (e^{\epsilon/k_B T} + 1) \approx N \epsilon e^{-\epsilon/k_B T}$ <p>iv) $\langle E \rangle$</p>  <p style="text-align: right;">$\frac{N\epsilon}{2}$</p>  <p style="text-align: center;">$T \sim \frac{\epsilon}{k_B}$</p> <p>High T — equally divided between wall/no wall states $\Rightarrow \langle E \rangle = \frac{N\epsilon}{2}$.</p> <p>Low T — very few walls & no thermal energy to create more $\therefore C_v$ small.</p>	<p>1</p> <p>2</p> <p>1</p> <p>1</p> <p>2</p> <p>1</p>

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a) i) Each independent quadratic degree of freedom has thermal energy $\langle E \rangle = \frac{1}{2} k_B T$ provided the energy level spacing $\hbar \omega \ll k_B T$. (i.e. T is high compared to $\frac{\hbar \omega}{k_B}$).

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[BW]

ii) A diatomic gas can store energy as follows:

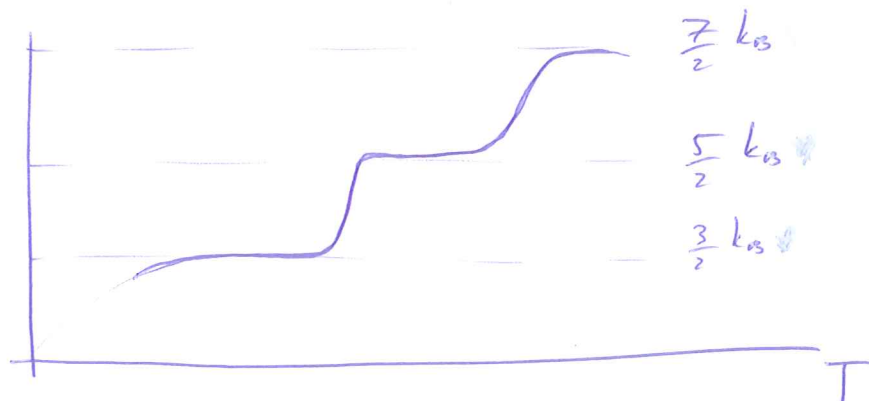
translational kinetic energy 3 components (x, y, z)
 $\Rightarrow \frac{3}{2} k_B T$

rotational kinetic energy 2 components
 $\Rightarrow k_B T$

vibrational kinetic & potential energy $\Rightarrow k_B T$

These reach their equipartition values provided temperature is high enough. Typically vibrational modes require the highest T , then rotational, then translation for equipartition to apply.

C_v
per
molecule



2

[BW]

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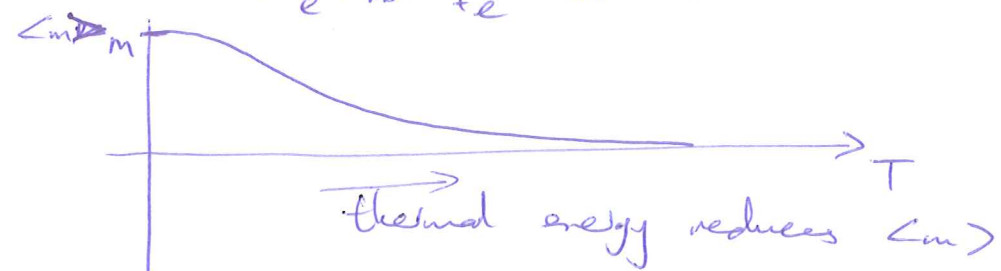
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<p>b) i) $Z = \sum_{states, i} e^{-\epsilon_i/k_B T}$</p> <p>here $\epsilon = +mB$ & $-mB$</p> <p>$\therefore Z = e^{mB/k_B T} + e^{-mB/k_B T}$</p> <p>ii) $p_p = \frac{e^{mB/k_B T}}{e^{mB/k_B T} + e^{-mB/k_B T}} = \frac{1}{1 + e^{-2mB/k_B T}}$</p> <p>$p_{ap} = \frac{1}{e^{2mB/k_B T} + 1}$ [or equivalent]</p>	<p>1</p> <p>1</p> <p>1</p> <p>1</p> <p>1</p>
<p>iii) $\langle m \rangle = +m p_p + -m p_{ap}$</p> <p>$= m \left(\frac{e^{mB/k_B T} - e^{-mB/k_B T}}{e^{mB/k_B T} + e^{-mB/k_B T}} \right) = m \tanh \frac{mB}{k_B T}$</p> 	<p>2</p> <p>1</p> <p>2</p>
<p>iv) $B \rightarrow \frac{B}{2}$ effective level spacing halved.</p> <p>$\Rightarrow p_{ap} = \frac{1}{e^{\frac{2mB}{k_B (2T)}} + 1} = \frac{1}{e^{\frac{2mB}{2k_B T}} + 1}$</p> <p>$\therefore T_{old} = 2T$ i.e. temperature half of old value.</p>	<p>2</p>

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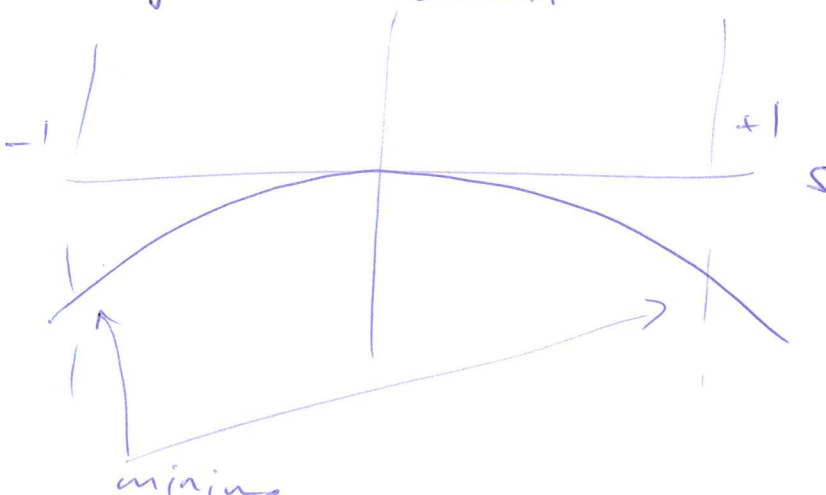
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<p>a) i) Order parameter — A quantity that is zero in the disordered (high T) phase & non-zero in the ordered (low T) phase. that describes the phase transition</p> <p>ii) Paramagnet \rightarrow ferromagnet — magnetisation liquid \rightarrow vapor — density difference :</p>	<p>2. [BW]</p> <p>2</p> <p>2</p> <p>[BW]</p>
<p>b) i) First term $\frac{zJ}{2} (1-s)(1+s)$</p> <p>Interaction energy of average spins with neighbours. $s = \pm 1$ (all aligned) favored</p> <p>term A</p> 	<p>1 [BW]</p> <p>2</p>

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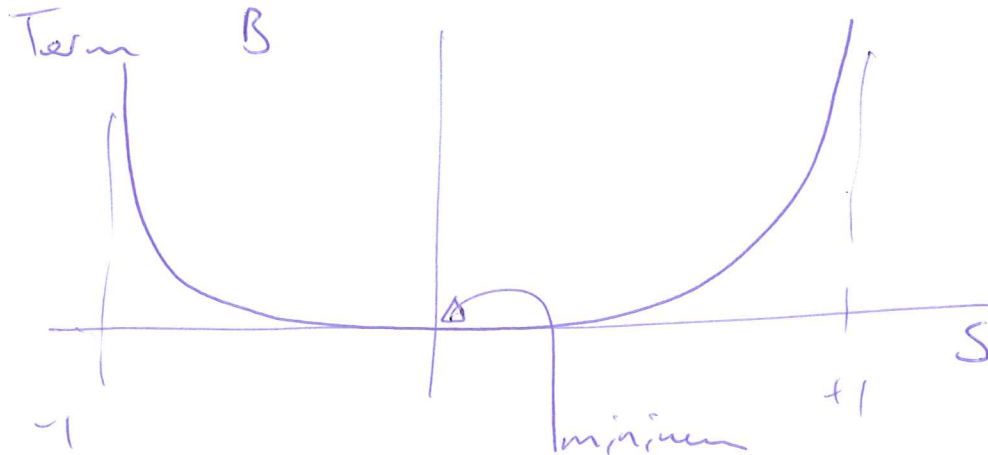
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\Rightarrow favors $s \approx 0$. (entropic term)

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[RW]

ii) At equilibrium $\frac{\partial F}{\partial s} = 0$

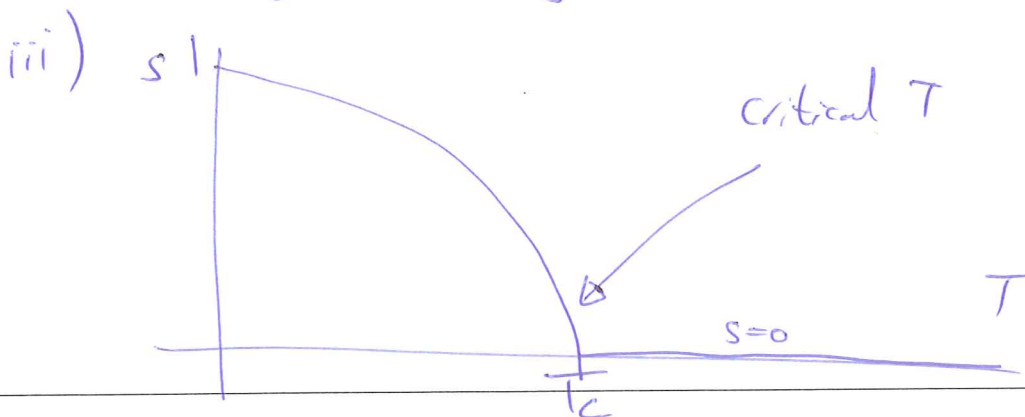
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$$\Rightarrow 0 = \frac{zJ}{z} (-2s) - \frac{k_B T}{z} (-\ln(1+s) + \ln(1-s))$$

2

$$\Rightarrow \frac{1}{k_B T} = \frac{1}{2zJ} \ln \frac{1+s}{1-s}$$

2



2.