

The final exam covers the entire course but emphasizes the material since the second midterm: over half the questions will come from chaps. 6 and 7. You are allowed a 1-page cheat sheet.

Review homework problems

Boltzmann factor, partition func'n Z , probabilities $\mathcal{P}(s) = Z^{-1} \exp(-\beta E(s))$; $Z = \sum_s \exp(-\beta E(s))$; $\beta \equiv 1/k_B T$

Z for specific cases: paramagnet: $Z = 2 \cosh \beta \mu B$; oscillator: $Z = (1 - e^{-\beta \epsilon})^{-1}$; ideal gas $Z_{tr} = (L/\lambda_Q)^3$

$$U = N\bar{E} = -N \partial \ln Z / \partial \beta \quad C_V = (\partial U / \partial T)_V \quad S(T) = \int_0^T \frac{C_V(T')}{T'} dT'$$

Application to paramagnetism, diatomic-molecule rotation, proof of equipartition theorem

Maxwell speed and velocity distributions in 3D, also 1D, 2D. Mean, mode, rms

$$F \equiv U - TS = -k_B T \ln Z; \quad S = -(\partial F / \partial T)_N \quad p = -(\partial U / \partial V)_{S,N}; \quad B = -V(\partial p / \partial V)_{T,N}$$

$$\mu = -T(\partial S / \partial N)_{U,V} = (\partial U / \partial N)_{S,V} = (\partial F / \partial N)_{T,V} = G/N|_{T,p}$$

Partition function for composite systems: distinguishable vs. indistinguishable

Ideal gas $Z_{tr} = (L/\lambda_Q)^3$, $Z = (V Z_{int}/v_Q)^N / N!$, counting modes in box, quantum length $\lambda_Q = h/\sqrt{(2\pi m k_B T)}$

Gibbs factor, grand partition function \mathcal{Z} , probabilities $\mathcal{P}(s) = \mathcal{Z}^{-1} \exp[-\{E(s) - \mu N(s)\}/k_B T]$

Langmuir adsorption isotherm

Density of states $\mathcal{G}(\epsilon) \propto \epsilon^{(d-2)/2}$ for $\epsilon \propto n^2$ and how it relates to N , U , etc.; for $\epsilon \propto n$, $\mathcal{G}(\epsilon) \propto \epsilon^{(d-1)}$

$$\epsilon_F = (\hbar^2/2m)(3\pi^2 N/V)^{2/3} \text{ (for } d=3) \quad \mathcal{G}(\epsilon) \text{ has units } 1/\text{energy}$$

Fermions: Pauli exclusion principle, partition function, Fermi-Dirac dist. $\bar{n}_{FD}(\epsilon) = 1/[e^{\beta(\epsilon - \mu)} + 1]$

Degenerate Fermi gas at $T=0$: definitions T_F , ϵ_F ; quantities U , p , B

Sommerfeld expansion, meaning, usage to get lowest order (in T) contributions to shift of chemical potential μ from ϵ_F , increase in U , $C_V \propto T$

$$\int_{-\infty}^{\infty} H(\epsilon) \bar{n}_{FD}(\epsilon) d\epsilon = \int_{-\infty}^{\mu} H(\epsilon) d\epsilon + \frac{\pi^2}{6} (k_B T)^2 H'(\mu) + O(k_B T)^4$$

Bosons: Planck distribution for simple harmonic oscillator, photons, mode counting (again)

Planck spectrum ($\mu = 0$) plotted vs ϵ or vs. λ ; total power radiated (eq. 7.99) (emissivity) (area) σT^4

Wien displacement law; $\lambda_{\max} T = \text{const.}$

Debye theory of lattice vibrations of solids: deficiency of Einstein model at low T , phonons and their [acoustic] dependence on n ; idea of n_{\max} ; $C_V \propto T^3$

Bose-Einstein condensation: N_0 and N_0/N ; role of chemical potential and of convergence of integral of

$$\mathcal{G}(\epsilon) \bar{n}_{BE}(\epsilon) \quad \bar{n}_{BE}(\epsilon) = 1/[e^{\beta(\epsilon - \mu)} - 1]$$

$N_{\text{exc}} = (T/T_c)^{3/2} N$ and $N_0 = N - N_{\text{exc}}$ (for $T < T_c$); consequent U , C_V , p

Ising model: energy = $J \sum_{\langle i,j \rangle} s_i s_j$ where $s_i = \pm 1$ (up or down) Mean field $k_B T_C = J \times (\# \text{ nearest neighbors})$

Important Figs.: 6.11, 6.12, 6.16, 7.6, 7.7, 7.9, 7.14, 7.16, 7.19, 7.26, 7.29, 7.31, 7.32, 7.33, 7.34, 7.37; and what they mean/how to use them