



PHY 410 - Reference Sheet

Stirling’s approximation - for very large N the factorial can be very accurately approximated with the following

log N! ≈ N log N − N
N! ≈ √(2πN) N^N e^(-N)

Fractional uncertainty of \mathbb{X} is uncertainty of expected value per particle.

(X)/N = (sqrt(X^2) - (X)^2)/N

Boltzmann’s constant
k_B = 1.380649 × 10^−23 m^2 s^−2 K^−1
Entropy S = k_B σ, σ_TOT = σ_1 + σ_2
Temperature T = τ/k_B

Microcanonical Ensemble

Multiplicity function

g = # of microstates, P(n) = 1/g

Expected value of \mathbb{X} is the average across all microstates.

(X) = sum(n) X(n)P(n) = 1/g sum(n) X(n)

Entropy can be written in terms of the multiplicity function.

σ(N, T, U, V, P) ≡ log[g(N, T, U, V, P)]

Binary System

A **binary system** is a system of N particles where each particles has two possible states. Let N_{\uparrow} is the number of particle in the up state and N_{\downarrow} be the number of particles in the down state.

g(N, N_{\uparrow}) = N! / (N_{\uparrow}!(N - N_{\uparrow})!), sum_{N_{\uparrow}=0}^N g(N, N_{\uparrow}) = 2^N

The binary system can be rewritten in terms of the difference between up states and down states this is the **spin excess**.

2S = N_{\uparrow} - N_{\downarrow}
g(N, S) = N! / ((N/2 + S)!(N/2 - S)!)
sum_{S=-N/2}^{S=N/2} g(N, N_{\uparrow}) = 2^N

Applying Stirling’s approximation to the binary model, for large N the multiplicity function and fractional uncertainty are

g(N, S) ≈ g(N, 0)e^(-2s^2/N)
g(N, S) ≈ sqrt(2/(πN)) 2^N e^(-2s^2/N)
(ΔS)/N ≈ 1/sqrt(N)

An example of a binary system is N spin 1/2 particles in an external **magnetic field** B . The total energy U and magnetization M of the system are

U = sum_{i=1}^N -m_i · B = -(N_{\uparrow} - N_{\downarrow})mB = -2SmB
M = 2Sm = -U/B
g(N, U) = N! / ((N/2 - U/(2mB))!(N/2 + U/(2mB))!)
σ(N, S) ≈ -((N/2 + S) log(N/2 + S/N) - (N/2 - S) log(N/2 - S/N))
M = Nm tanh(mB/τ)

Einstein Solid

An **einstein solid** is a system of N atoms where each atom is modeled as a harmonic oscillator the energy of the system is determined by the number of atoms n oscillating at frequency ω .

U = nħω
g(N, n) = (n + N - 1)! / n!(N - 1)!
g(N, n) ≈ ((n+N)/n)^n ((n+N)/N)^N / sqrt(2πn(n+N)/N)

Thermal Equilibrium

Temperature

1/τ = (∂σ/∂U)_{N,V}

Thermal Equilibrium

((∂σ_1/∂U_1)_{N_1,V_1} = (∂σ_2/∂U_2)_{N_2,V_2})
1/τ_1 = 1/τ_2

2nd law of thermo - Change in entropy ≥ 0.
Sharpness of Equilibrium For a two binary systems, the number of states in a configuration of deviation δ from equilibrium is

g_1 g_2 = (g_1 g_2)_{max} e^(-2δ^2/N_1 - 2δ^2/N_2)

Canonical Ensemble

Partition Function - partition by energy levels for a fixed temperature

z = sum_n e^(-ε_n/τ), P(n) = 1/z e^(-ε_n/τ)
z = sum_α g(ε_α) e^(-ε_α/τ), for degeneracy g(ε_α)

Expected Value of \mathbb{X} is the average across all energies (Thermal Average).

(X) = sum_n X(n)P(n) = 1/z sum_n X(n)e^(-ε_n/τ)

Expected Energy in the canonical ensemble is

U = (ε) = 1/z sum_n ε_n e^(-ε_n/τ)
U = (ε) = τ^2 1/z ∂z/∂τ = τ^2 ∂/∂τ log z

The total partition function and expected energy for N non-interacting particles is simply

z_N = z_1^N
U_N = (ε)_N = NU_1 = N(ε)_1
(this also applies for expected value of any \mathbb{X})

Theromodynamic Relations

1st Law of Thermo

dU = dQ + dW = τdσ - PdV
dσ = 1/τ dU + P/τ dV

Temperature τ = (∂U/∂σ)_V
Quasi-static Compression Equilibrium
the equilibrium condition for quasi-static compression is

((∂U_1/∂V_1)_{σ_1} = (∂U_2/∂V_2)_{σ_2})

Helmholtz Free Energy

F = U - τσ = U - ST = -τ log z
dF = -σdτ - PdV

Entropy σ = - (∂F/∂τ)_V, S = k_B σ
Pressure

P = - (∂U/∂V)_σ = τ (∂σ/∂V)_U = - (∂F/∂V)_τ

Energy

U = -τ^2 ∂/∂τ (F/τ)

Ideal Gas

DeBroglie Thermal Wavelength is the wavelength of the wave functions of matter at a given temperature.

λ_T = sqrt(2πħ^2/mτ)

Concentration of a system is the inverse of the volume

n = 1/V

Quantum Concentration is the density of quantum state per particle. It is used to define when a system will behave classically ($n < n_Q$) and when a system will be dominated by quantum effects ($n > n_Q$).

n_Q = 1/λ_T^3

Single Particle Ideal Gas is a system in the canonical ensemble consisting of a signle particle in a box of side lengths L . The energy levels , partition function and average energy are

E_{n_x, n_y, n_z} = h^2/(2m) * (π/L)^2 * (n_x^2 + n_y^2 + n_z^2)
z_1 = V/λ_T^3
U_1 = 3/2 τ
σ_1 = log(V/λ_T^3) + 3/2

Gibbs Resolution states that for systems in the classical regime the partition function for an ideal gas with N particles is

z_N = 1/N! (z_1)^N
U_N = 3/2 Nτ
σ_N = N [log(V/(Nλ_T^3)) + 5/2]

N-Particle Ideal Gas - by applying Gibbs resolution and properties of expected values we can find the classical ideal gas results

PV = Nτ
U = 3/2 Nτ
σ = N [log(V/(Nλ_T^3)) + 5/2]

DOG (bork)

