

# Statistical Thermodynamics

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” Put a cool quote here or your lame ”

-Thomas Brosnan

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# 1 Concepts and Terminology

## 1.1 System macrostate

- Properties at large scale of the system when we know constraining thermodynamic parameters such as  $P, V$  and  $T$ .

## 1.2 Quantum microstate

- Each quantum state is a separate and distinct microstate of the system.

## 1.3 Weakly coupled systems

- We assume weakly coupled systems. These are isolated systems such that the total macro-parameters  $E, V, N$ , are constant. Weak coupling implies that the energy levels of a single particle are effectively unchanged by particle interactions, but the interaction is sufficiently large to allow energy exchange. This means that at equilibrium the system must have a common temperature.

This assumption makes the quantum mechanics involved easier, as coupled systems in quantum mechanics can be quite complex where as simple systems like a gas of  $H_2$  can be solved analytically (eigenvalues of vibration, rotation and translation).

## 1.4 Locality

- Localised particles are particles that are restricted to a certain place like a lattice for example. This way each particle of the lattice is *distinguishable*, i.e. it can be told apart from the other particles in the gas due to its position. If particles in a system are non-localised they are *indistinguishable*.

## 1.5 Bosons and Fermions

- In the case of indistinguishable particles we have two cases. Consider a two state system with two single particle states (orbitals)  $a$  and  $b$  and a wave function  $\Psi$  describing the system. We assume (weakly coupled systems) that the particles are non-interacting so that we can write the wavefunction as a product of the two states:

$$\Psi(1, 2) = \psi_a(1)\psi_b(2) \quad \text{or} \quad \Psi(2, 1) = \psi_a(1)\psi_b(2) \quad (1.1)$$

Where here there are two possibilities for the states as the particles are indistinguishable. Quantum mechanics tells us that the wavefunction must be a linear combination of all the (equally likely thus same amplitude) possible states, so we write:

$$\Psi(1, 2) \propto \psi_a(1)\psi_b(2) \pm \psi_a(2)\psi_b(1) \quad (1.2)$$

The  $\pm$  here indicates the two possibilities with respect to particle exchange. If we swap two particles and the system is symmetric (i.e. we get a  $+$ ) then these particles are *Bosons*. And we can have any number of them in a single state, as if the states  $a = b$  1.2 is not 0.

However if we swap two particles and the system is anti-symmetric (now we get a  $-$  sign) then the particles are *Fermions* and we can only have one fermion per state, as 1.2 is 0 if  $a = b$ . This is the Pauli exclusion principle.

