

LECTURE 6: Degenerate bosons

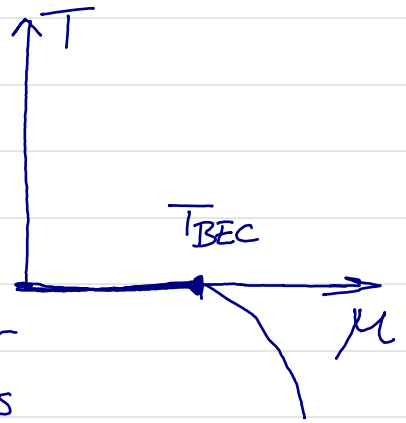
1° Understanding chemical potential of bosons

$$n(\epsilon) = \frac{1}{e^{\beta(\epsilon-\mu)} - 1} \Rightarrow \mu \leq \epsilon_0 = 0 \text{ for } \epsilon_k = \frac{\hbar^2 k^2}{2m}$$

in order to keep $n(\epsilon) > 0$

$$N = \sum_{\mathbf{k}} \frac{1}{e^{\beta(\epsilon_{\mathbf{k}}-\mu)} - 1} \Leftrightarrow N = \int_0^{\infty} \frac{D(\epsilon) d\epsilon}{e^{\beta(\epsilon-\mu)} - 1}$$

→ as T decreases, μ MUST decrease in absolute value as well to keep N constant ⇒ when μ becomes zero, further decrease of T leads to loss of particles



→ integral does not change if you change its value at discrete set of points of measure zero; also $D(\epsilon) \propto \sqrt{\epsilon}$ means $D(\epsilon) = 0$ so any number of particles in $\epsilon = 0$ is not included:

$$N = N_0 + N_* = \frac{g}{e^{\beta\mu} - 1} + g \frac{V}{4\pi^2} \left(\frac{2m}{\hbar^2}\right)^{3/2} \int_0^{\infty} \frac{\sqrt{\epsilon} d\epsilon}{e^{\beta(\epsilon-\mu)} - 1}$$

grand canonical ensemble applies

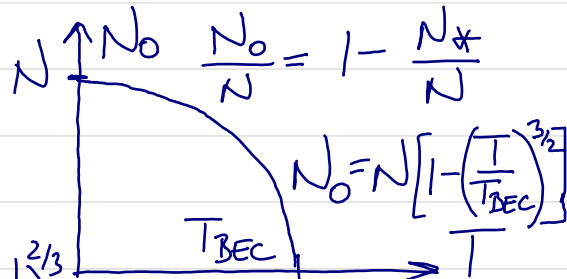
macroscopic number of particles in Bose-Einstein condensate

only to particles above the condensate

$$N_* = N \Rightarrow T = T_{BEC}$$

$$\frac{N}{V} = g \frac{(2mk_B T_{BEC})^{3/2}}{4\pi^2 \hbar^3} \int_0^{\infty} \frac{x^{1/2} dx}{e^x - 1}$$

$$\int_0^{\infty} \frac{x^{m-1} dx}{e^x - 1} = \Gamma(m) \zeta(m) \Rightarrow T_{BEC} = \frac{3.31 \hbar^2}{m k_B} \left(\frac{N}{gV}\right)^{2/3} \left[\zeta(3/2) = 2.6124 \right]$$



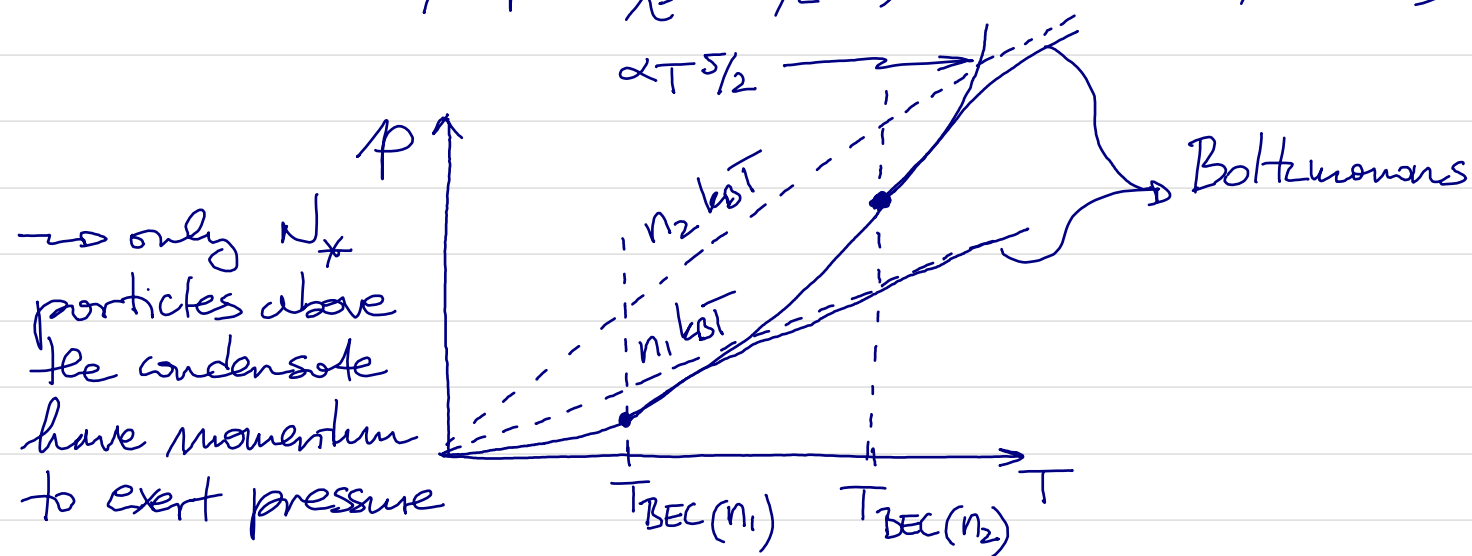
→ role of thermodynamic limit: $N_0 = \frac{1}{e^{\beta\mu} - 1} \approx \frac{-k_B T}{\mu}$

$\left. \begin{array}{l} \mu \\ T_{BEC} \\ T \end{array} \right\} \text{for finite } N \leftarrow \mu \sim \frac{1}{N} \text{ as } \begin{array}{l} N \rightarrow \infty \\ V \rightarrow \infty \\ \frac{N}{V} = \text{const.} \end{array}$

2° pressure of BEC

$$T < T_{BEC} \Rightarrow \beta p = \frac{g}{\lambda^3} \zeta_{5/2}^+(1) \approx 1.341 \frac{g}{\lambda^3} \neq F(n)$$

$z = e^{\beta\mu} = 1$ in the condensate



3° Is BEC of noninteracting particles first or second order phase transition?

$$n_* = \frac{g}{\lambda^3} \zeta_{3/2}^+(1) \Rightarrow V^* = \frac{1}{n_*} = \frac{\lambda^3}{g \zeta(3/2)}$$

so transition can be induced by reducing volume $V < V_*$ or increasing density at fixed T

→ Clausius - Clapeyron relation for latent heat Q_L :

$$\mu_0(T, p) = \mu_*(T, p) \Leftrightarrow d\mu_0 = d\mu_* \Rightarrow d\mu_0 = -S_0 dT + V_0 dp$$

$$\left(\frac{\partial \mu}{\partial T} \right)_p = -S, \quad \left(\frac{\partial \mu}{\partial p} \right)_T = V \Rightarrow \frac{\partial p}{\partial T} \Big|_{\text{coexist}} = \frac{S_* - S_0}{V_* - V_0} = \frac{Q_L}{T_{BEC}(V_* - V_0)}$$

$$dp/dT = \frac{5}{2} p/T = \frac{\Delta S}{\Delta V} = \frac{Q_L}{T_{BEC}(V_* - V_0)} \rightarrow V_0 \equiv 0$$

unrealistic due to no interparticle interactions

$$Q_L = T_{BEC} V_* \frac{dp}{dT} = T_{BEC} V_* \frac{5}{2} \frac{p}{T_{BEC}} =$$

$$= \frac{5}{2} \frac{g}{\lambda^3} \left(\frac{5}{2} \right) k_B T_{BEC} \left[\frac{g}{\lambda^3} \right]^{-1} = \frac{5}{2} \frac{\zeta(5/2)}{\zeta(3/2)} k_B T_{BEC} \approx 1.28 k_B T_{BEC}$$

$\Rightarrow Q_L \neq 0 \Rightarrow$ first order phase transition

■ compressibility $\alpha_T = \left(\frac{\partial n}{\partial p} \right)_T / n$

$$\left. \begin{aligned} dp/dz &= \frac{g k_B T}{\lambda^3} \frac{1}{z} f_{3/2}^+(z) \\ dn/dz &= \frac{g}{\lambda^3} \frac{1}{z} f_{1/2}^+(z) \end{aligned} \right\} \text{using } \frac{d}{dz} f_m^+(z) = \frac{1}{z} f_{m-1}^+(z)$$

take ratio $\Rightarrow \alpha_T = f_{1/2}^+(z) [n k_B T f_{3/2}^+(z)]^{-1} \rightarrow \infty$ as $T \rightarrow T_{BEC}$

$\lim_{z \rightarrow 1} f_{1/2}^+(z) \rightarrow \infty$

\rightarrow since α_T diverges as $T \rightarrow T_{BEC}$ this would be continuous or second order phase transition

■ heat capacity $C_V = \left(\frac{\partial E}{\partial T} \right)_{V,N}$

$$E = \frac{3}{2} p \cdot V = \frac{3}{2} V \frac{g}{\lambda^3} k_B T f_{5/2}^+(z) \propto T^{5/2} f_{5/2}^+(z)$$

$$C_V = \frac{3}{2} V \frac{g}{\lambda^3} k_B T \left[\frac{5}{2T} f_{5/2}^+(z) + \frac{1}{z} f_{3/2}^+(z) \frac{\partial z}{\partial T} \Big|_{V,N} \right]$$

N is fixed $\Rightarrow dN/dT|_V = \frac{g}{\lambda^3} V \left[\frac{3}{2T} f_{3/2}^+(z) + \frac{1}{z} f_{1/2}^+(z) \frac{dz}{dT} \Big|_{V,N} \right]$

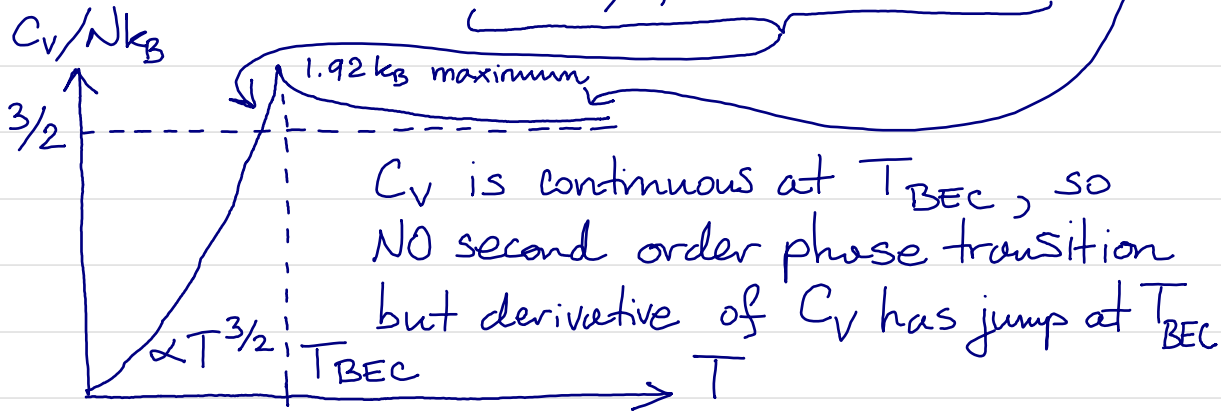
$$\frac{T}{z} \frac{dz}{dT} \Big|_{V,N} = - \frac{3}{2} \frac{f_{3/2}^+(z)}{f_{1/2}^+(z)}$$

$$\frac{C_V}{V k_B} = \frac{3}{2} \frac{g}{\lambda^3} \left[\frac{5}{2} f_{5/2}^+(z) - \frac{3}{2} \frac{f_{3/2}^+(z)}{f_{1/2}^+(z)} \right]$$

at high T $\approx \frac{3}{2} \left[1 + \underbrace{n \lambda^3 / 2^{7/2}}_{\text{larger than the classical value}} + \dots \right]$

→ at low T $z = e^{\beta\mu} = 1$:

$$\frac{C_V}{N k_B} = \frac{15}{4} \frac{g}{n \lambda^3} \zeta_{5/2} = \frac{15}{4} \zeta_{3/2} \left(\frac{T}{T_{BEC}} \right)^{3/2}$$



- INTUITIVE interpretation of $C_V \propto T^{3/2}$:
- at $T=0$ K all particles are in state with $\vec{k}=0$
 - at T slightly above 0 K, particles occupy states with $\leq k_{max}$ where $\frac{1}{2} k_{max}^2 / m = k_B T$ and each of these states has energy $\propto k_B T$
 - $E \propto V k_{max}^d k_B T$ for $V k_{max}^d$ as the density of states in d -dimensions $\Rightarrow C_V \propto V k_B T^{d/2}$