

Statistical Mechanics
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Lecture - 56
Pressure of an Ideal Bose Gas

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Handwritten derivation on a slide:

$$\beta V = k_B T \ln Q_+$$

$$\beta P V = \ln Q_+$$

$$\ln Q_+ = \frac{gV}{\lambda_T^3} \int_{S_2}^+ f_2(z) - g \ln(1-z)$$

$$\beta P V = \frac{gV}{\lambda_T^3} \int_{S_2}^+ f_2(z) - g \ln(1-z)$$

Approximation for $z \ll 1$:

$$z \ll 1$$

$$z = \frac{N}{N+g}$$

$$1-z = 1 - \frac{N}{N+g} = \frac{g}{N+g}$$

$$N \sim N_0 = \frac{gz}{1-z}$$

$$\sim \ln(N+g)$$

$$\sim \ln N$$

$$\beta P = \frac{g}{\lambda_T^3} \int_{S_2}^+ f_2(z) - \frac{g}{V} \ln(1-z) \sim \frac{1}{V} \ln N \sim \frac{1}{N}$$



Now, we know that $P V$ in the grand canonical ensemble was $k_B T \ln Q$ plus. So, that $\beta P V$ is going to be \ln of Q plus. So, we write down $\beta P V$ if you recall this expression of $\ln Q$ plus that we have \ln of Q plus was $\frac{gV}{\lambda_T^3} \int_{S_2}^+ f_2(z) - g \ln(1-z)$. So, this expression of $\beta P V$ takes the form $\frac{gV}{\lambda_T^3} \int_{S_2}^+ f_2(z) - g \ln(1-z)$.

Now, if you compare the first term in this expression which is this one, and the second term. You see that the second term can always be ignored in the thermodynamic limit, when Z is far away from 1 which means Z is less than very very less than 1 the high temperature limit.

Then I know that this quantity is 0 in contrast, when Z approaches 1 then I know that Z is going to be N over N plus g . We did that in our earlier lectures, when we said that the total particle number is compared to N naught which is equal to $g Z$ over 1 minus Z .

And this gives this gives us this expression for Z . So, therefore, 1 minus Z is going to be 1 minus N divided by N plus g which is going to be g over N plus g . So, that this term then as Z approaches 1 becomes the order of $\ln N$ plus g and since g is very very small compared to N , then this term is $\ln N$. In contrast the first term since there is a volume factor, I can divide throughout the volume fact from the left hand side and the right hand side.

And I am going to have βP is equal to g over $\lambda^3 T^{5/2}$ minus Z minus 1 by $V g$ times; \ln of 1 minus Z . And you see when Z tends to 1; then this goes as 1 by $V \ln N$ which is of the order of 1 over N . So, this in the thermodynamic limit vanishes, as well as when Z is much much less than 1, then when Z is much much less than 1 than this is just value then this 1 also vanishes is just 1 by V right.

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$$\begin{aligned}
 & \downarrow \\
 & \sim \ln(N+g) \quad 1-z = 1 - \frac{N}{N+g} = \frac{g}{N+g} \quad T > T_c \\
 & \sim \ln N \quad T < T_c \\
 \beta P &= \frac{g}{\lambda_T} \int_{S/2}^+ f_{S/2}(z) - \frac{g}{V} \frac{\ln(1-z)}{z} \quad \sim \frac{1}{V} \ln N \sim \frac{1}{N} \\
 \beta P &= \frac{g}{\lambda_T} \int_{S/2}^+ f_{S/2}(z) \\
 \text{For } T < T_c \quad \beta P &= \frac{g}{\lambda_T} \int_{S/2}^+ f_{S/2}(z) = \frac{g}{\lambda_T} \zeta(5/2) \\
 P &= g \zeta(5/2) \frac{k_B T}{\lambda_T} \quad \lambda_T \sim T^{3/2}
 \end{aligned}$$



So, therefore, in both the limiting cases when T is greater than T_c as well as when T is less than T_c , I have βP is equal to g over λ_T $\zeta(5/2)$. Now, for $T < T_c$ I have βP is equal to g over λ_T $\zeta(5/2)$ plus $\frac{1}{N}$ right. And this quantity is g over λ_T $\zeta(5/2)$.

So, that the thermodynamic pressure is $k_B T$ g well we will just write down like this $g \zeta(5/2) k_B T$ over λ_T recall that λ_T will go as T to the power 3 half.

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For $T < T_c$

$$\beta P = \frac{g}{\lambda_T} f_{5/2}^+(z=1) = \frac{g}{\lambda_T} \zeta(5/2)$$

$$P = g \zeta(5/2) \frac{k_B T}{\lambda_T}$$

$$= g \frac{\zeta(5/2) k_B}{\alpha} T^{5/2} \quad P \sim T^{5/2}$$


Does not depend on N (n)

$T > T_c$

$$f_m^+(z) = z + \frac{z^2}{2^m} + \dots$$

$$f_{5/2}^+(z) = z + \frac{z^2}{2^{5/2}} + \dots$$

$\alpha = \frac{N \lambda_T^3}{g V}$




And effectively therefore, you have $g \zeta(5/2) k_B$ some constant. So, I can write down λ_T is equal to αT to the power $3/2$ αT to the power $5/2$; which means that the thermodynamic pressure as for temperatures. When you are below the convince condensate Bose Einstein condensation searching, you have P going as T to the power $5/2$. What is interesting to note that this pressure does not depend on the number density N .

So, this does not depend on number density or rather we will say that it does not depend on N or n or 1 by or ρ that is also fine the density right. Now, for T greater than T_c I know the expansion of this function and that is going to be Z plus Z to the power 1 have Z square divided by m square sorry, it was k to the power n .

So, this is going to be 2 to the power m plus higher order terms, which means f plus of 5 by 2 Z is going to be Z plus Z square over 2 to the power 5 by 2 plus higher order terms.

And I have seen how the chemical potential depends on x equal to N lambda T over g V, we did this we saw that if I plot Z as a function of x, then for small x this goes linearly Z equal to x and beyond x equal to zeta 3 half this is 1. So, we are not looking in this part of the regime, but rather we are looking in the high temperature regime, where we are close to 0 Z 0 value of Z.

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$$f_{5/2}^+(z) \approx z = \frac{N \lambda T}{g V}$$

$$\beta P = \frac{g}{\lambda T} f_{5/2}^+(z) = \frac{g}{\lambda T} \cdot z = \frac{g}{\lambda T} \frac{N \lambda T}{g V} = \frac{N}{V}$$

$P = n k_B T$

→ High temperature limit for an ideal box Gas!!

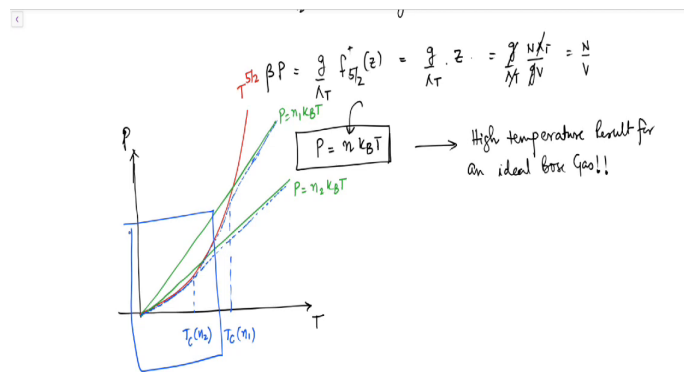


So, therefore, what we will do is we will substitute f plus 5 by 2 Z is equal is approximately equal to Z which is going to be N lambda T over g V. So, that I have pressure beta P which

was g over $\lambda^3 T^3$ times f plus $5/2$ by Z becomes g over $\lambda^3 T^3$ times Z , which is equal to g over $\lambda^3 T^3 N \lambda^3 T^3$ over $g V$.

So, that $\lambda^3 T^3$ and g and g cancels out and you have N over V . And you see you nicely recall recover the ideal gas result pressure is going to be $n k_B T$. This is the high temperature result for an Ideal Bose Gas, interesting in the high temperature limit you see that the pressure depends on the number density right.

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So, now, if I want to plot P as a function of temperature, then the first thing I have to fix is that this is the dependence when T is less than T_c . And I have no dependence on number density or density itself. So, if I do it like this, T to the power $5/2$ at in the high temperature limit I have $n k_B T$. So, I can for 2 densities I can draw it like no. So, let us draw it oops so,

let us draw T to the power $5/2$ like this way I have 2 densities in which case the pressure is this.

And in the other case the pressure P/V this is P is equal to $n^{1/2} k_B T$ and this is P is equal to $n^{3/2} k_B T$. And, then if I really measure the pressure, you will see that the pressure is going to follow this red line. And, then it is going to crossover to this line in the asymptotic limit of high temperature. And, similarly for the other one how do I do that? Well, let us do it like this it is going to follow this line up to here, when it starts to deviate and then go to the asymptotic limit.

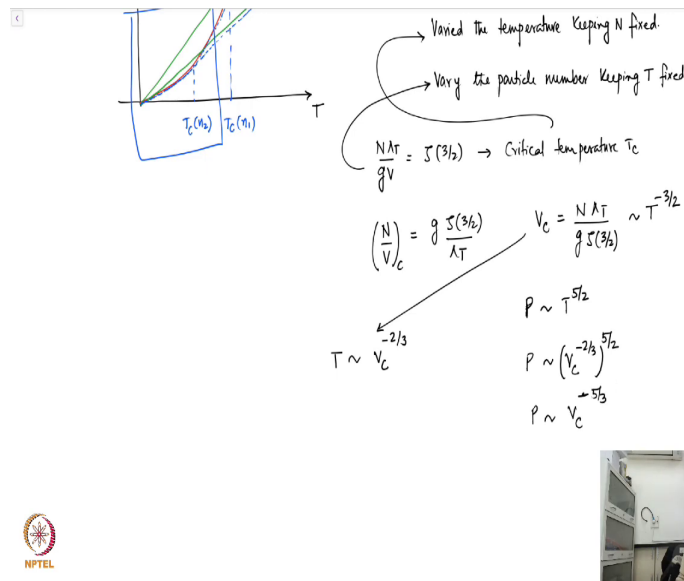
So, these would be the critical temperatures T_C of $n^{1/2}$ and these would be the this would be the critical temperature T_C of $n^{3/2}$. So, this is how your pressure temperature diagram would look for an ideal Bose gas in the high temperature limit, you are going to recover your ideal gas result.

And, in the low temperature limit below the critical temperature, you are going to recover that this pressure thermodynamic pressure of the Ideal Bose Gas is essentially independent of the density.

Now, this is reflective of the fact that below the critical temperature, if you keep on in decreasing the temperature then more and more number of particles see you have an upper limit on the number of particles you can put in right in the excited levels. So, you reach that limit and if you keep on decreasing the temperature, then more and more number of particles from the excited states come to the ground state.

But, in the ground state ϵ is equal to 0, they do not have any kinetic energy. And therefore, you do not get essentially any contribution to the pressure. And that is essentially reflected; here in this part of the diagram, where your pressure is independent of the density of the gas because, your condenser has started to appear in the system. Now just as. So, what did we do it?

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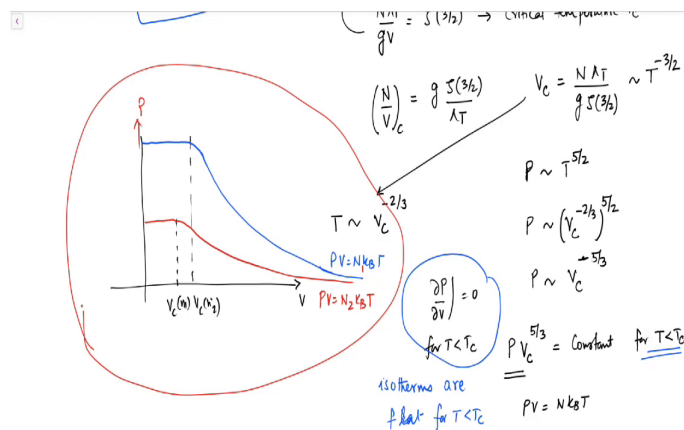


So, you so, essentially we varied the temperature keeping N fixed one can also vary the particle number keeping T fixed right, in which case the criteria that $N \lambda T$ over $g V$ which gave us $\zeta(3/2)$. This was the criteria which gave us the critical temperature T_c for this case. Now I can use the same relation to determine a density critical density in the second case, when I am varying the particle number keeping the temperature fixed.

So, that I will have N/V is equal to $g \zeta(3/2)$ and we will write this critical over λT . So, that V_c becomes $N \lambda T$ divided by $g \zeta(3/2)$. And this one you see scales as T to the power minus $3/2$, because λT scales as T to the power minus $3/2$ as well. Now, when I have this critical V_c scaling as T to the power minus $3/2$ and I have this expression for pressure, where pressure scaled as T to the power $5/2$ I can easily eliminate this.

So, that from this expression I can easily see that temperature will scale as V_C raised to the power minus two-third so, that the pressure will scale as V_C minus 2/3 raised to the power 5 by 2 and you have pressure scaling as V_C raised to the power 5 minus of five-third.

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So, that you have P of V_C raised to the power five-third is equal to constant for T less than T_C the moment. Therefore, the isotherms are very very curious here, why is curious? Because you see pressure does not depend on the volume again.

So, $\frac{\partial P}{\partial V}$ is going to be 0 for T less than T_C on the other hand for high temperatures you still have $P V$ is equal to $N K B T$. So, that if you plot the isotherms and let us say this is 1 volume V , V_C n 1 and the other 1 over here is V_C N of 2, then in the high temperature limit you know that this has to follow $P V$ is equal to $N K B T$. So, let it be so, this goes as P

P is equal to $\frac{N k_B T}{V}$, but below the critical temperature this is the pressure is nearly constant. So, it goes and saturates to this value.

This essentially means that the isotherms are flat for T less than T_C . Similarly here also you will see the sorry, you are going to get a flat region up to this value after which it slowly goes to P V is equal to so, we will call this N_1 and this will be $N_2 k_B T$. So, this is going to be your pressure volume diagram. So, this is how your P V diagram looks for an ideal Bose gas and this is how your P T diagram looks for an ideal Bose gas.