

PHYS 351 #4

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1 Q1

Recall that the van der Waals equation of state is $(P + \frac{a}{v^2})(v - b) = RT$, where v is the molar volume and a and b depend only on the type of gas.

1.1 a

The coefficient of thermal expansion is defined as follows:

$$\beta \equiv \frac{1}{V} \left(\frac{\partial V}{\partial T} \right) \Big|_P$$

Find β for a van der Waals gas. Show that this reduces to the ideal gas result, $\beta_{ideal} = \frac{1}{T}$, when $a = 0$ and $b = 0$.

Lets start by taking a $\frac{\partial}{\partial T}$ holding P as a constant. Noting that $v = V/n$ we begin taking a implicit derivative on both sides.

$$\begin{aligned} \frac{\partial}{\partial T} (P + \frac{an^2}{V^2})(V/n - b) &= \frac{\partial}{\partial T} RT \\ \left(\frac{-2an^2}{V^3} \frac{\partial V}{\partial T} \Big|_P \right) (V/n - b) + (P + \frac{an^2}{V^2}) (1/n \frac{\partial V}{\partial T} \Big|_P) &= R \\ \frac{\partial V}{\partial T} \Big|_P &= \frac{R}{\left(\frac{-2an^2}{V^3} \right) (V/n - b) + (P + \frac{an^2}{V^2}) (1/n)} \\ \frac{\partial V}{\partial T} \Big|_P &= \frac{Rv^2V}{(-2a)(v - b) + (Pv^2 + a)(v)} = \frac{Rv^2V}{2ab + (Pv^2 - a)v} \\ \beta &= \frac{Rv^2}{2ab - av + Pv^3} \end{aligned}$$

(Cool β is size independent, not dependent on V) If we take the ideal gas limit ($a=0$ $b=0$) we get:

$$\beta = \frac{R}{Pv}$$

Noting that $Pv=RT$ we see immediately:

$$\beta = \frac{1}{T}$$

Verifying the expected value for an ideal gas.

1.2 b

The coefficient of compression is defined as follows:

$$\kappa \equiv -\frac{1}{V} \left(\frac{\partial V}{\partial P} \right) \Big|_T$$

Find κ for a van der Waals gas. Show that this reduces to the ideal gas result, $\kappa_{ideal} = \frac{1}{P}$, when $a = 0$ and $b = 0$.

Lets start by taking a $\frac{\partial}{\partial P}$ holding T as a constant. Noting that $v = V/n$ we begin taking a implicit derivative on both sides.

$$\begin{aligned} \frac{\partial}{\partial P} (P + \frac{an^2}{V^2})(V/n - b) &= \frac{\partial}{\partial P} RT \\ (1 + \frac{-2an^2}{V^3} \frac{\partial V}{\partial P} \Big|_T)(V/n - b) + (P + \frac{an^2}{V^2})(1/n \frac{\partial V}{\partial P} \Big|_T) &= 0 \\ (1 + \frac{2a}{v^2} \kappa)(v - b) - (P + \frac{a}{v^2})(v\kappa) &= 0 \\ 2a\kappa(v - b) - (Pv^2 + a)(v\kappa) &= (b - v)v^2 \\ \kappa &= \frac{(v - b)v^2}{2ab + Pv^3 - av} \end{aligned}$$

This term is also size independent. We note that as a and b become 0 the above becomes $\kappa_{ideal} = \frac{1}{P}$.

2 Q2

We have an elastic band. The Youngs modulus Y is defined by the linear relation between a differential change in length, dL , that results from the application of a differential increment of force, dF : $dF = \frac{YA}{L_o} dL$, where the length is L_o when no force is applied, and the cross-sectional area is A .

2.1 a

Let the tension on the band be increased quasi-statically from T_1 to T_2 . Calculate the work done on the band, assuming that the cross-section does not change appreciably.

Basically we are to assume that Y , A , L_o are all constant. this would mean that $F(L)$ the only two variables left in the system.

$$\int dF = \int \frac{YA}{L_o} dL \implies F = \frac{YA}{L_o} * L + C \implies F = \frac{YA}{L_o} * (L - L_o)$$

The constant is determined such that at $L = L_o$, $F = 0$. We also know that:

$$dw = FdL = \frac{FL_o}{YA} dF$$

We can now integrate our differential work:

$$\int dw = \int_{T_1}^{T_2} \frac{FL_o}{YA} dF = \frac{F^2 L_o}{2YA} \Big|_{T_1}^{T_2} = \frac{((T_2)^2 - (T_1)^2)L_o}{2YA}$$

2.2 b

Assume that the heat capacity of the elastic band does not change during the stretching process described in part (a), and assume that process occurs adiabatically. How does the temperature of the band change during the stretching process? Does this agree with observation?

Our model has all of the energy being stored in the system, and being perfectly reversible. We would expect no change in the temperature of the band as it is stretched as all of the energy is going into the stretching of the material. This is different from what we observe in the laboratory as stretching and retracting a material will not be a perfectly reversible process and we will see temperature increase.