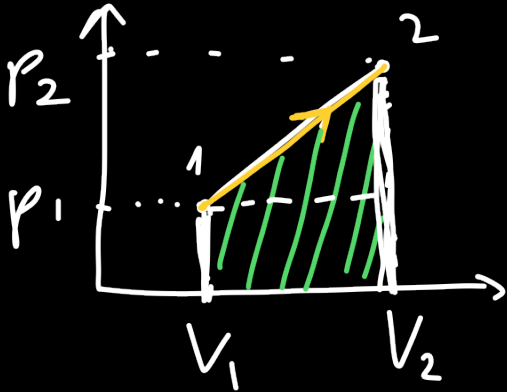


Homework 28

1.



certain amount
of gas.

$$p_1 = 10^5 \text{ Pa}$$

$$p_2 = 2 \cdot 10^5 \text{ Pa}$$

$$V_2 = 2 \cdot V_1 = 0.2 \text{ m}^3$$

$$W_{\text{gas}} = \frac{1}{2} (V_2 - V_1) \cdot (p_2 - p_1) + p_1 (V_2 - V_1)$$

$$= (V_2 - V_1) \left(\frac{1}{2} (p_2) - \frac{1}{2} p_1 + p_1 \right)$$

$$= \frac{1}{2} (p_1 + p_2) (V_2 - V_1)$$

$$= \frac{3}{2} \cdot 10^5 \cdot 0.1 \text{ Pa} \cdot \text{m}^3$$

$$W_{\text{gas}} = 15 \text{ kJ}$$

$$\textcircled{2} \quad Q = 1000 \text{ cal} = 4.186 \text{ kJ}$$

$$Q = W + 0$$

$$W_1 = F_{\text{grav}} \cdot h = mgh$$

$$W_{\text{total}} = mgh \cdot N$$

$$N = \frac{4.186 \cdot 10^3 \text{ J}}{20 \text{ kg} \cdot 9.8 \frac{\text{N}}{\text{kg}} \cdot 2 \text{ m}} \approx 7.12$$

$$\textcircled{3} \quad p = 10^5 \text{ Pa}, \quad \Delta V = 1.0 \text{ liters.}$$

$$Q = 300 \text{ J}, \quad \Delta E_{\text{int.}} = ?$$

$$W_{\text{gas}} = p \cdot \Delta V = 10^5 \text{ Pa} \cdot 10^{-3} \text{ m}^3 = 100 \text{ J}$$

$$Q = \Delta E_{\text{int}} + W_{\text{gas}} \Rightarrow$$

$$\Delta E_{\text{int}} = 300 \text{ J} - 100 \text{ J} = 200 \text{ J}$$

Classwork The first law of thermodynamics.

Specific heat of an ideal gas

$$Q = m \cdot c \cdot \Delta T = n \cdot c^{\text{molar}} \cdot \Delta T$$

c^{molar} — molar specific heat.

First law of thermodynamics:

$$Q = W_{\text{gas}} + \Delta E_{\text{int.}} \quad \textcircled{\text{I}}$$

Focus on the ideal gas!

$W_{\text{gas}} \rightarrow p \cdot \Delta V$ when $p = \text{const}$
or area under the graph.
with $\pm$ sign.

Work done on the gas $\leftrightarrow W_{\text{gas}}$

$$W_{\text{on gas}} = -W_{\text{gas}}$$

Second form:

$$Q = \Delta E_{\text{int}} - W_{\text{on gas}}$$

$$Q + W_{\text{on gas}} = \Delta E_{\text{int}}$$

heat
transferred
to gas

work done on gas

change
in the
internal
energy.

What is ΔE_{int} ?

$$E_{\text{internal}} = PE + KE$$

In ideal gas: KE PE

$$PE = 0, KE > 0$$

$\begin{matrix} < \\ > \\ = \end{matrix} \begin{matrix} I \\ III \\ I \end{matrix}$

$$E_{\text{int}} = KE_{\text{tot}}$$

$$KE_{av.} = \frac{KE_1 + \dots + KE_N}{N} = \frac{KE_{tot}}{N}$$

$N \leftarrow \# \text{ of molecules}$

$$KE_{av.} = \frac{3}{2} K_B \cdot T$$

monoatomic gases.

$$K_B = 1.38 \cdot 10^{-23} \text{ J/K}$$

$$KE_{tot} = \frac{3}{2} N \cdot K_B \cdot T$$

$$n = N/N_A \Rightarrow N = n \cdot N_A$$

$$KE_{tot} = \frac{3}{2} n \cdot N_A \cdot K_B \cdot T$$

$$N_A \cdot K_B = 6.02 \cdot 10^{23} \frac{1}{\text{mole}} \cdot 1.38 \cdot 10^{-23} \frac{\text{J}}{\text{K}}$$

$$= 8.31 \frac{\text{J}}{\text{mole} \cdot \text{K}} = R$$

universal gas
constant!

Finally:

$$KE_{tot} = \frac{3}{2} n R T$$

$$\Delta E_{int} = \Delta KE_{tot} = \frac{3}{2} n R \Delta T$$

$$Q = W_{\text{gas}} + \Delta E_{\text{int.}}$$

univ.
gas const.

$$\hookrightarrow \boxed{n \cdot C^{\text{molar}} \cdot \Delta T = W_{\text{gas}} + \frac{3}{2} n R \Delta T}$$

molar specific heat

C^{molar} depends on the process

if $V = \text{const.} : W_{\text{gas}} = 0$

$$\Rightarrow \boxed{\cancel{n} \cdot C_v^{\text{molar}} \cdot \cancel{\Delta T} = 0 + \frac{3}{2} \cancel{n} R \cancel{\Delta T}}$$

$$\boxed{C_v^{\text{molar}} = \frac{3}{2} R}$$

$$\boxed{C_p^{\text{molar}} = R + \frac{3}{2} R = \frac{5}{2} R}$$

$$\uparrow W_{\text{gas}} = p \cdot \Delta V = n R \Delta T$$

