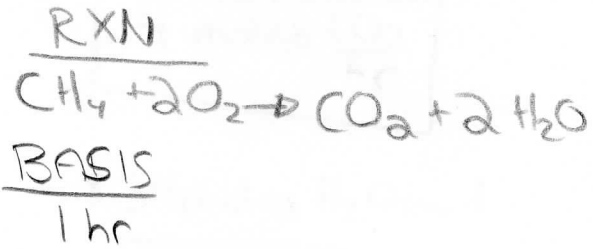
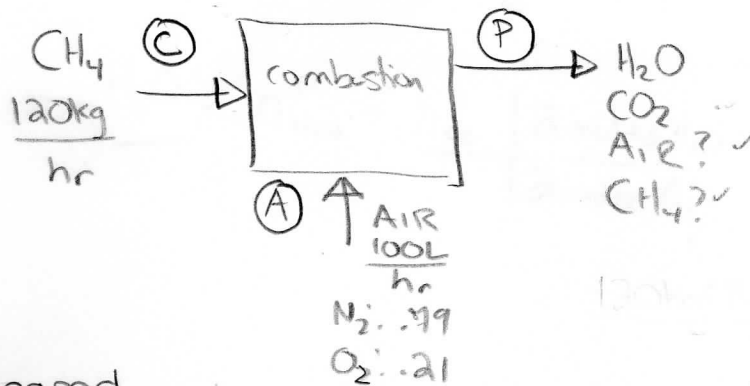


Problem #1 (25 points)

If 2kg/min of CH_4 and 100L/hour of air enter and exit from a combustion engine at room temperature and react, what is the flow rate of each of the molecular species exiting the combustion engine?

Assumptions

- AIR is an ideal gas
- reaction goes to completion

Legend

\dot{n}_i = moles of species i in stream j

Find Limiting reactant

is $\frac{\dot{n}_{\text{CH}_4}^{\text{C}}}{\dot{n}_{\text{O}_2}^{\text{A}}} >, <, \text{ or } = \frac{1}{2}$?

$$\dot{n}_{\text{CH}_4}^{\text{C}} = \frac{2 \text{ kg}}{16 \text{ kg/mol}} = 0.125 \text{ kmol} = 125 \text{ moles}$$

$$\dot{n}_{\text{O}_2}^{\text{A}} = \frac{PV}{RT} = \frac{(101,300 \text{ Pa})(100 \text{ L} \cdot \frac{10^{-3} \text{ m}^3}{1 \text{ L}})(.21)}{8.314 \frac{\text{Pa} \cdot \text{m}^3}{\text{mol} \cdot \text{K}} 298 \text{ K}} = 0.859 \text{ moles O}_2$$

$\rightarrow \text{O}_2$ is limiting reagent $\dot{n}_{\text{O}_2}^{\text{A}} < \dot{n}_{\text{CH}_4}^{\text{C}}$

Material Balance

$$\dot{n}_{\text{CH}_4}^{\text{C}} - \dot{n}_{\text{CH}_4}^{\text{reacted}} = \dot{n}_{\text{CH}_4}^{\text{P}}$$

$$\dot{n}_{\text{CH}_4}^{\text{reacted}} = \frac{\dot{n}_{\text{O}_2}^{\text{A}}}{2} \cdot 1 \text{ mol CH}_4 = 0.4295 \text{ moles CH}_4 \text{ react}$$

$$125 \text{ moles} - 0.4295 \text{ moles} = \boxed{124.57 \text{ moles CH}_4 \text{ exit}}$$

$$\text{O}_2: \dot{n}_{\text{O}_2}^{\text{A}} - \dot{n}_{\text{O}_2}^{\text{reacted}} = \dot{n}_{\text{O}_2}^{\text{P}}$$

$$\dot{n}_{\text{O}_2}^{\text{A}} = \dot{n}_{\text{O}_2}^{\text{reacted}} \rightarrow \boxed{0 \text{ moles O}_2 \text{ exit}}$$

$$\text{N}_2: \dot{n}_{\text{N}_2}^{\text{A}} = \dot{n}_{\text{N}_2}^{\text{P}} =$$

$$\rightarrow \boxed{3 \text{ moles N}_2 \text{ exit}}$$

$$\dot{n}_{\text{N}_2}^{\text{A}} = \frac{PV}{RT} = \frac{(101,300 \text{ Pa})(100 \text{ L} \cdot \frac{10^{-3} \text{ m}^3}{1 \text{ L}})(.79)}{8.314 \frac{\text{Pa} \cdot \text{m}^3}{\text{mol} \cdot \text{K}} 298 \text{ K}} = 3.2 \text{ moles N}_2$$

Problem #1 (cont'd)

$$\underline{\text{CO}_2}: \eta_{\text{CO}_2}^P = \frac{\eta_{\text{O}_2}^A}{2 \text{ moles O}_2} \left| \frac{1 \text{ mol CO}_2}{1 \text{ mol CO}_2} \right| = 0.4295 \text{ moles CO}_2 \text{ out}$$

$$= \boxed{0.4 \text{ moles } \frac{\text{CO}_2}{\text{hr}}}$$

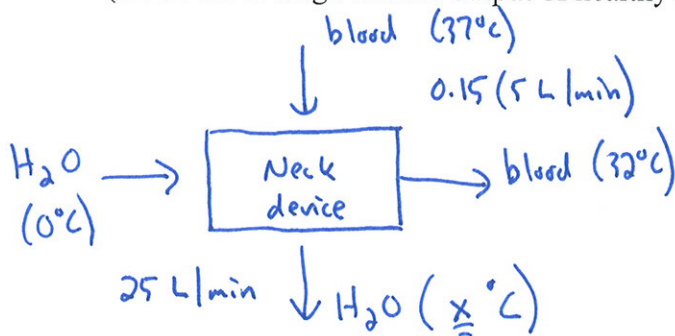
$$\underline{\text{H}_2\text{O}}: \eta_{\text{H}_2\text{O}}^P = \frac{\eta_{\text{O}_2}^A}{2 \text{ moles O}_2} \left| \frac{2 \text{ moles H}_2\text{O}}{2 \text{ moles O}_2} \right| = 0.859 \text{ moles H}_2\text{O out}$$

$$= \boxed{0.9 \text{ moles } \frac{\text{H}_2\text{O}}{\text{hr}}}$$

Problem #2 (25 points)

In order to cryoprotect the brain prior to neurosurgery, ice-cold water is pumped at a rate of 25 L/min through a device that wraps around the neck and is 75% efficient. If the brain is to be cooled by 5°C and 15% of the total blood volume goes to the head, what is the temperature of the water as it exits the device?

(Note: the average cardiac output of healthy adults is 5 L/min & $C_v \text{H}_2\text{O}(l) = 4.18 \text{ J/g}\cdot\text{K}$)



Basic: 1 min

open, steady-state

Assume: $C_{v \text{H}_2\text{O}} \approx C_{v \text{blood}} (= 4.18 \frac{\text{J}}{\text{g}\cdot\text{K}})$
 $\rho_{\text{H}_2\text{O}} \approx \rho_{\text{blood}} (= 1.0 \text{ g/mL})$

$$C_v = C_p \quad \text{or} \quad \Delta H = \Delta U$$

$$\Delta E = 0 = \Delta (U + KE + PE) = \overset{\text{system stationary}}{\cancel{Q}} + \overset{\text{insulated}}{\cancel{W}} - \Delta (H + \overset{\text{no height diff}}{\cancel{PE}} + \overset{\text{no velocity change}}{\cancel{KE}})$$

$$\Rightarrow 0 = \Delta U, \quad 0 = \Delta H = \Delta H_{\text{blood}} + \underbrace{\Delta H_{\text{H}_2\text{O}}}_{\times 0.75 \text{ (efficiency)}}$$

$$\Rightarrow 0 = (0.75 \text{ kg}) \left(4.18 \frac{\text{J}}{\text{g}\cdot\text{K}} \right) (-5 \text{ K}) + 0.75 \left[(25 \text{ kg}) \left(4.18 \frac{\text{J}}{\text{g}\cdot\text{K}} \right) (x - 273 \text{ K}) \right]$$

$$0 = m \int_{T_i}^{T_f} C_p^{\text{blood}} dT + \left[m \int_{T_i}^{T_f} C_p^{\text{H}_2\text{O}} dT \right] 0.75$$

$$15.675 \text{ kJ} = 78.375 \frac{\text{kJ}}{\text{K}} (x - 273 \text{ K})$$

$$\boxed{0.2^\circ\text{C}} = x - 273 \text{ K}$$

Switch
order

Problem #3 (25 points)

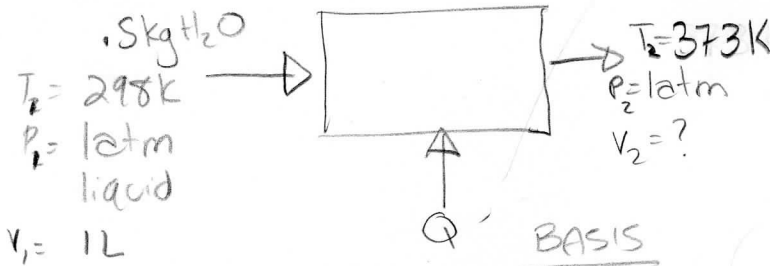
A piston chamber originally at 1 L and containing 0.5 kg of water at room temperature is heated to the point that all of the water evaporates and the chamber expands to maintain constant atmospheric pressure. In this case, how much energy enters the chamber?

$$\Delta \hat{H}_{\text{vaporization}} (100^\circ\text{C}) = 40.65 \text{ kJ/(g mol)}$$

$$\Delta \hat{H}_{\text{fusion}} (0^\circ\text{C}) = 60.1 \text{ kJ/(g mol)}$$

$$C_p \text{ H}_2\text{O (l)} = 75.4 \text{ J/(g mol)}$$

$$C_p \text{ H}_2\text{O (v)} = 33.9 \text{ J/(g mol)}$$



Assume

- energy change due to heating bare dryer in vessel is negligible
- no heat lost to surroundings
- kinetic energy, potential energy changes are negligible

Energy Balance

$$Q = \Delta U \quad (\text{closed, unsteady})$$

$$\Delta U = \Delta H - \Delta(PV)$$

$$Q = \Delta U = n_{\text{H}_2\text{O}} \int_{T_1}^{T_2} C_{p \text{ H}_2\text{O (l)}} dT + n_{\text{H}_2\text{O}} \Delta \hat{H}_{\text{vap}} - (P_2 V_2 - P_1 V_1)$$

Find $n_{\text{H}_2\text{O}}$ (moles H_2O)

$$n = \frac{500 \text{ g H}_2\text{O}}{18 \text{ g/mol}} = 27.8 \text{ moles}$$

Find ΔV

$$V_2 = \frac{nRT}{P} = \frac{27.8 \text{ moles} \left(8.314 \frac{\text{J}}{\text{mol K}} \right) (373 \text{ K})}{101,300 \text{ Pa}} = 0.851 \text{ m}^3$$

$$V_1 = \frac{100 \text{ L} \cdot 0.001 \text{ m}^3}{1 \text{ L}} = 0.1 \text{ m}^3$$

$$\Delta U = 27.8 \text{ moles} \left(75.4 \frac{\text{J}}{\text{g mol}} \right) (373 - 298 \text{ K}) + 27.8 \text{ moles} \left(40,656 \frac{\text{J}}{\text{g mol}} \right) - 101,300 \text{ Pa} (0.851 \text{ m}^3 - 0.1 \text{ m}^3)$$

$$= 1,572,095 \text{ J} + 1,130,000 \text{ J} - 760,766 \text{ J}$$

$$= 1,211,000 \text{ J} = 1211 \text{ kJ} \Rightarrow \boxed{Q = 1000 \text{ kJ}}$$

Problem #4 (25 points)

If the heat of combustion for 1 mole of glucose ($C_6H_{12}O_6$) at standard conditions is -2825.8 kJ/g mol , and the heats of formation for water and carbon dioxide are -285.8 kJ/g mol and -393.5 kJ/g mol (respectively), what is the heat of formation for glucose?



Basic: 1 g/mol
 $C_6H_{12}O_6$

$$\Delta H_{rxn} = \sum w_i \cdot \hat{H}_f(\text{prod}) - \sum w_i \cdot \hat{H}_f(\text{react})$$

$$-2825.8 \text{ kJ} = \left[\underbrace{6(-285.8 \text{ kJ})}_{-1714.8} + \underbrace{6(-393.5)}_{-2361} \right] - \left[\hat{H}_f^{\text{glucose}} + 6(\cancel{x}) \right]$$

$$\Rightarrow \hat{H}_f^{\text{glucose}} = \boxed{1250 \text{ kJ/g mol}}$$