1. You have four identical 1.00 L **unbreakable** containers filled with gases. The molar masses of the gases are given in curly brackets {}. Assume all <u>gases are ideal</u>.

Flask A:	1.0 mol He	{4.00 g/mol}	100 K
Flask B:	0.40 mol CO	{28.01 g/mol}	400 K
Flask C:	1.0 atm Cl_2	{70.91 g/mol}	298 K
Flask D:	$1.0 atm NO_2$	{46.01 g/mol}	273 K

(a) Which flask has the greatest pressure?

Flasks C and D are both at 1 atm pressure. Determine the pressure of flasks A and B: $P_{A} = \frac{nRT}{v} = \frac{(1.0 \text{ mol})(0.08206 \frac{L \cdot atm}{mol \cdot K})(100 \text{ K})}{1.0 \text{ L}} = 8.2 \text{ atm} \quad P_{B} = \frac{nRT}{v} = \frac{(0.40 \text{ mol})(0.08206 \frac{L \cdot atm}{mol \cdot K})(400 \text{ K})}{1.0 \text{ L}} = 13 \text{ atm}$ Answer: Flask B

- (b) Which flask is at STP? Recall that STP is T = 0 °C = 273 K and P = 1.0 atm. Answer: Flask D
- (c) In which flask would the contents take up the smallest volume if brought to STP? Recall that at STP, the volume of 1 mol of any ideal gas is 22.4 L. Therefore, the volume is directly proportional to the number of moles of each gas (V_{STP} α n). Flask A has 1.0 mol of gas and flask B has 0.40 mol of gas. Determine the number of moles in flasks C and D: $n_{C} = \frac{PV}{RT} = \frac{(1.0 \text{ atm})(1.0 \text{ L})}{(0.08206 \frac{L \text{ atm}}{100 \text{ kK}})^{(298 \text{ K})}} = 0.041 \text{ mol Cl}_{2}$ $n_{D} = \frac{PV}{RT} = \frac{(1.0 \text{ atm})(1.0 \text{ L})}{(0.08206 \frac{L \text{ atm}}{100 \text{ kK}})^{(273 \text{ K})}} = 0.045 \text{ mol NO}_{2}$

Answer: Flask C

(d) In which flask will diffusion of the gas be fastest?



2. A gas tank has a volume of 32.0 L, a temperature of 27.0 °C, a pressure of 1.10×10^5 Torr, and contains 748 g of an <u>unknown</u> gas. What is the identity of the gas?

Determine the number of moles of unknown gas (X) using the ideal gas law:

$$n_{\rm X} = \frac{PV}{RT} = \frac{\left(2125 \text{ psi} \times \frac{1 \text{ atm}}{14.7 \text{ psi}}\right)(32 \text{ L})}{\left(0.08206 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right)(27 + 273.15 \text{ K})} = 18_{7.8} \text{ mol X}$$

We can determine the molar mass of gas X (MM_X)now:

$$MM_{X} = \frac{mass_{X}}{n_{X}} = \frac{748 \text{ g X}}{18_{7.8} \text{ mol } X} = 4.0 \frac{\text{g}}{\text{mol}}$$

The gas is <u>helium</u> (He).

 $6 O_2(g) + C_6 H_{12}O_6(aq) \rightarrow 36 \text{ ATP}(aq) + 6 CO_2(g) + 6 H_2O(l)$

What volume of O_2 (at STP) is consumed in 30.0 minutes? Determine number of moles of O_2 required:

 $30 \min \times \frac{1 \text{ hr}}{60 \min} \times \frac{31 \text{ g } \text{ O}_2}{1 \text{ hr}} \times \frac{1 \text{ mol } \text{ O}_2}{32.00 \text{ g } \text{ O}_2} = 0.48_4 \text{ mol } \text{ O}_2$

Determine volume of this much O_2 using the ideal gas law:

$$V_{O_2} = \frac{n_{O_2}RT}{P} = \frac{(0.48_4 \text{ mol } O_2) \left(0.08206 \frac{L \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right) (273.15 \text{ K})}{1.00 \text{ atm}} = 11 \text{ L}$$

4. Consider the following arrangement of two flasks at 400.0 K, connected by a stopcock. Assume that the gases are ideal and the tube connecting the two flasks has negligible volume.



(a) Assuming <u>no chemical reaction</u> between NO and O₂, calculate the partial pressures of NO and O₂ if the stopcock were opened. Assume no temperature changes.

Use pressure-volume gas relationship to find new partial pressures: $P_1V_1 = P_2V_2$ $P_{0_2} = \frac{(100.0 \text{ Torr})(2.0 \text{ L})}{6.0 \text{ L}} = 33 \text{ Torr}$ $P_{N0} = \frac{(600.0 \text{ Torr})(4.0 \text{ L})}{6.0 \text{ L}} = 4.0 \times 10^2 \text{ Torr}$

(b) Now assume that when the stopcock is opened, the NO and O₂ react to form NO₂. Assume that there is no temperature changes.

Calculate the partial pressures of NO and O₂ after the reaction is complete. $2NO(g) + O_2(g) \rightarrow 2NO_2(g)$

First, determine the number of moles of each reactant through the ideal gas law:

$$n_{NO} = \frac{PV}{RT} = \frac{\left(\frac{600.0 \text{ Torr} \times \frac{1 \text{ atm}}{760 \text{ Torr}}\right)(4.0 \text{ L})}{\left(0.08206 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right)(400.0 \text{ K})} = 0.096_2 \text{ mol NO}$$
$$n_{O_2} = \frac{PV}{RT} = \frac{\left(100.0 \text{ Torr} \times \frac{1 \text{ atm}}{760 \text{ Torr}}\right)(2.0 \text{ L})}{\left(0.08206 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right)(400.0 \text{ K})} = 0.0080_2 \text{ mol O}_2$$

Second, determine that the limiting reactant is O₂:

$0.096_2 \text{ mol NO} \times \frac{1 \text{ mol } O_2}{2 \text{ mol NO}} = 0.048_1 \text{ mol } O_2$	$0.096_2 \operatorname{mol} \mathrm{NO} \times \frac{2 \operatorname{mol} \mathrm{NO}_2}{2 \operatorname{mol} \mathrm{NO}} = 0.096_2 \operatorname{mol} \mathrm{NO}_2$
	$0.0080_2 \mod O_2 \times \frac{2 \mod NO_2}{1 \mod O_2} = 0.016_0 \mod NO_2$
\rightarrow We have less O ₂ than we need.	$\rightarrow O_2$ produces less NO ₂ product.

Since O_2 is limiting, none will be left after the reaction. However, some NO gas will be left over: $2 \mod NO$

$$0.0080_2 \text{ mol } 0_2 \times \frac{2 \text{ mol NO}}{1 \text{ mol } 0_2} = 0.016_0 \text{ mol NO reacted}$$

 $n_{\text{NO,left over}} = 0.096_2 \text{ mol} - 0.016_0 \text{ mol} = 0.080_2 \text{ mol NO left over}$

Third, find the partial pressures of NO gas leftover:

$$\mathbf{P_{NO}} = \frac{nRT}{V} = \frac{(0.080_2 \text{ mol}) \left(0.08206 \frac{\text{L} \cdot \text{atm}}{\text{mol} \cdot \text{K}}\right) (400.0 \text{ K})}{(2.0 \text{ L} + 4.0 \text{ L})} = 0.43_9 \text{ atm} \times \frac{760 \text{ Torr}}{1 \text{ atm}} = 330 \text{ Torr}$$