

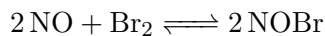
Chapter 6 Challenge Problem Solutions

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6.97

Problem: Nitric oxide and bromine at initial pressures of 98.4 and 41.3 torr, respectively, were allowed to react at 300. K. At equilibrium the total pressure was 110.5 torr. The reaction is as follows.



- a) Calculate the value of K_p
- b) What would be the partial pressures of all species if NO and Br₂, both at an initial partial pressure of 0.30 atm, were allowed to come to equilibrium at this temperature?

Solution: We model the change in pressure of each substance through an ICE table and note the concentrations.

	[2 NO]	+	[Br ₂]	\rightleftharpoons	[2 NOBr]
I	98.4		41.3		
C	-2x		-x		+2x
E	(98.4 - 2x)		(41.3 - x)		(+2x)

We are told that $P_E = 110.5$ torr, meaning that we can solve for x

$$\begin{aligned}P_E &= P_{\text{NO}} + P_{\text{Br}_2} + P_{\text{NOBr}} \\110.5 &= (98.4 - 2x) + (41.3 - x) + (2x) \\x &= 29.2 \text{ torr}\end{aligned}$$

Solving for each equilibrium concentration gives us a K_p value of

$$K_p = \frac{58.4^2}{40.0^2 \times 12.1} = 0.176 \text{ torr}^{-1} = \boxed{134 \text{ atm}^{-1}}$$

Solving for the second part of the problem becomes a bit more convoluted. We are given that the initial concentrations of the reactants NO and Br₂ are 0.3 atm each.

Writing an ICE table for the following reaction is trivial and is omitted. We end with a K_p of

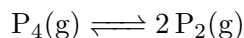
$$134 = \frac{(2x)^2}{(0.3 - 3x)^2(0.3 - x)}$$

Solving for x is complicated to do algebraically, but graphically, we see that $x = 0.125$ atm. Knowing that the equilibrium concentrations of NO and Br₂ are $(0.3 - 2x)$ and $(0.3 - x)$, and NOBr is $2x$, we see that the partial pressures of each of the substances are

$$P_{\text{NO}} = 0.052 \text{ atm} \quad P_{\text{Br}_2} = 0.18 \text{ atm} \quad P_{\text{NOBr}} = 0.25 \text{ atm}$$

6.99

Problem: Consider the reaction



where $K_p = 1.00 \times 10^{-1}$ at 1325 K. In an experiment where P₄(g) was placed in a container at 1325 K, the equilibrium mixture of P₄(g) and P₂(g) has a total pressure of 1.00 atm. Calculate the equilibrium pressures of P₄(g) and P₂(g). Calculate the fraction (by mole) of P₄(g) that has dissociated to reach equilibrium.

Solution: In this problem, we have a few unknowns, including the original pressure of P₄(g) and of course the equilibrium pressures of P₄(g) and P₂(g). Our basic ICE table gives us two unknowns, the initial concentration of P₄(g) y , and the change variable x . I have omitted the ICE table again for this problem, but writing it out would give us the equilibrium concentrations of the two substances in terms of y and x .

$$P_4(\text{g}) = y - x$$

$$P_2(\text{g}) = 2x$$

The problem statement also gives us the total pressure as $P = 1.00$ atm, which we can rewrite to get our system of equations

$$K_p = \frac{(2x)^2}{(y - x)}$$

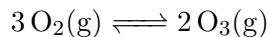
$$P = (y - x) + (2x)$$

Solving this system of equations is simply algebraic manipulation, which gives us $y = 0.865$ atm, and $x = 0.135$ atm. Finding our answers is simply arithmetic.

$$P_{P_4} = 0.73 \text{ atm} \quad P_{P_2} = 0.270 \text{ atm} \quad 16\% \text{ of } P_4 \text{ decomposed}$$

6.101

Problem: Consider the reaction



At 175 °C and a pressure of 128 torr an equilibrium mixture of O₂ and O₃ has a density of 0.168 g/L. Calculate the K_p for the above reaction at 175 °C.

Solution: We first must come up with a method of finding density in terms of pressure. Density's units are g L⁻¹, and the equation that relates pressure and volume is

$$PV = nRT$$

However, we must manipulate this equation to introduce mass. We do this by introducing molar mass (M) and obtain

$$PM = M \frac{n}{V} RT = \rho RT$$

$$\frac{PM}{RT} = \frac{P_{\text{O}_2} M_{\text{O}_2} + P_{\text{O}_3} M_{\text{O}_3}}{RT} = \rho = 0.168$$

We also know that $P = 128$ torr, or 0.168 atm. Our second equation is

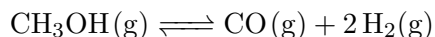
$$P = P_{\text{O}_2} + P_{\text{O}_3} = 0.168$$

This is a simple system of equations with two unknowns (P_{O_2} and P_{O_3}). Once solved, we find that $P_{\text{O}_2} = 0.118$ atm and $P_{\text{O}_3} = 0.05$ atm. Which gives us a final answer

$$K_p = \frac{P_{\text{O}_3}^2}{P_{\text{O}_2}^3} = 1.5 \text{ atm}^{-1}$$

6.103

Problem: a 4.72 g sample of methanol (CH_3OH) was placed in an otherwise empty 1.00 L flask and heated to 250°C to vaporize the methanol. Over time the methanol vapor decomposed by the following reaction:



After the system has reached equilibrium, a tiny hole is drilled in the side of the flask allowing gaseous compounds to effuse out of the flask. Measurements of the effusing gas show that it contains 33.0 times as much $\text{H}_2(\text{g})$ as $\text{CH}_3\text{OH}(\text{g})$. Calculate the K for this reaction at 250°C .

Solution: We are given that the effusing gas shows 33 times as much H_2 as CH_3OH . However, this does not account for the the differing rate of effusion for the two gasses. Using Graham's law of effusion gets us an effusion ratio of

$$\frac{\text{Rate}_{\text{H}_2}}{\text{Rate}_{\text{CH}_3\text{OH}}} = \sqrt{\frac{M_{\text{CH}_3\text{OH}}}{M_{\text{H}_2}}} = \sqrt{\frac{32}{2}} = 4.$$

We now know that H_2 effuses at 4 times the rate of CH_3OH , however, we are told that there is 33 times as much H_2 . Since the effusion rate does not account for enough of an increase, the other $\frac{33}{4} = 8.25$ factor comes from the greater concentration of H_2 at equilibrium. The mole ratio of H_2 to CH_3OH is thus

$$\frac{n_{\text{H}_2}}{n_{\text{CH}_3\text{OH}}} = 8.25$$

We start with 4.72 grams of CH_3OH , or 0.147 mol. The ICE Table is trivial and is omitted. Once written however, we see at equilibrium

$$8.28 = \frac{n_{\text{H}_2}}{n_{\text{CH}_3\text{OH}}} = \frac{2x}{0.147 - x}$$

$$x = 0.118$$

Note that because the reaction takes place in a 1 liter vessel, the number of moles of any substance is identical in number to the concentration.

$$K = \frac{4x^3}{0.147 - x} = 0.23$$