

Sample problem

K_p for the following reaction is 0.16 at 25 °C. What is the value of K_c ?



- $K_P = K_C(RT)^{\Delta n}$
- $0.16 = K_c(0.08206 \text{ L atm mol}^{-1} \text{ K}^{-1} \times 298.15 \text{ K})^{3-2}$
- **$K_c = 6.5 \times 10^{-3}$**

Sample problem

$K_c = 5.6 \times 10^{-12}$ at 500 K for the dissociation of iodine molecules to iodine atoms.



A mixture has $[\text{I}_2] = 0.020 \text{ M}$ and $[\text{I}] = 2.0 \times 10^{-8} \text{ M}$. Is the reaction at equilibrium at 500 K? If not, which way must the reaction proceed to reach equilibrium?

- $Q = \frac{[\text{I}]^2}{[\text{I}_2]} = \frac{(2.0 \times 10^{-8})^2}{0.020} = 2.0 \times 10^{-14}$
- $Q < K_c \rightarrow$ too many reactant particles
- Reaction will shift to the **right** (to make more product)

Sample problem

In aqueous solution Fe^{3+} ions react with I^- ions to give Fe^{2+} ions and I_3^- ions. Suppose the initial concentration of Fe^{3+} is 0.200 M, the initial I^- concentration is 0.300 M, and the equilibrium concentration of I_3^- ions is 0.0866 M. What is the value of K_c ?

	2Fe^{3+}	$+ 3 \text{I}^-$	\rightleftharpoons	2Fe^{2+}	$+ \text{I}_3^-$
I	0.200 M	0.300 M		0	0
C	$- 2x$	$- 3x$		$+ 2x$	$+ x$
E	$0.200 - 2x$	$0.300 - 3x$		$2x$	$x = 0.0866 \text{ M}$

$$K_c = \frac{[\text{Fe}^{2+}]^2 [\text{I}_3^-]}{[\text{Fe}^{3+}]^2 [\text{I}^-]^3} = \frac{(2 \times 0.0866)^2 (0.0866)}{(0.200 - 2 \times 0.0866)^2 (0.300 - 3 \times 0.0866)^3} = 5.6 \times 10^4$$

Sample problem

The equilibrium constant K_c ($=55.64$) for the reaction $\text{H}_2 (\text{g}) + \text{I}_2 (\text{g}) \rightleftharpoons 2 \text{HI} (\text{g})$ has been determined at 425°C . If 0.130 mol each of H_2 and I_2 is placed in a 25.0-L flask at 425°C , what are the concentrations of H_2 , I_2 and HI when equilibrium has been achieved?

	H_2	$+ \text{I}_2$	\rightleftharpoons	2HI
I	$5.20 \times 10^{-3} \text{ M}$	$5.20 \times 10^{-3} \text{ M}$		0
C	$-x$	$-x$		$+ 2x$
E	$5.20 \times 10^{-3} - x$	$5.20 \times 10^{-3} - x$		$2x$

$$[\text{HI}] = 8.20 \times 10^{-3} \text{ M}$$

$$[\text{H}_2] = [\text{I}_2] = 1.10 \times 10^{-3} \text{ M}$$

$$K_c = \frac{[\text{HI}]^2}{[\text{H}_2][\text{I}_2]} = \frac{(2x)^2}{(5.20 \times 10^{-3} - x)(5.20 \times 10^{-3} - x)} = \left(\frac{2x}{5.20 \times 10^{-3} - x} \right)^2 = 55.64$$

$$\frac{2x}{5.20 \times 10^{-3} - x} = 7.459 \rightarrow 2x = 0.0388 - 7.459x \rightarrow 9.459x = 0.0388 \rightarrow x = 4.10 \times 10^{-3}$$

Equilibrium calculations: quadratic expression involving approximation

The reaction $\text{N}_2(\text{g}) + \text{O}_2(\text{g}) \rightleftharpoons 2 \text{NO}(\text{g})$ has $K_c = 1.0 \times 10^{-5}$ at 1500 K. Suppose a sample of air has $[\text{N}_2] = 0.080 \text{ M}$ and $[\text{O}_2] = 0.020 \text{ M}$ before any reaction occurs. Calculate the equilibrium concentration of reactants and products after the mixture has been heated to 1500 K.

	N_2	$+ \text{O}_2$	\rightleftharpoons	2NO
I	0.080 M	0.020 M		0
C	$-x$	$-x$		$+ 2x$
E	$0.080 - x$	$0.020 - x$		$2x$

$$K_c = \frac{[\text{NO}]^2}{[\text{N}_2][\text{O}_2]} = \frac{(2x)^2}{(0.080 - x)(0.020 - x)} = 1.0 \times 10^{-5}$$

$$4x^2 - (1.0 \times 10^{-5})x - 1.6 \times 10^{-8} = 0 \quad \text{Quadratic expression}$$

Equilibrium calculations: quadratic expression involving approximation

Use approximation to update the table and make your life easier
[N₂] and [O₂] don't significantly change during the reaction.

	N ₂	+ O ₂	⇌	2 NO
I	0.080 M	0.020 M		0
C	~ 0	- x		+ 2x
E	~0.080 M	~0.020 M		2x

$$K_c = \frac{[NO]^2}{[N_2][O_2]} = \frac{(2x)^2}{(0.080)(0.020)} = 1.0 \times 10^{-5}$$

$$[NO] = 1.3 \times 10^{-4} \text{ M}$$

$$x^2 = 4.0 \times 10^{-9} \rightarrow x = 6.3 \times 10^{-5}$$

Sample problem

Calculate K for the reaction



given the following information:

- $\text{SnO}_2 (\text{s}) + 2 \text{H}_2 (\text{g}) \rightleftharpoons \text{Sn} (\text{s}) + 2 \text{H}_2\text{O} (\text{g})$ $K = 8.12$
- $\text{H}_2 (\text{g}) + \text{CO}_2 (\text{g}) \rightleftharpoons \text{H}_2\text{O} (\text{g}) + \text{CO} (\text{g})$ $K = 0.771$
- ~~$\text{SnO}_2 (\text{s}) + 2 \text{H}_2 (\text{g}) \rightleftharpoons \text{Sn} (\text{s}) + 2 \text{H}_2\text{O} (\text{g})$~~ $K_1 = 8.12$
- ~~$2 \text{H}_2\text{O} (\text{g}) + 2 \text{CO} (\text{g}) \rightleftharpoons 2 \text{H}_2 (\text{g}) + 2 \text{CO}_2 (\text{g})$~~ $K_2 = \left(\frac{1}{0.771}\right)^2$
- $K_{\text{net}} = K_1 K_2 = (8.12) \left(\frac{1}{0.771}\right)^2 = 13.7$

Sample problem

N_2O_3 decomposes to NO and NO_2 is an endothermic process ($\Delta H^\circ = 40.5 \text{ kJ}$)



Predict the effect of the following changes on the equilibrium position.

- Adding more $\text{N}_2\text{O}_3 (\text{g}) \rightarrow$
- Adding more $\text{NO}_2 (\text{g}) \leftarrow$
- Increasing the volume of the reaction flask \rightarrow
 - System feels lower P (because V increased)
 - System needs to increase P by shifting to the side with more moles of gas
 - Reaction will shift to right (2 moles of gas on the right vs. 1 mole of gas on the left)
- Lowering the temperature \leftarrow
 - Reaction is endothermic so heat is a reactant
 - Lower T = less heat = less reactant