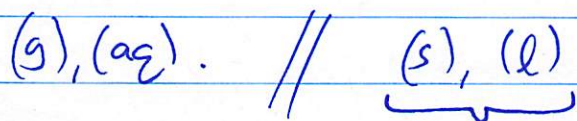


3/8/2019

What about solids + liquids?

K_c , molar concs (ignore 'em!)

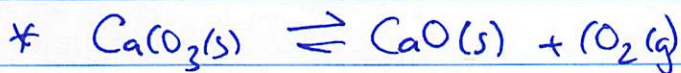
K_p , gas pressures (atm, ignore units).



effective conc/pressure = 1

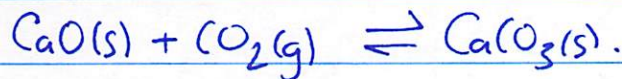


$$K_c = \frac{[CO_2][C]}{[CO]^2} = \frac{[CO_2]}{[CO]^2}$$

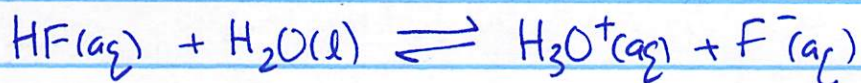


$$K_p = \frac{(P_{CaO})(P_{CO_2})}{(P_{CaCO_3})} = P_{CO_2}$$

reverse

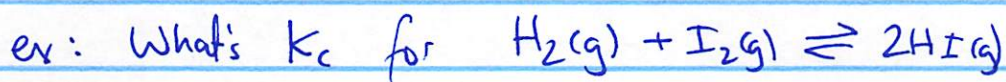


$$K_p = \frac{(P_{CaCO_3})}{(P_{CaO})(P_{CO_2})} = \frac{1}{P_{CO_2}}$$



$$K_c = \frac{[\text{H}_3\text{O}^+][\text{F}^-]}{[\text{HF}][\text{H}_2\text{O}]^?} = \frac{[\text{H}_3\text{O}^+][\text{F}^-]}{[\text{HF}]}$$

Calculating K_c from eqm concs



if $[\text{H}_2] = [\text{I}_2] = 0.11\text{M}$ and $[\text{HI}] = 0.78\text{M}$ @EQM

$$K_c = \frac{[\text{HI}]^2}{[\text{H}_2][\text{I}_2]} = \frac{(0.78)^2}{(0.11)(0.11)} = 50. \text{ or } 5.0 \times 10^1$$

What happens if we know INITIAL concs, but only know some of the final eqm concs? Can we find K_c ? (YES!)

-use stoichiometry + (R)ICE chart.

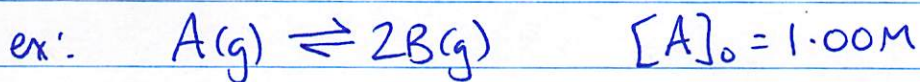
(Rxn) init' | | \ eqm
 change

Note: no matter the initial concentrations of H₂, I₂, or HI — at equilibrium the ratio of HI squared to H₂ × I₂ is always a constant (the equilibrium constant)

Typo! The squared should be outside of the []

TABLE 15.1 Initial and Equilibrium Concentrations for the Reaction
 $\text{H}_2(\text{g}) + \text{I}_2(\text{g}) \rightleftharpoons 2 \text{HI}(\text{g})$ at 445 °C

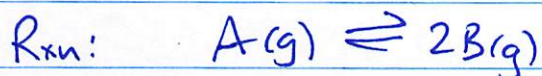
Initial Concentrations			Equilibrium Concentrations			Equilibrium Constant
[H ₂]	[I ₂]	[HI]	[H ₂]	[I ₂]	[HI]	$K_c = \frac{[\text{HI}]^2}{[\text{H}_2][\text{I}_2]}$
0.50	0.50	0.0	0.11	0.11	0.78	$\frac{(0.78)^2}{(0.11)(0.11)} = 50$
0.0	0.0	0.50	0.055	0.055	0.39	$\frac{(0.39)^2}{(0.055)(0.055)} = 50$
0.50	0.50	0.50	0.165	0.165	1.17	$\frac{(1.17)^2}{(0.165)(0.165)} = 50$
1.0	0.50	0.0	0.53	0.033	0.934	$\frac{(0.934)^2}{(0.53)(0.033)} = 50$
0.50	1.0	0.0	0.033	0.53	0.934	$\frac{(0.934)^2}{(0.033)(0.53)} = 50$



$[B]_0 = 0.00M$

$[A]_{eq} = 0.75M$

$K_c = ?$



Initial 1.00 0 +0.50

Change -0.25 $+ 2 \times 0.25$

Equilibrium 0.75 0.50

$K_c = \frac{[B]^2}{[A]_{eq}}$

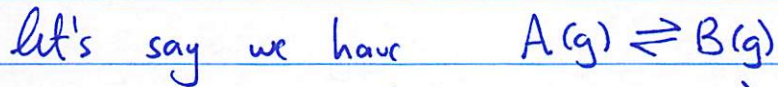
$= \frac{(0.50^2)}{(0.75)} = 0.33$

p 690

Ex: 15.5, 15.6

Do!

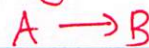
Predicting direction of change



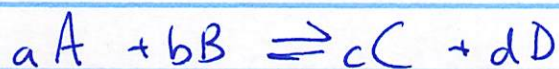
$K_c = 1.45$ (25°C)

$[A]_0 = [B]_0 = 1.00M$

Q: Which way will rxn proceed?



We need to calculate the
Reaction Quotient, Q



$$Q_c = \frac{[C]^c [D]^d}{[A]^a [B]^b}$$

↑ reaction quotient ← use current concs! (init)

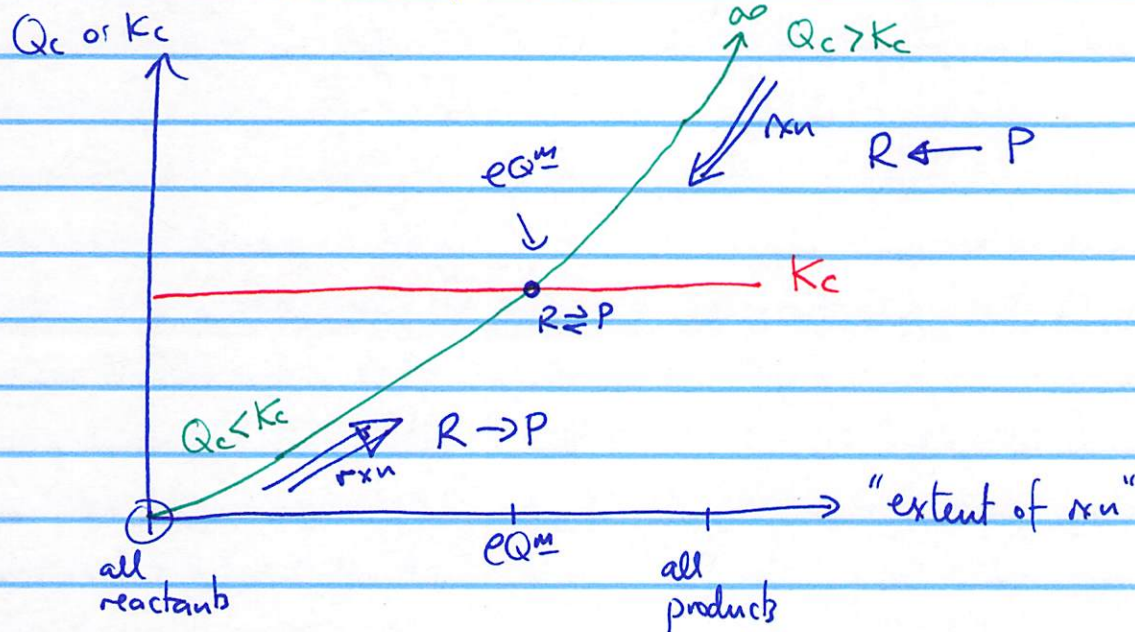
ex: @ beginning of rxn: $[C] = [D] = 0$, so $Q_c = 0$

→

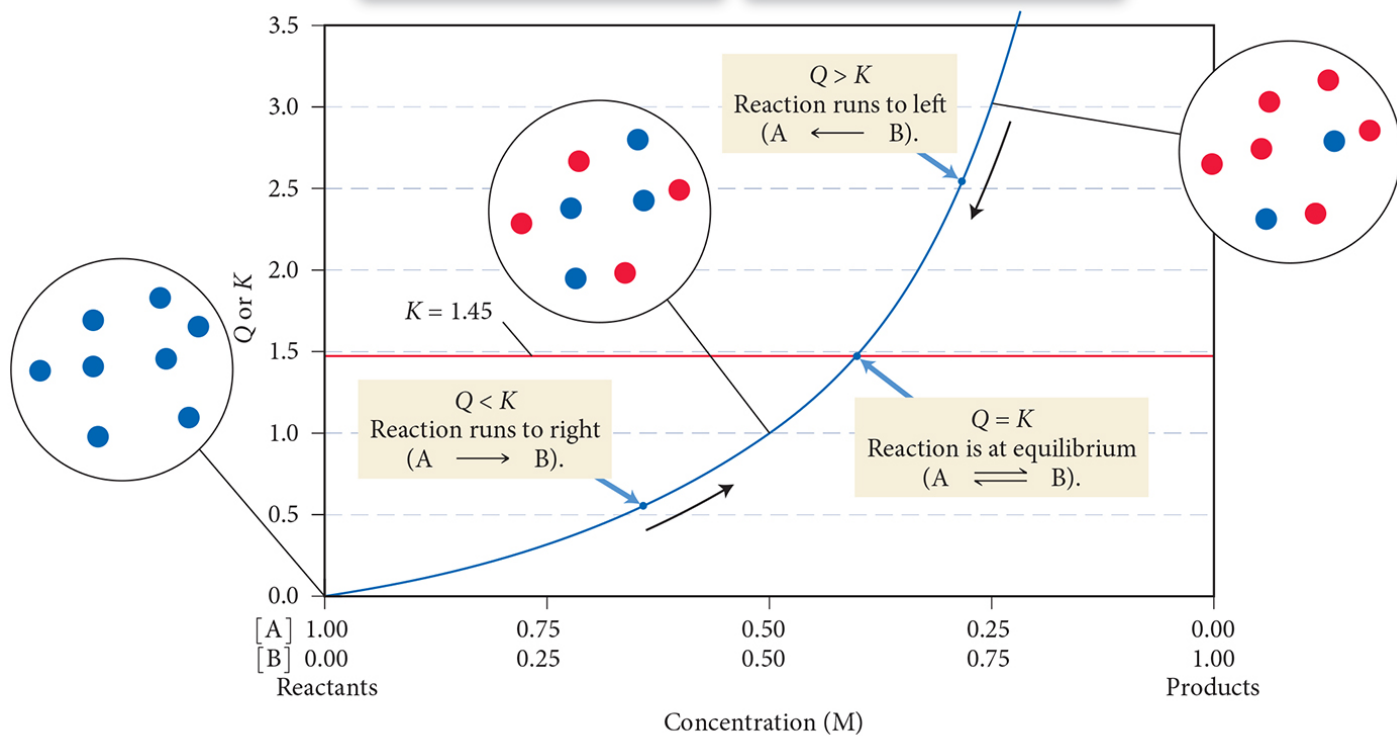
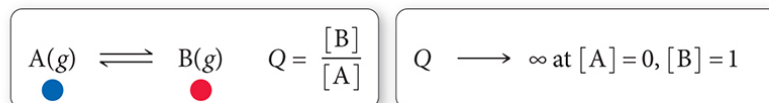
@ end of rxn: $[A] = [B] = 0$, so $Q_c = \infty$

←

@ eqm, $Q_c = K_c$



Q, K, and the Direction of a Reaction



our last situation: $A(g) \rightleftharpoons B(g)$; $K_c = 1.45$

$$[A]_0 = [B]_0 = 1.00M$$

$$Q_c = ?$$

Which way will it shift?

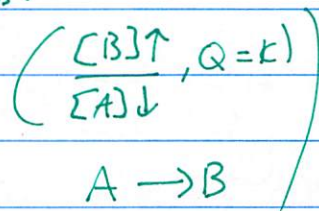
$$Q_c \neq K_c \text{ (not @ eqm)}$$

$$Q_c = \frac{[B]}{[A]} = \frac{1.00}{1.00} = 1.00 \quad Q_c < K_c: \text{Shift to RHS}$$

\leftarrow instantaneous/init values.

Yes, in theory you can memorize the direction of shift as follows:

- $Q < K$, shift to RHS
- $Q = K$, no shift (@ eqm)
- $Q > K$, shift to LHS



But if you understand that in order for the reaction to come to equilibrium the value of Q must increase or decrease to equalize K — and that only happens by causing more A to be converted to B or vice versa — then you will never forget on an exam!

Understanding takes a lot more time, energy, and effort than memorizing. Unfortunately that's the price for not forgetting. 😊