

Chemical Equilibria

Why do we care?!!

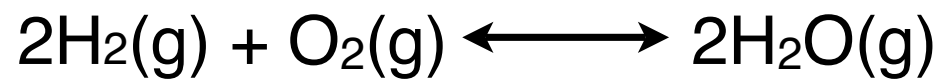
Put stuff in a beaker and what do you get?

We can use thermodynamics to predict the molecular concentrations at equilibrium (very powerful!)



$$\Delta_r G^\circ = +113.4 \text{ kJ mol}^{-1}$$

What about the opposite reaction?



$$\Delta_r G^\circ = -113.4 \text{ kJ mol}^{-1}$$

Does everything go to equilibrium as predicted?

What happens if you mix H_2 and O_2 at 298K?

- A. The explode and form water
- B. They explode and form hydrogen peroxide (H_2O_2)
- C. Nothing

Why didn't I get to equilibrium?

Kinetics

Other Reactions

Chemical Equilibria

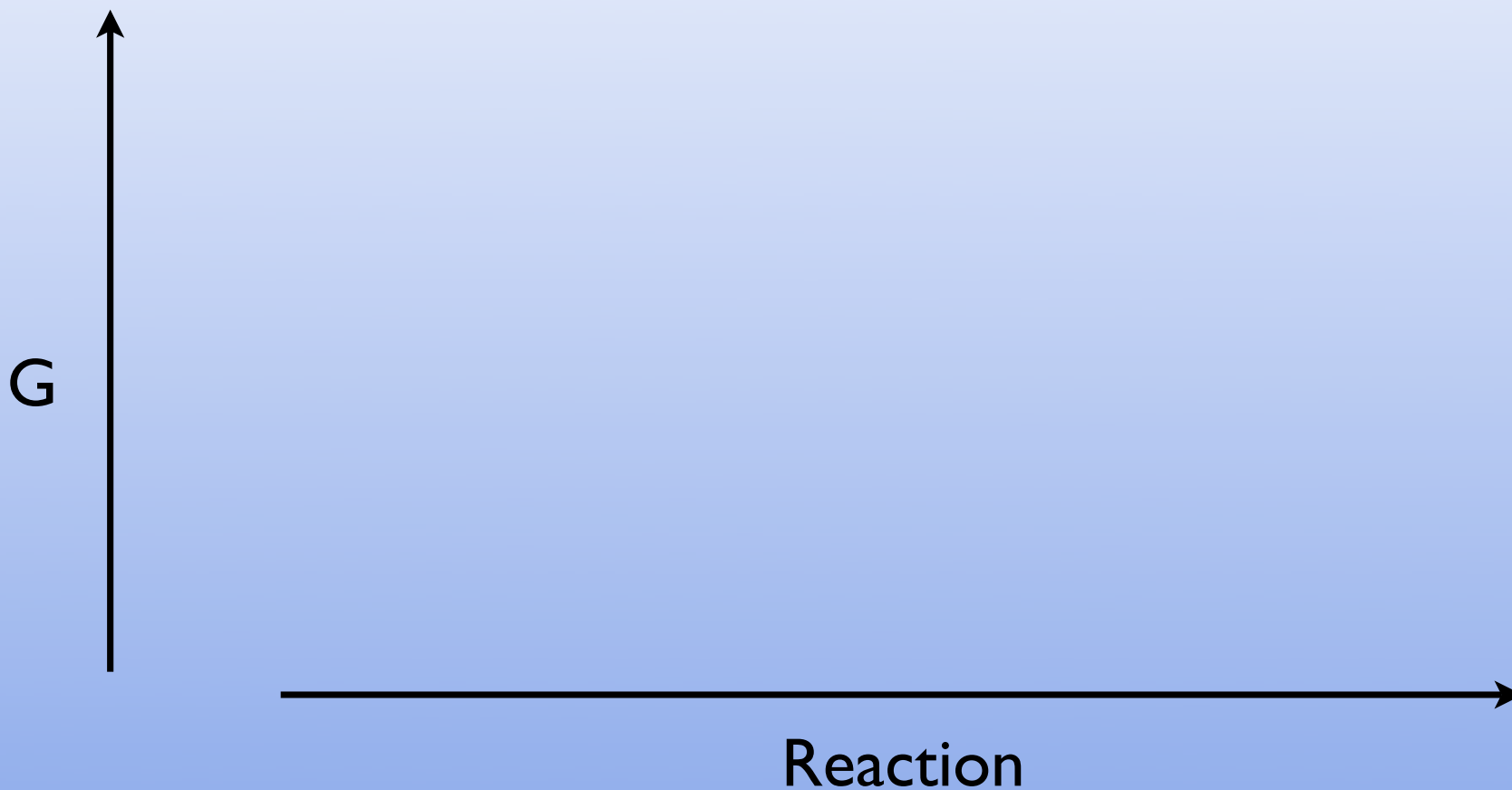
Why do we care?!!

Things might not get to equilibrium
but they never move away from it

Why does K depend on $\Delta_r G^\circ$?

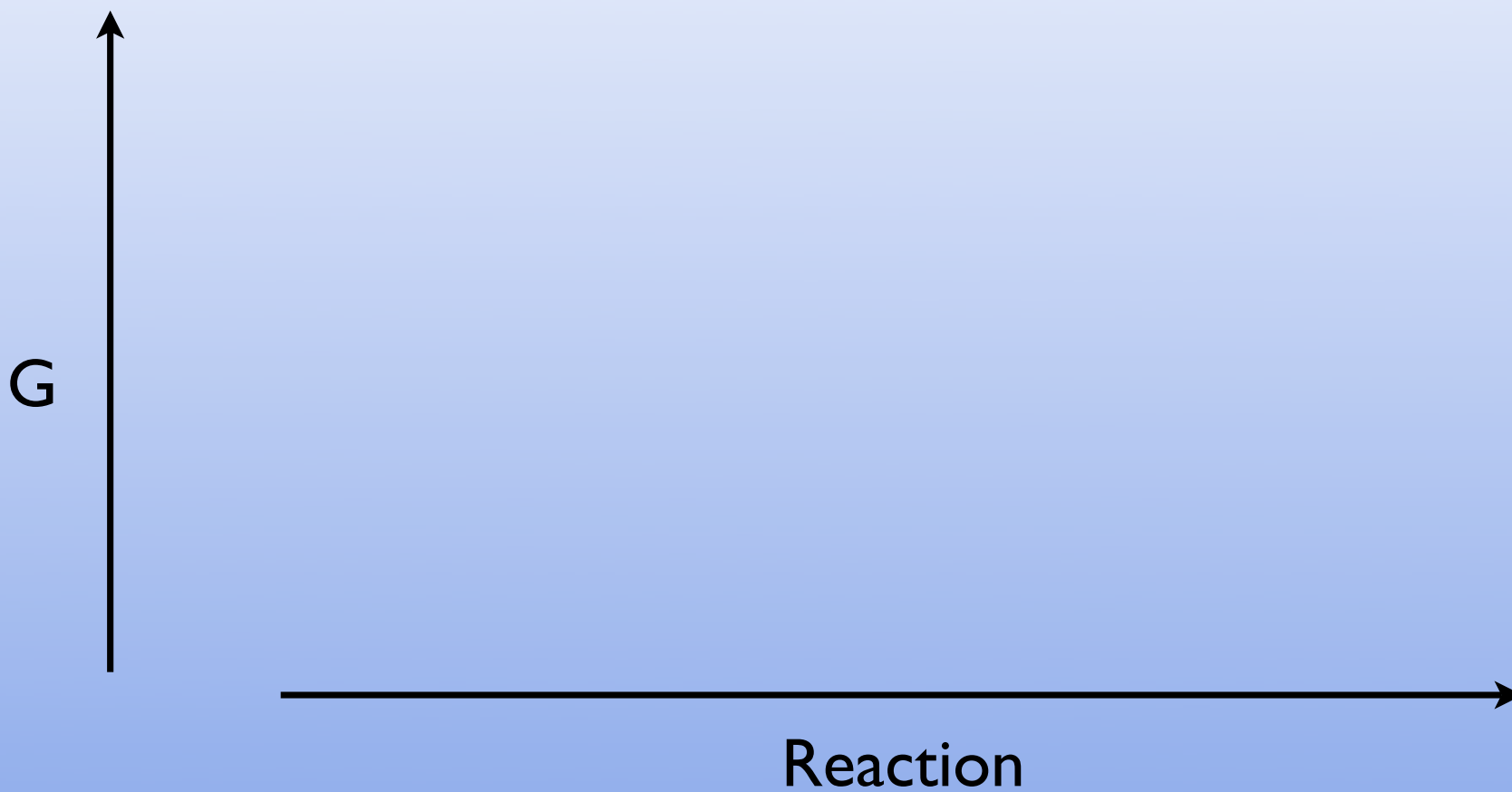


imagine
 $\Delta_r G^\circ = 0$



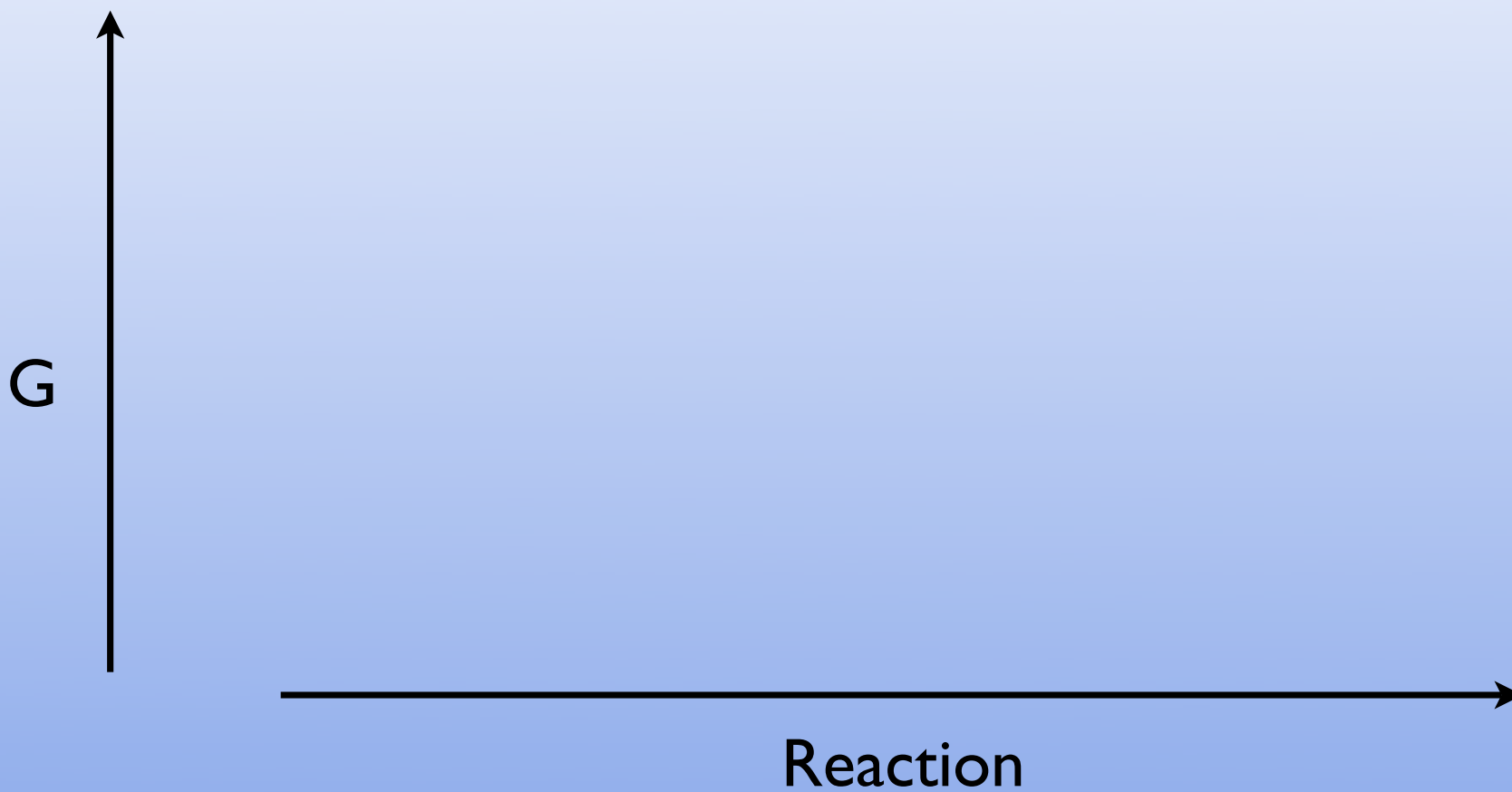


imagine
 $\Delta_r G^\circ = 0$





imagine
 $\Delta_r G^\circ > 0$



Equilibrium does not depend on starting conditions

TABLE 6.1 Results of Three Experiments for the Reaction $\text{N}_2(g) + 3\text{H}_2(g) \rightleftharpoons 2\text{NH}_3(g)$

Experiment	Initial Concentrations	Equilibrium Concentrations	$K = \frac{[\text{NH}_3]^2}{[\text{N}_2][\text{H}_2]^3}$
I	$[\text{N}_2]_0 = 1.000 \text{ M}$ $[\text{H}_2]_0 = 1.000 \text{ M}$ $[\text{NH}_3]_0 = 0$	$[\text{N}_2] = 0.921 \text{ M}$ $[\text{H}_2] = 0.763 \text{ M}$ $[\text{NH}_3] = 0.157 \text{ M}$	$K = 6.02 \times 10^{-2} \text{ L}^2/\text{mol}^2$
II	$[\text{N}_2]_0 = 0$ $[\text{H}_2]_0 = 0$ $[\text{NH}_3]_0 = 1.000 \text{ M}$	$[\text{N}_2] = 0.399 \text{ M}$ $[\text{H}_2] = 1.197 \text{ M}$ $[\text{NH}_3] = 0.203 \text{ M}$	$K = 6.02 \times 10^{-2} \text{ L}^2/\text{mol}^2$
III	$[\text{N}_2]_0 = 2.00 \text{ M}$ $[\text{H}_2]_0 = 1.00 \text{ M}$ $[\text{NH}_3]_0 = 3.00 \text{ M}$	$[\text{N}_2] = 2.59 \text{ M}$ $[\text{H}_2] = 2.77 \text{ M}$ $[\text{NH}_3] = 1.82 \text{ M}$	$K = 6.02 \times 10^{-2} \text{ L}^2/\text{mol}^2$

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Each equilibrium has different concentrations,
but the same value for K_c

A convention to keep things straight

we'll be doing a lot of aqueous problems

C denote concentrations initially

[] denote concentrations at equilibrium

Reaction	$3\text{H}_2(\text{g}) + \text{N}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$
Initial	
Change	
Equilibrium	

Really Easy problems

At equilibrium you find

$$[\text{H}_2] = .1 \text{ M}, [\text{N}_2] = 0.2 \text{ M}, \text{ and } [\text{NH}_3] = .2\text{M}$$

K =

Reaction	$3\text{H}_2(\text{g}) + \text{N}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$
Initial	
Change	
Equilibrium	

Fairly Easy problem

Given $K = 200$ and
 $[H_2] = .2 \text{ M}$, $[N_2] = 0.4 \text{ M}$, and $C_{NH_3} = .1 \text{ M}$
fill in the rest

Reaction	$3H_2(g) + N_2(g)$	\longleftrightarrow	$2NH_3(g)$
Initial			
Change			
Equilibrium			

Typical problem

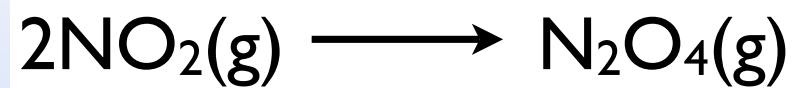
Given $K = 200$ and
 $C_{H_2} = .2 \text{ M}$, $N_2 = 0.2 \text{ M}$ what are the
concentrations at equilibrium

Reaction	$3H_2(g) + N_2(g)$	\longleftrightarrow	$2NH_3(g)$
Initial			
Change			
Equilibrium			

Reaction	$3\text{H}_2(\text{g}) + \text{N}_2(\text{g}) \rightleftharpoons 2\text{NH}_3(\text{g})$		
Initial	.2	.2	0
Change	-3x	-x	+2x
Equilibrium	.2-3x	.2-x	+2x

I thought you said we need to use
 K_p for gases
and
 K_c for solutions?

Relating K_p and K_c



$$K_c = \text{_____}$$

$$K_p = \text{_____}$$

$$P_{\text{N}_2\text{O}_4} = \text{_____}$$

Relating K_p and K_c



$$K_p = \frac{P_{\text{N}_2\text{O}_4}}{P_{\text{NO}_2}^2} = \frac{[\text{N}_2\text{O}_4]RT}{[\text{NO}_2]^2(RT)^2} =$$

In general $K_p = K_c(RT)^{\Delta n}$

Δn is the change in the number of moles of gas

Time out for activities

That is what we are actually putting
into the equilibrium constant

Gas

$$a_i =$$

Free Energy
Changes with P

Compound
in solution

$$a_i =$$

Free Energy
Changes with
Concentration

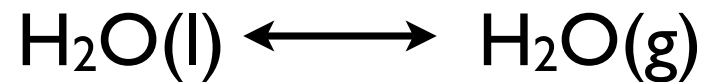
What about the activity of a pure liquid or solid?

The pure compound is the reference state!

$a_i =$

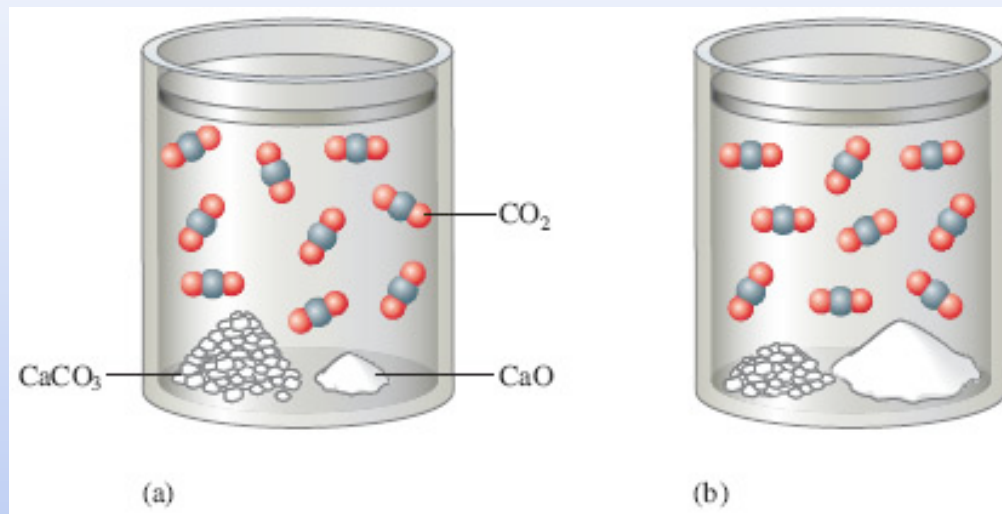
Pure solids and liquids “don’t show up”
in the equilibrium constant
(they are there. they are just always = 1)

What is the equilibrium constant for this reaction?



K =

Equilibria with more than one phase are called Heterogeneous Equilibria



K=

For the following reaction $\Delta_{\text{R}}G^{\circ} = +740 \text{ kJ mol}^{-1}$ at 298K
In air will I form any solid iron?



- A. all the iron oxide will convert to iron
- B. about half of the iron oxide will convert to iron
- C. a very small amount of the iron oxide will convert to iron
- D. not a single atom of iron will form

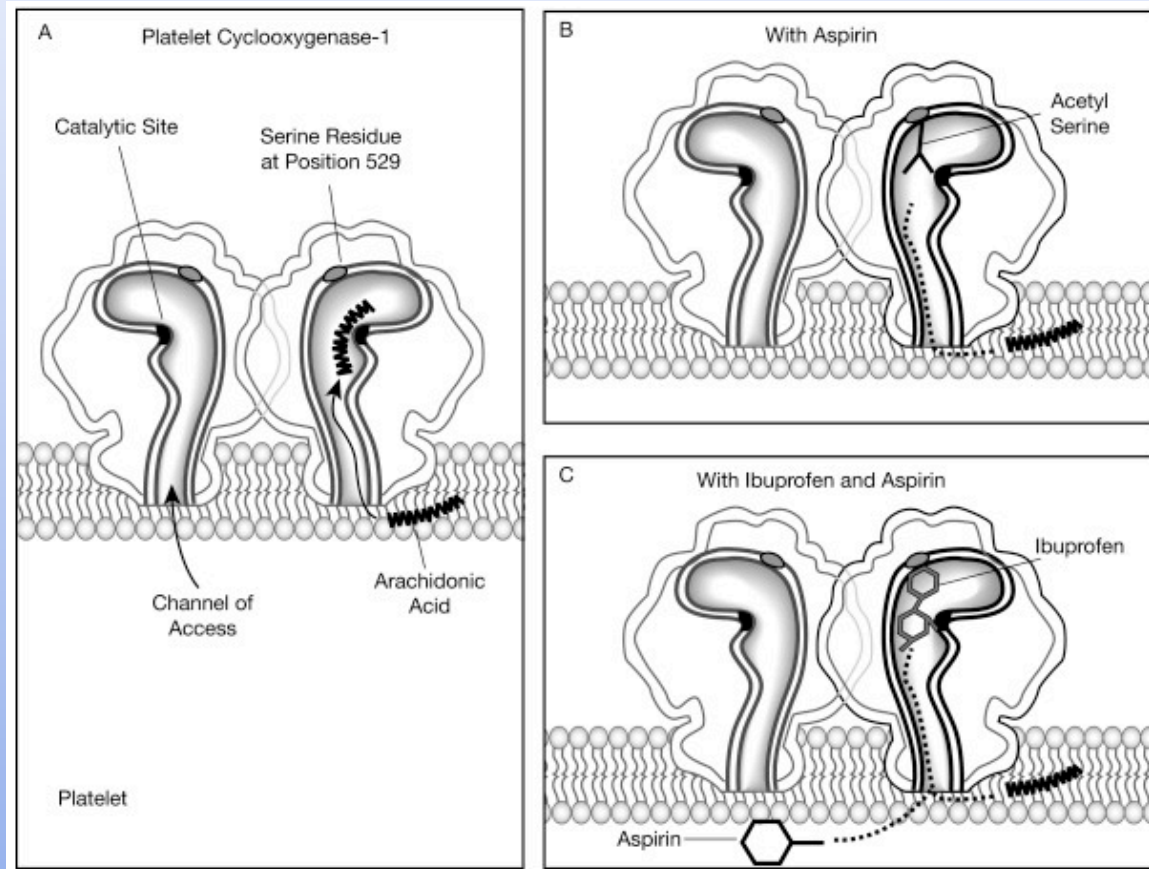
For the following reaction $\Delta_{\text{R}}G^{\circ} = +740 \text{ kJ mol}^{-1}$ at 298K
In air will I form any solid iron?



$$K = \exp[-740,000 / (8.314)(298)] = 2 \times 10^{-130}$$

What is Q?

What is equilibrium good for?



Fendrick *et al.* *Osteopathic Medicine and Primary Care* 2008 **2**:2 doi:10.1186/1750-4732-2-2

Drug Binding

How “strongly” should it bind to work?



$K =$

Do an experiment
Measure K



For **ibuprofen** binding to the **COX channel**

$$K \sim 10^8$$

If we want 100x more complexed protein than free protein
what concentration of drug do we need?

A. 10^{-8} M

B. 10^{-6} M

C. 10^{-4} M

D. 10^{-2} M

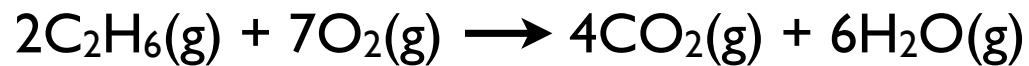
$$K = \frac{[\text{complex}]}{[\text{drug}][\text{protein}]}$$

How much is that?

Person is 50 kg = 50 L of water

MW of ibuprofen is $\sim 200 \text{ g mol}^{-1}$

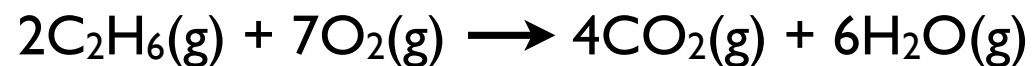
For the following reaction what is the change value for H₂O?



R	C ₂ H ₆	O ₂	CO ₂	H ₂ O
I	1.0	1.4	1.8	0
C	-2x	?	?	?

- A. -2x
- B. +2x
- C. +3x
- D. +6x

For the following reaction what is the equilibrium value for CO₂?



R	C ₂ H ₆	O ₂	CO ₂	H ₂ O
I	1.0	1.4	1.8	0
C	-2x	?	?	?

- A. 1.8 - 2x
- B. 1.8 + 2x
- C. 1.8 + 4x
- D. 1.0 + 6x