

# C12 - 2.0 - Reaction Rates/Graphs

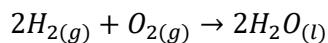
$$\boxed{\text{Rate} = \frac{\text{amount}}{\text{time}}}$$

Find rate if 8g used up in 4 minutes.

$$\text{Rate} = \frac{8\text{ g}}{4\text{ min}}$$

$$\boxed{\text{Rate} = 2\frac{\text{ g}}{\text{ min}}}$$

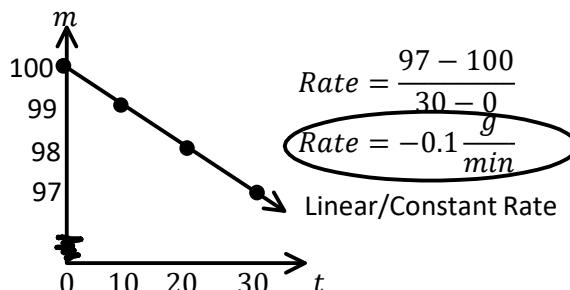
If rate of water production is  $0.5 \frac{\text{ mol }}{\text{ min }}$ , find the rate of oxygen gas consumption.



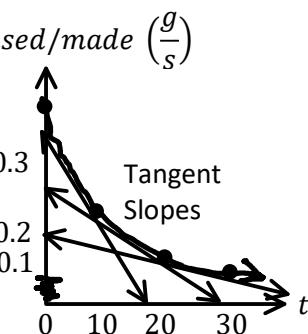
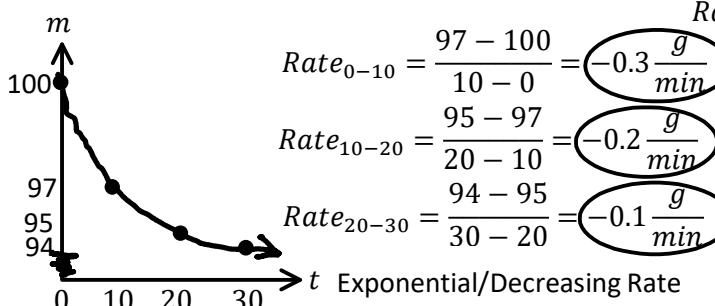
$$0.5 \frac{\text{ mol } H_2O}{\text{ min }} \times \frac{1 \text{ mol } O}{2 \text{ mol } H_2O} = \boxed{0.25 \frac{\text{ mol } O_2}{\text{ min }}} \quad (\text{Coefficients}^*)$$

Rates :  
 Moles Reactants used up =  
 Moles Products are made  
 (Coefficients\*)

Time(s)	Mass(g)
0	100
10	99
20	98
30	97

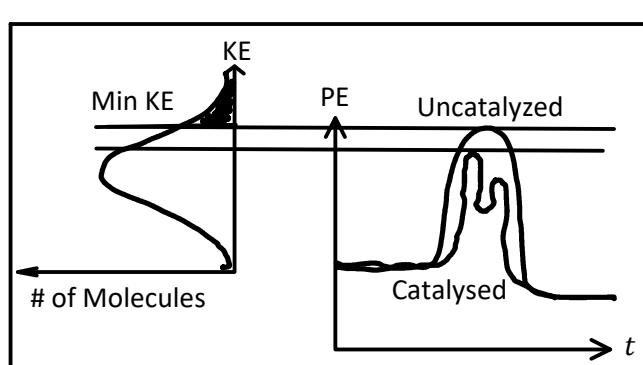
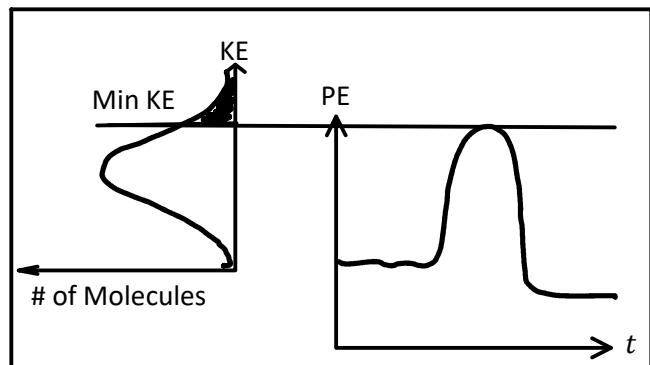
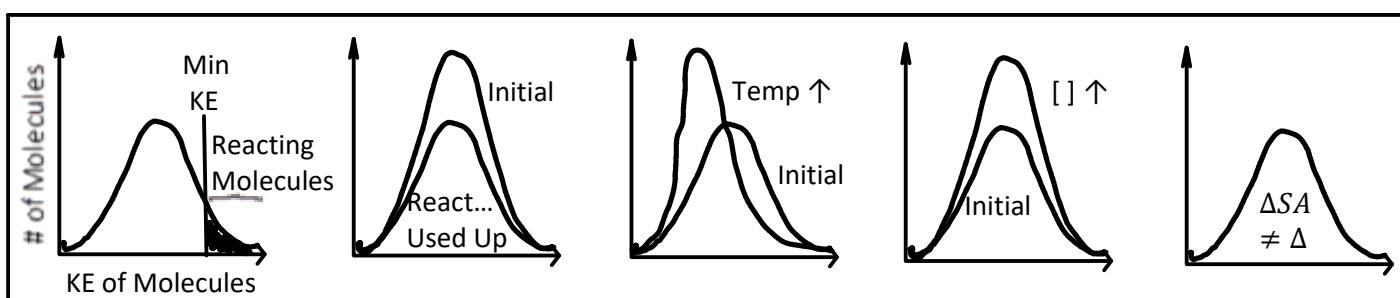


Time(s)	Mass(g)
0	100
10	97
20	95
30	94



Faster  
 $Li^+ + F^- \rightarrow LiF$   
 -1 bonds made

Slower  
 $H - H + O = O \rightarrow 2H - O - H$   
 -2 bonds broken, 4 bonds made



# C12 - 2.0 - Equilibrium (Eq)

Don't have to calculate the rates\*

Equilibrium -  $Rate_{forward} ; K_f = K_r ; Rate_{reverse}$ . (Microscopic changes\*)

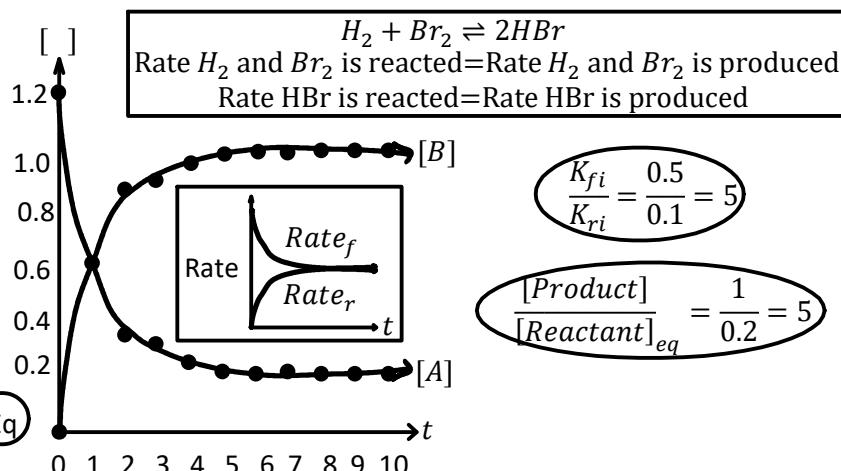
**K : Rate constant**

$K_f \propto [Reactant]$   
 $K_r \propto [Product]$

Rate of consumption of reactants = rate of production of reactants. (A system will tend towards Eq.)  
[Reactants] & [Products] constant in time. [Reactants] differ from [Products] in general.



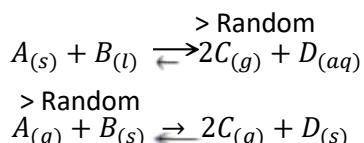
Time (min)	$R_f$	$R_r$	[A]	[B]
0	0.600	0.000	1.200	0.000
1	0.300	0.060	0.600	0.600
2	0.180	0.084	0.360	0.840
3	0.132	0.094	0.264	0.936
4	0.113	0.097	0.226	0.974
5	0.105	0.099	0.210	0.990
6	0.102	0.100	0.204	0.996
7	0.101	0.100	0.202	0.998
8	0.100	0.100	0.201	0.999
9	0.100	0.100	0.200	1.000
10	0.100	0.100	0.200	1.000



Spontaneous Change- A change that occurs by itself, without outside assistance. (Exothermic\*)

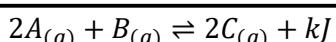
Entropy - Amount of randomness in a system.  $gases \gg solutions \gg liquids \gg solids$

Tendency for a reaction to go to the side of Maximum Entropy (Most particles of most random phase)/Minimum Energy (Enthalpy).



Le Chatelier's Principle - If a closed system (Nothing can enter or leave) at equilibrium is subject to a change, processes will occur that tend to counteract that change. (Whatever we do, nature tends to undo\*.)

## Effects



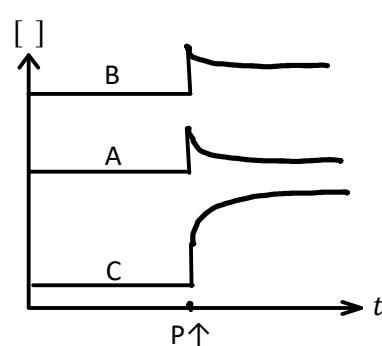
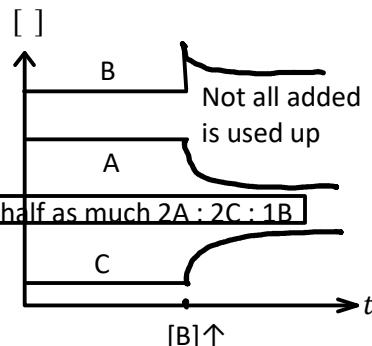
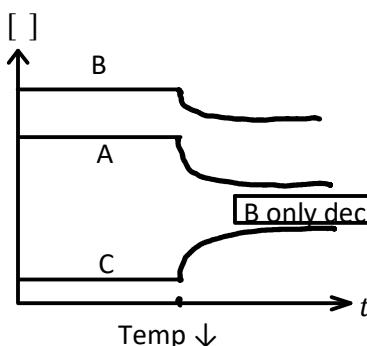
$\uparrow [B]$  Shifts to use up more [A].

Initially :  $\uparrow r_f, \downarrow r_r$ , Then :  $\downarrow r_f, \uparrow r_r$

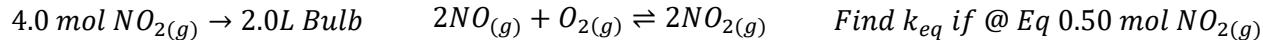


$\downarrow V, \uparrow P, \uparrow [\text{gases}]$   
Shifts to lower overall pressure.  
(Side with lower moles.)

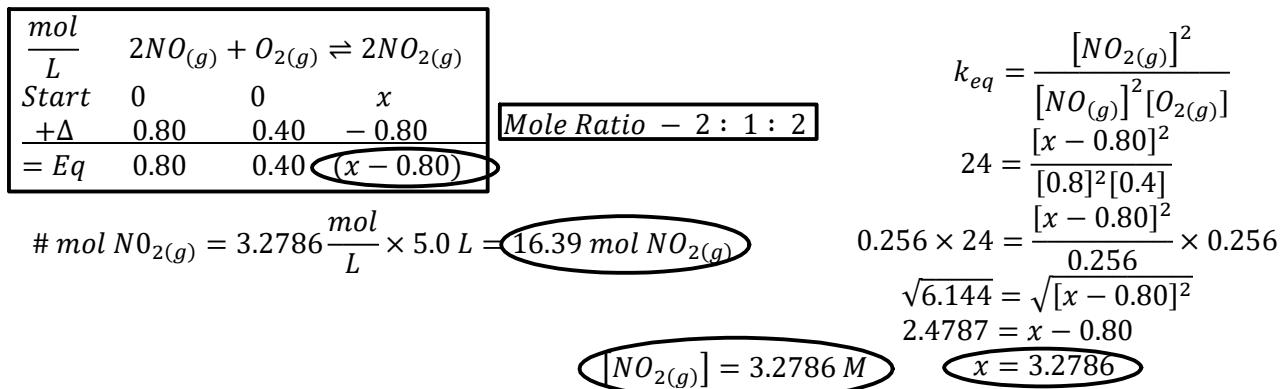
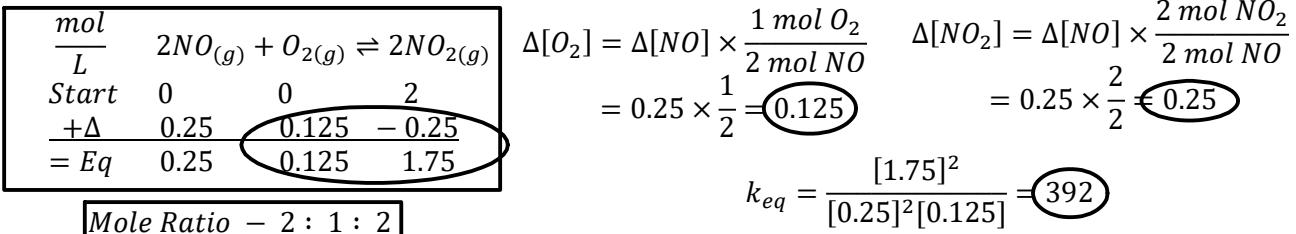
$\downarrow \text{heat.}$  Shifts to produce  $A \rightleftharpoons B + kJ$  more heat.



## C12 - 2.0 - Equilibrium



$$k_{eq} = \frac{[NO_{2(g)}]^2}{[NO_{(g)}]^2 [O_{2(g)}]} \quad [NO_{2(g)}]_{start} = \frac{4.0 \text{ mol}}{2.0 \text{ L}} = 2.0 \text{ M} \quad [NO_{(g)}]_{eq} = \frac{0.5 \text{ mol}}{2.0 \text{ L}} = 0.25 \text{ M}$$



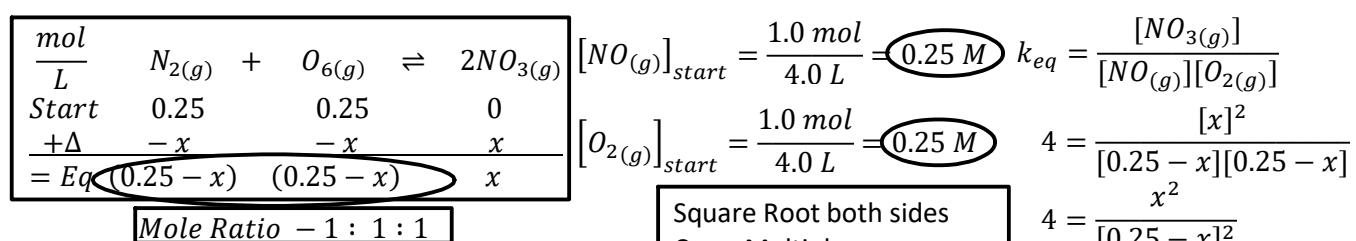
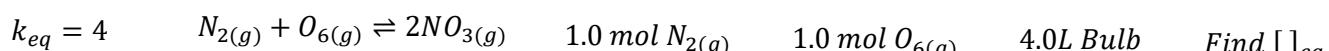
Which direction will  
the reaction shift?

$Q = \text{Reaction Quotient}$

$$Q = \frac{[NO_{2(g)}]^2}{[NO_{(g)}]^2 [O_{2(g)}]} \quad k_{eq} = \frac{[NO_{2(g)}]^2}{[NO_{(g)}]^2 [O_{2(g)}]}$$

$$[NO_{(g)}]_{start} = \frac{2.0 \text{ mol}}{2.0 \text{ L}} = 1.0 \text{ M} \quad [O_{2(g)}]_{start} = \frac{0.2 \text{ mol}}{2.0 \text{ L}} = 0.1 \text{ M} \quad [NO_{2(g)}]_{start} = \frac{0.4 \text{ mol}}{2.0 \text{ L}} = 0.2 \text{ M}$$

$$Q = \frac{[0.2]^2}{[1.0]^2 [0.1]} = 0.4 = \frac{[\text{Products}]}{[\text{Reactants}]} < k_{eq} = 49 \quad \text{Shift towards products}$$



$$[NO_3(g)] = 0.17 \text{ M}$$

$$[N_2(g)] = [O_6(g)] = (0.25 - x) = (0.25 - 0.167) = 0.08$$

$$2(0.25 - x) = x \\ 0.5 - 2x = x \\ 0.5 = 3x \\ x = 0.167$$