General Chemistry II Chapter 14 Study Guide

Types of reaction rates:

- If the concentration of a reactant is plotted against time:
 - If the graph has a curved line = rate changes, will reach equilibrium
 - If the graph is linear = rate is constant; will run out of reactant
- Average rate of reaction: change in concentration of a reactant or product over time; reaction rate = slope
- instantaneous rate: slope of a line tangent to the curve; can pick any time
- Initial rate: rate within t= 0 to t= given time; generally portion of the curve that is linear

Rate laws:

- Rate laws: only reactants appear in the rate law; the exponents tells the order of the rxn with respect to each reactant → sum of exponents is the overall order of the rxn; once we know the order of each reactant, we can plug in the concentrations and reaction rate to calculate the rate constant, k
- rate = k[A]"[B]"

Determining reaction orders:

- Suppose A and B are two reactants in a reaction:
 - If double [A] will cause no change to rate, the order on A is 0 (m=0), e.g. [2]0 = 1.1410 - 1
 - If double [A] will cause the rate to double, the order on A is 1 (m=1), e.g. [2]1 = 2, [4]1 - 4
 - If double [A] will cause the rate to quadruple, the order on A is 2 (m=1), e.g. [2]2. -4, [4]2 - 16
 - The order on reactant is not related to its coefficient in the chemical equation.
 - the exponents must be determined experimentally.

Determining rate laws; isolation method

- For rxns with more than one reactant, the rate law can be determined by the isolation method; approach:
 - Fix the concentration of all reactants except for one
 → determine the rxn rates with varied concentrations of that reactant
 - 2. The change in rate is only attributed to that one reactant, enabling determination of the order of that reactant in the rate law
 - Apply similar method to determine the order of other reactants

Zero-order process: overall order = 0:

Zero-Order Process: overall order = 0 For reaction A→ Product: Units of k: k = rate = M/s. Rate = $k[A]^2 = k$ 104

d[A] = -kdtTake the integral:

= $[A]_t = -k\varepsilon + C$ Plug in initial condition (A) =(A), at t=0: G = fAL

Plot of IAL vs. f. 3

 $[A]_t = -kt + [A]_\alpha$ Equivalent to a straight-line function:

y = mx + b

- Plot of [A], vs. t is a straight line with k as slope.
- b is the y-intercept (i.e. the initial concentration [AL)

Zero-Order Process

Eventually [A] will go to zero (all reactants will be used up)

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T. W. Graham Solomons, Craig B. Fryhle, Scott A. Snyder

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