

During Class Invention

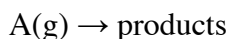
Name _____

Integrated Rate Law Part I

TA Name _____

Lab Section # _____

1. The reaction:



follows simple first order kinetics. When the initial concentration of A is 0.500 M the initial rate of the reaction is determined to be $4.20 \times 10^{-3} \text{ M s}^{-1}$. If the initial concentration of A is tripled, what would be the new initial rate of the reaction?

The rate law for the reaction that follows first order kinetics is

$$\text{Rate} = k[A]^1$$

If [A] is equal to 3[A] then substitute into the rate law,

$$\text{Rate} = k[3[A]]^1$$

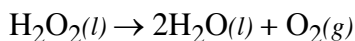
The initial rate will triple.

$$\text{Rate} = 3 \cdot 4.20 \times 10^{-3} \text{ M s}^{-1} = 1.26 \times 10^{-2} \text{ M s}^{-1}$$

2. Write the integrated rate law for a reaction that follows simple first order kinetics.

$$\ln \left(\frac{[A]}{[A]_0} \right) = -kt$$

3. The decomposition of H_2O_2 to H_2O and O_2 follows first order kinetics with a rate constant of 0.0410 min^{-1} at a particular temperature.



Calculate the $[\text{H}_2\text{O}_2]$ after 10 mins if $[\text{H}_2\text{O}_2]_0$ is 0.200 M.

$$\ln \frac{[\text{H}_2\text{O}_2]}{[\text{H}_2\text{O}_2]_0} = -kt$$

$$\ln \frac{[\text{H}_2\text{O}_2]}{0.200 \text{ M}} = -(0.0410 \text{ min}^{-1})(10 \text{ mins})$$

$$\ln \frac{[\text{H}_2\text{O}_2]}{0.200 \text{ M}} = -0.410$$

$$e^{\left(\ln \frac{[\text{H}_2\text{O}_2]}{0.200 \text{ M}} \right)} = e^{-0.410}$$

$$\frac{[\text{H}_2\text{O}_2]}{0.200 \text{ M}} = 0.6637$$

$$[\text{H}_2\text{O}_2] = 0.6637 \cdot (0.200 \text{ M})$$

$$[\text{H}_2\text{O}_2] = 0.133 \text{ M}$$

4. The decomposition of N_2O_5 to O_2 and NO_2 follows first order kinetics. If a sample at 25°C with the initial concentration of N_2O_5 of $1.25 \times 10^{-3} \text{ M}$ falls to $1.02 \times 10^{-3} \text{ M}$ in 100. minutes, calculate the rate constant for the reaction.

$$\ln \frac{[\text{N}_2\text{O}_5]}{[\text{N}_2\text{O}_5]_0} = -kt$$

$$\ln \frac{[1.02 \times 10^{-3} \text{ M}]}{[1.25 \times 10^{-3} \text{ M}]_0} = -k(100 \text{ min})$$

$$\ln 0.816 = -k(100 \text{ min})$$

$$-0.203 = -k(100 \text{ min})$$

$$\frac{-0.203}{100 \text{ min}} = -k$$

$$2.03 \times 10^{-3} \text{ min}^{-1} = k$$

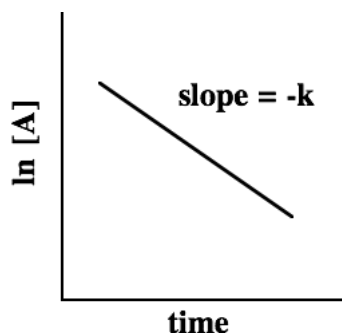
5. Show how a plot of $\ln[\text{concentration}]$ versus time can provide the rate constant for a reaction which follows simple first order kinetics.

$$\ln \left(\frac{[\text{A}]}{[\text{A}]_0} \right) = -kt$$

$$\ln[\text{A}] - \ln[\text{A}]_0 = -kt$$

$$\ln[\text{A}] = -kt + \ln[\text{A}]_0$$

this equation fits the general equation for a line, $y = mx + b$, where
 $y = \ln[\text{A}]$: $m = -k$: $x = t$



6. Using the following data, establish that the decomposition N_2O_5 according to the reaction,



follows first order kinetics. Determine the rate constant for the reaction.

Time (sec)	$[\text{N}_2\text{O}_5] \text{ (M)}$	$\ln[\text{NO}_2]$
0	1.50×10^{-3}	-6.50
2000	1.40×10^{-3}	-6.57
5000	1.27×10^{-3}	-6.67
7000	1.18×10^{-3}	-6.74
11000	1.03×10^{-3}	-6.88
15000	9.00×10^{-4}	-7.01

