

C. 27183
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Quiz 1 (Chapter 12)

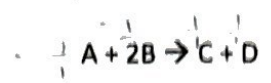
Kelsey

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	Zero-Order	First-Order	Second-Order
rate law	rate = k	rate = k[A]	rate = k[A] ²
units of rate constant	M s ⁻¹	s ⁻¹	M ⁻¹ s ⁻¹
integrated rate law	[A] = -kt + [A] ₀	ln[A] = -kt + ln[A] ₀	$\frac{1}{[A]} = kt + \left(\frac{1}{[A]_0}\right)$

Arrhenius Equation: $k = Ae^{-E_a/RT}$

1. Consider the following reaction in aqueous solution:



a) Write the equation that relates the rate expressions for this reaction in terms of the disappearance of A and the disappearance of B.

$$-\frac{1}{1} \frac{\Delta[A]}{\Delta t} = -\frac{1}{2} \frac{\Delta[B]}{\Delta t} = \frac{1}{1} \frac{\Delta[C]}{\Delta t} = \frac{1}{1} \frac{\Delta[D]}{\Delta t}$$

Not part of the Q.

b) If the rate of disappearance of A at a particular moment during the reaction is 1.4×10^{-4} mol L⁻¹ s⁻¹, what is the rate of disappearance of B at that moment?

$$(1.4 \times 10^{-4} \frac{\text{mol}}{\text{L}\cdot\text{s}}) \times -2 = \boxed{0.0002} \text{ unit.}$$

$$\begin{aligned} -\frac{1}{1} \frac{\Delta[A]}{\Delta t} &= -\frac{1}{2} \frac{\Delta[B]}{\Delta t} \\ \times 2 & \quad \times 2 \\ -2 \frac{\Delta[A]}{\Delta t} &= \frac{\Delta[B]}{\Delta t} \end{aligned}$$

2. The following data have been determined for the reaction:



	[NO] initial (M)	[Br ₂] initial (M)	Rate (mol L ⁻¹ s ⁻¹)
1	0.02 ^{x2}	0.02	9.6 × 10 ⁻²
2	0.04	0.02 ^{x2}	3.8 × 10 ⁻¹ ^{x2}
3	0.02	0.04 ^{x2}	1.9 × 10 ⁻¹

Determine 1) the rate law and 2) the rate constant for this reaction.

[NO]

$$\frac{\text{rate 1}}{\text{rate 2}} = \frac{k[\text{NO}]^m + [\text{Br}_2]^n}{k[\text{NO}]^n + [\text{Br}_2]^n}$$

$$\frac{9.6 \times 10^{-2}}{3.8 \times 10^{-1}} = \left(\frac{0.02}{0.04}\right)^m \rightarrow \ln(0.2526) = m \ln(0.5) \rightarrow m = \ln\left(\frac{0.2526}{0.5}\right)$$

$$m = -0.683$$

1

2.

[Br₂]

Between rate 2 & 3, the initial (M) and the rate are both doubled meaning it is the first order. [Br₂]¹

✓

1).

$$\left(\text{rate} = k[2\text{NO}]^m + [\text{Br}_2]^n \right)$$

$$3.8 \times 10^{-1} \frac{\text{mol}}{\text{L s}} = k[0.04\text{M}]^m + [0.02\text{M}]^n$$

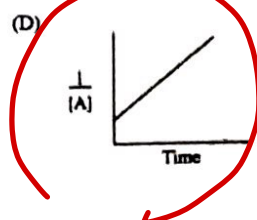
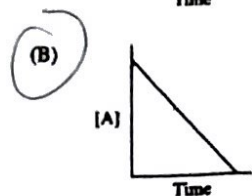
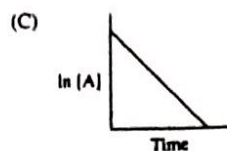
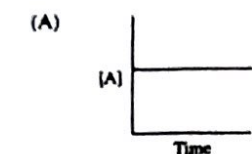
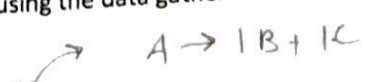
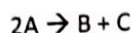
$$3.8 \times 10^{-1} \frac{\text{mol}}{\text{L s}} = k(0.06\text{M})$$

ok.

$$\left(k = 6.33 \frac{1}{\text{L s}} \right)$$

✱ Box your answer.

3. Which of the following graphs may have been created using the data gathered from the following reaction? Assume this is a single step reaction:



-2

4. Dinitrogen pentoxide gas decomposes according to the equation: $2 \text{N}_2\text{O}_5(\text{g}) \rightarrow 4 \text{NO}_2(\text{g}) + \text{O}_2(\text{g})$. The first-order reaction was allowed to proceed at 40.0°C . The initial concentration of N_2O_5 was 0.400 M and after 20.0 minutes, the concentration changed to 0.289 M .

$$0.400 - 0.289 = 0.111$$

$$20.0 \text{ min} \times \frac{60 \text{ sec}}{1 \text{ min}} = 1200$$

- (a) Calculate the rate constant for the reaction. $k = ?$

$$k t = \frac{\ln[A]_0}{\ln[A]} \rightarrow k t = \ln\left(\frac{0.400 \text{ M}}{0.289 \text{ M}}\right) \rightarrow k t = 1.2819$$

-1

$$k = \frac{1.2819}{20.0 \text{ min}}$$

$$k = 0.0641 \frac{1}{\text{min}} \times \frac{1 \text{ min}}{60 \text{ sec}} = 1.07 \times 10^{-3} \frac{1}{\text{sec}}$$

- (b) After how many minutes will $[\text{N}_2\text{O}_5]$ be equal to 0.350 M ?

$$k = 0.0641 \frac{1}{\text{min}}$$

$$t = ?$$

$$[A] = 0.350 \text{ M}$$

$$[A]_0 = 0.400 \text{ M}$$

$$0.400 \text{ M} - 0.350 \text{ M} = 0.05$$

$$k t = \ln\left(\frac{0.400 \text{ M}}{0.350 \text{ M}}\right)$$

$$(0.0641 \frac{1}{\text{min}})(t) = 2.079$$

$$t = \frac{2.079}{0.0641 \frac{1}{\text{min}}}$$

ok.

$$t = 32.4 \frac{1}{\text{min}}$$

5. The rate constant at 550 °C for the decomposition reaction $2\text{H}_2\text{O}_2 \rightarrow 2\text{H}_2\text{O} + \text{O}_2$ is $6.0 \times 10^{-7} \text{ s}^{-1}$, and the frequency factor (A) is $1.2 \times 10^{12} \text{ s}^{-1}$. Determine the activation energy for the reaction.

$$R = 8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}}$$

$$k = A e^{-\frac{E_a}{RT}}$$

$$e = 2.7183$$

$$T = 550^\circ\text{C} + 273.15 = 823.15 \text{ K}$$

$$k = 6.0 \times 10^{-7} \frac{1}{\text{s}}$$

$$A = 1.2 \times 10^{12} \frac{1}{\text{s}}$$

$$E_a = ?$$

Correct

$$6.0 \times 10^{-7} \frac{1}{\text{s}} = (1.2 \times 10^{12} \frac{1}{\text{s}}) (2.7183)^{\frac{-E_a}{(8.314 \frac{\text{J}}{\text{mol} \cdot \text{K}})(823.15 \text{ K})}}$$

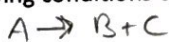
$$5.0 \times 10^{-19} = 2.7183^{\frac{-E_a}{6843.67 \frac{\text{J}}{\text{mol}}}}$$

$$1.839 \times 10^{-19} = 1^{\frac{-E_a}{6843.67 \frac{\text{J}}{\text{mol}}}}$$

x,

(-)

6. At 600 K, compound A decomposes to form compounds B and C via a first-order reaction. Discuss the effect of each of the following conditions on the half-life of A:



$$t_{\frac{1}{2}} = \frac{\ln 2}{k}$$

$$kt = \frac{100}{50}$$

- (a) Increasing the initial concentration of A

increasing the concentration of A would overall increase more reacting molecules and would then ~~reduce~~ the half life of A

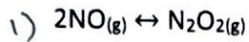
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- (b) Increasing the temperature at which the reaction occurs

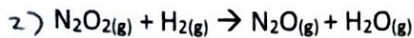
increasing the temperature would cause the collisions to happen more rapidly making the half life of A even less.

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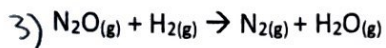
7. Consider the following:



(fast, k_1 represents the forward rate constant, k_{-1} the reverse rate constant)

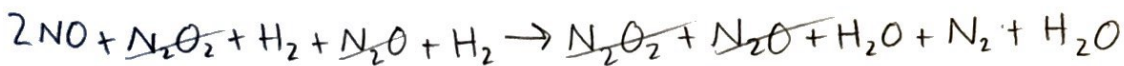


(slow, k_2 the rate constant)



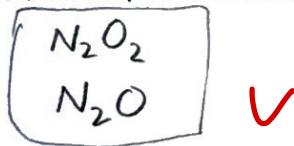
(fast, k_3 the rate constant)

(a) Write the overall reaction.



✓

(b) Identify all intermediates.



(c) Write the overall rate law.

$$1) \text{rate}_1 = k_1[2\text{NO}]^2 = k_{-1}[\text{N}_2\text{O}_2]$$

$$\text{N}_2\text{O}_2 = \frac{k_1[2\text{NO}]^2}{k_{-1}}$$

$$2) \text{rate}_2 = k_2[\text{N}_2\text{O}_2][\text{H}_2]$$

$$k_2 \left(\frac{k_1[2\text{NO}]^2}{k_{-1}} \right) + [\text{H}_2]$$

(-)

$$k_2 \left(\frac{k_1}{k_{-1}} [2\text{NO}]^2 \right) + [\text{H}_2]$$

$$\text{rate} = \boxed{k[2\text{NO}]^2 + [\text{H}_2]}$$