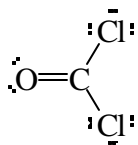


2001 CHEM1405 (Vet. Science)

2001-J-2

- $\text{Mn}(\text{ClO}_4)_2(\text{s}) \rightarrow \text{Mn}^{2+}(\text{aq}) + 2\text{ClO}_4^{-}(\text{aq})$
- iron(III) chloride-6-water
 $(\text{NH}_4)_2\text{CO}_3$
potassium dichromate
 PbO_2
- -835 kJ mol^{-1}
- Particles in a negatively charged sol repel each other. Addition of a cation overcomes the interparticle repulsions and allows the particles to get closer to each other and hence promotes coagulation.

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2001-J-3

- 666 g mol^{-1}
- $1.45 \times 10^{-3} \text{ M}$
- $\text{HPO}_4^{2-}(\text{aq}) + \text{H}_3\text{O}^+(\text{aq}) \rightarrow \text{H}_2\text{PO}_4^{-}(\text{aq}) + \text{H}_2\text{O}$
 $\text{H}_2\text{PO}_4^{-}(\text{aq}) + \text{OH}^{-}(\text{aq}) \rightarrow \text{HPO}_4^{2-}(\text{aq}) + \text{H}_2\text{O}$

2001-J-4

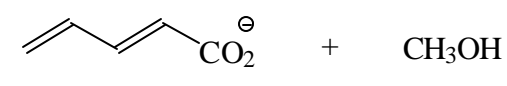
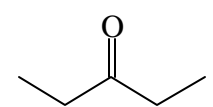
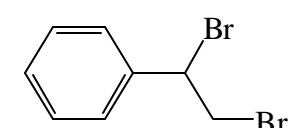
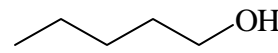
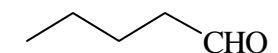
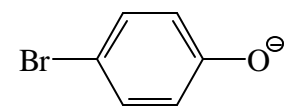
- $k = 0.210 \text{ hour}^{-1}$ $t_{1/2} = 3.30 \text{ hour}$
6.60 hour

The appropriate second order rate law is $\text{Rate} = k[\text{sucrose}][\text{water}]$.

The concentration of water (present in vast excess as the solvent) does not change over time, thus giving a pseudo first order rate law: $\text{Rate} = k_1[\text{sucrose}]$ where $k_1 = k[\text{water}]$.

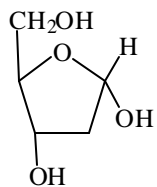
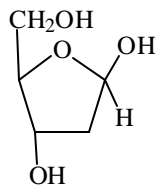
2001-J-5

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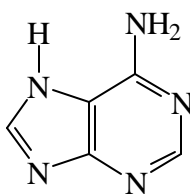
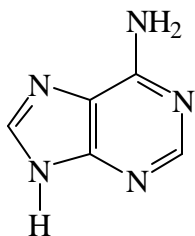
	conc. H ₂ SO ₄	
		
	excess HN(CH ₃) ₂	
		
		
 or 		
		

2001-J-6

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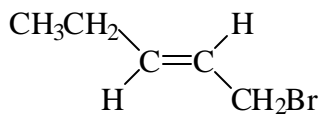
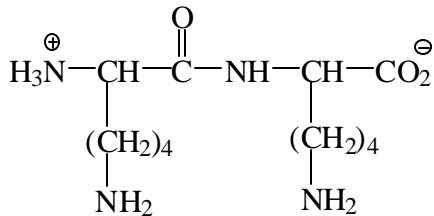
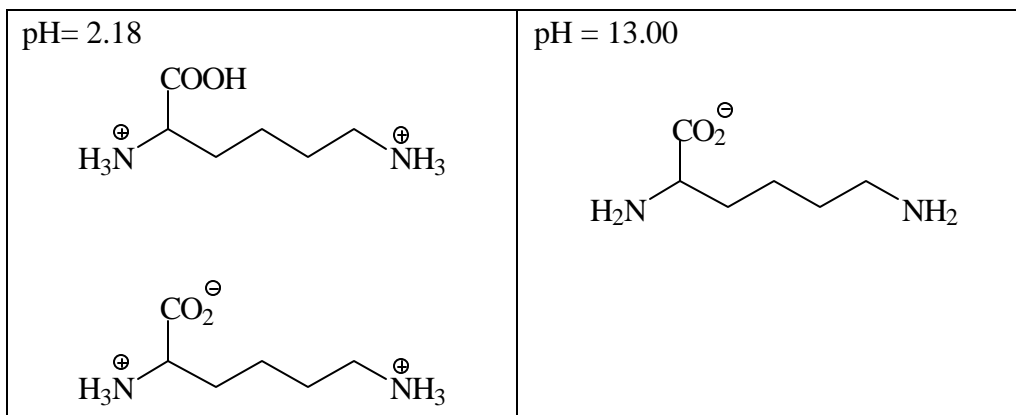


α -D-2-deoxyribofuranose



2001-J-7

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- 4-ethyl-2,3-dimethyl-1-hexanol

2001-J-8

