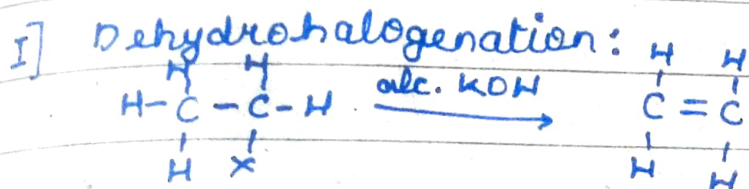
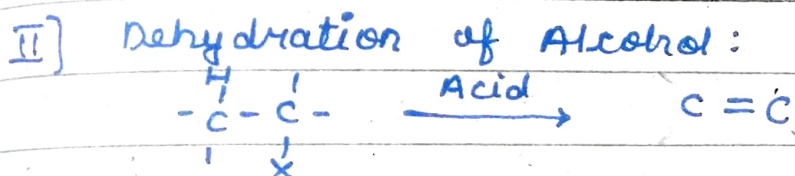


# ELIMINATION REACTIONS - 303



Also known as: 1,2 elimination,  $\beta$  elimination



$\hat{=}$   $E_2$  Reaction:

- II<sup>nd</sup> order Reaction
- depends on conc<sup>n</sup> of Both reagents (alkyl halide and base) Single Step reac<sup>n</sup>.
- strong base should be used

$\hat{=}$   $E_1$  Reaction:

- I<sup>st</sup> order Reaction
- depends only the conc<sup>n</sup> of alkyl halide
- Two step Reac<sup>n</sup>.
- weak base is used.

10/3/18

\* Kinetics of elimination Reaction:

A concentrated sol<sup>n</sup> of strong base follows II<sup>nd</sup> order kinetics i.e. the rate of alkene formation depends upon the concentra<sup>n</sup> of both substrate i.e. alkyl halide and base.

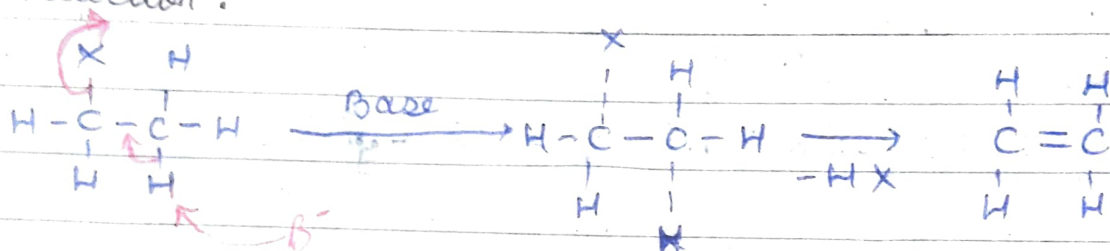
$$\text{Rate} = k [\text{R-X}] [\text{B}]$$

BASE

If we reduce the strength / concentration of base, it follows first order kinetics. The rate of elimination depends only upon the conc<sup>n</sup> of alkyl halide.

$$\text{Rate} = k[\text{R-X}]$$

\* E<sub>2</sub> Reaction:



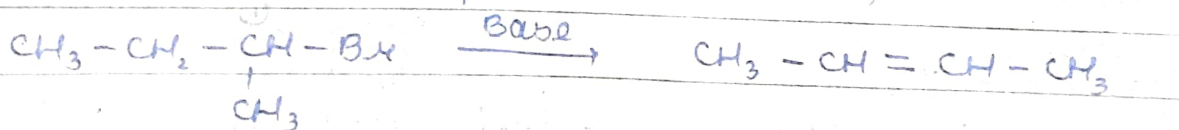
\* Heterolytic bond dissociation energy: shows the strength of carbon <sup>halogen</sup> hydrogen bonds.



\* Reactivity <sup>of alkyl halide</sup> towards elimination:  $\text{R-I} > \text{R-Br} > \text{R-Cl} > \text{R-F}$

Saytzeff's Rule: In dehydrohalogenation the preferred product is the alkene that has the greater no. of alkyl groups attached to the doubly bonded carbon atoms. OR

Poorer becomes more poorer.



The structure of more branched <sup>at</sup> the carbon carrying the halogen provided a greater no. of  $\beta$ -hydrogen for attack by base and hence more favourable factor towards elimination.

Reactivity of alkyl halide towards elimination: (E<sub>2</sub>)



Case of formation of alkene -



The more stable the alkene, the faster its formation.