



Chapter- Organic chemistry some basic principle and techniques and halo alkanes and halo arenes

Topic- Reaction intermediate θ attacking reagents and chemical properties of halo alkanes θ halo arenes

CLASS - XI, XII

DESCRIPTION

The neucleophillic substitution reactions is very common in alkyl halides but these reactions can takes place by these mechanisms based on Substrate and Solvents

1) SN1 reactions (Substitution Neucleophillic Unimolecular Reactions)

In this mechanism carbocation is the inter mediate.

Reactivity order of alkylhalides - $3^{\circ} > 2^{\circ} > 1^{\circ}$

Sterco chemistry of products - Retention and Inversion

Solvent - Polar protic solvents (eq alcohols).

 $\ln SN^{3}$ reaction since the slow step involves only on the substrate, So the molecularity of the reaction is one.

2) SN² Reactions (Substitution Neucleophillic Bimolecular Reactions)

In this mechanism transition state is the intermediate.

Due tot steric hinderance the neucleophillic attacks from the back side to the product have inversion of canfiguration.

Reactivity order alkylhalides - $1^{\circ} > 2^{\circ} > 3^{\circ}$.

Solvents - Polar aprotic solvents (DMSO, acetone etc).

3) SN, Reaction (Substitution Neucleophillic Internal)

The solvents used in SN₄ reaction is diethylether or THF.

During the SN^I reaction there occurs an internal transformation through a cyclic transition state to which attack of chloride from front side leads to retention in configuration.

Note: If pyridine is used as a solvent it will react with thionyl chloride to Form pyridinium chloride. Hence the chloride ion attack the carbon atom from back side. As a result inversion of configuration of product takes place. So the mechanism is not SN_1 but is actually SN_2 .



