

## 7.8: THE S<sub>N</sub>1 REACTION

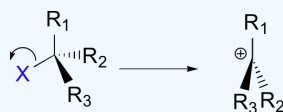
### Learning Objective

- determine the rate law & predict the mechanism based on its rate equation or reaction data for S<sub>N</sub>1 reactions
- predict the products and specify the reagents for S<sub>N</sub>1 reactions with stereochemistry
- propose mechanisms for S<sub>N</sub>1 reactions
- draw and interpret Reaction Energy Diagrams for S<sub>N</sub>1 reactions

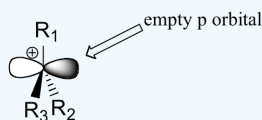
### THE S<sub>N</sub>1 MECHANISM WITH STEREOCHEMISTRY

A second model for a nucleophilic substitution reaction is called the '**dissociative**' or '**S<sub>N</sub>1**' mechanism. In many cases, the nucleophile is the solvent, so this mechanism can also be called "solvolysis".

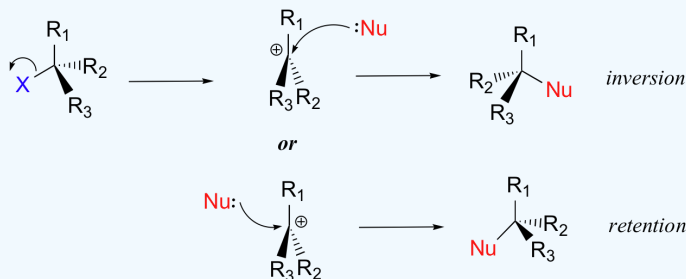
Step 1: In the S<sub>N</sub>1 mechanism, the carbocation forms when the C-X bond breaks *first*, before the nucleophile approaches



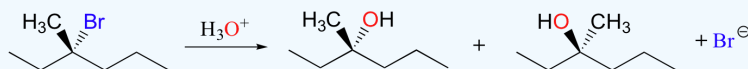
The carbocation has a central carbon with only three bonds and bears a formal charge of +1. Recall that a carbocation should be pictured as  $sp^2$  hybridized, with trigonal planar geometry. Perpendicular to the plane formed by the three  $sp^2$  hybrid orbitals is an empty, unhybridized  $p$  orbital.



Step 2: The nucleophile reacts with the empty, 'electron hungry'  $p$  orbital of the carbocation to form a new bond and return the carbon to tetrahedral geometry. Because of this trigonal planar geometry, the nucleophile can approach the carbocation from either lobe of the empty  $p$  orbital (aka either side of the carbocation). This means that about half the time the product has the same stereochemical configuration as the starting material (retention of configuration), and about half the time the stereochemistry has been inverted. In other words, *racemization* of the product occurs during S<sub>N</sub>1 reactions if the electrophilic carbon is chiral. If the intermediate from a chiral alkyl halide survives long enough to encounter a random environment, the products are expected to be racemic (a 50:50 mixture of enantiomers). On the other hand, if the departing halide anion temporarily blocks the front side, or if a nucleophile is oriented selectively at one or the other face, then the substitution might occur with predominant inversion or even retention of configuration.



As an example, the tertiary alkyl bromide below, (S)-3-bromo-3-methylhexane, would be expected to form a racemic mix of *R*- and *S*-3-methyl-3-hexanol after an S<sub>N</sub>1 reaction with water as the nucleophile.

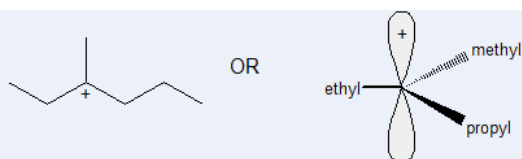


### Exercise

- Draw the structure of the intermediate in the two-step nucleophilic substitution reaction (S<sub>N</sub>1) above.

Solution

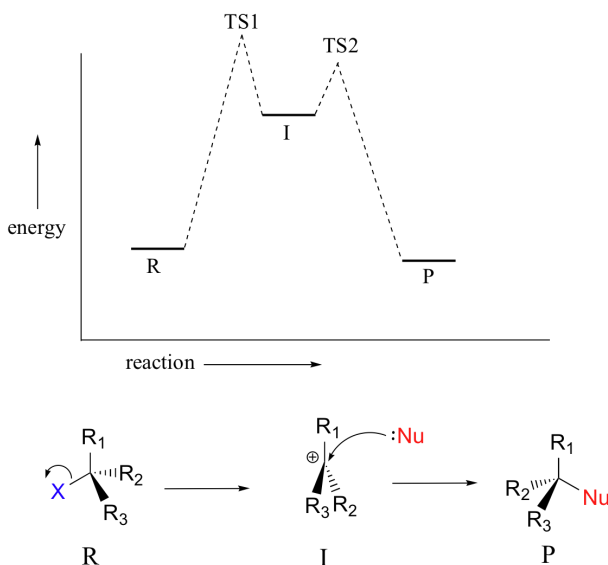
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When interpreting bond-line structures, it is useful to visualize the electronic structure with an unhybridized, empty p orbital extending perpendicular to the alkyl groups.

## THE S<sub>N</sub>1 REACTION ENERGY DIAGRAM

The S<sub>N</sub>1 reaction is an example of a two-step reaction with a reaction intermediate. Evaluating reactive intermediates is a very important skill in the study of organic reaction mechanisms. Many important organic reactions do not occur in a single step; rather, they are the sum of two or more discrete bond-forming / bond-breaking steps, and involve transient intermediate species that go on to react very quickly. In the S<sub>N</sub>1 reaction, the carbocation species is a reaction intermediate. A potential energy diagram for an S<sub>N</sub>1 reaction shows that the carbocation intermediate can be visualized as a kind of valley in the path of the reaction, higher in energy than both the reactant and product but lower in energy than the two transition states.

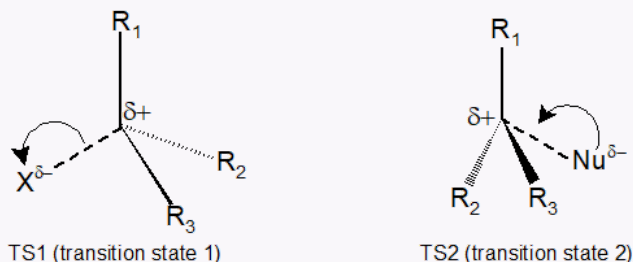


### Exercise

2. Draw structures representing transition state 1 (TS1) and transition state 2 (TS2) in the reaction above. Use the solid/dash wedge convention to show three dimensions.

Solution

2.



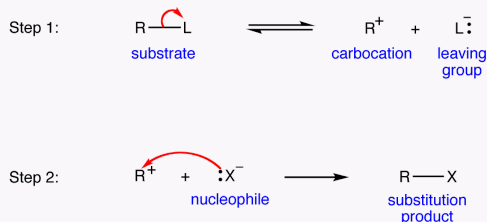
## S<sub>N</sub>1 REACTION KINETICS

In the first step of an S<sub>N</sub>1 mechanism, two charged species are formed from a neutral molecule. This step is much the slower of the two steps, and is therefore rate-determining. In the reaction energy diagram, the activation energy for the first step is higher than that for the second step indicating that the S<sub>N</sub>1 reaction has *first order* kinetics because the rate determining step involves one molecule splitting

apart, not two molecules colliding. It is important to remember that first order refers to the rate law expression where the generic term substrate is used to describe the alkyl halide.

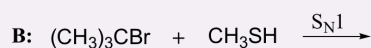
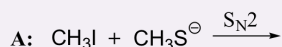
$$\text{rate} = k [\text{substrate}]$$

Because an  $S_N1$  reaction is first order overall the concentration of the nucleophile does not affect the rate. The implication is that the nucleophile does not participate in the rate limiting step or any prior steps, which suggests that the first step is the rate limiting step. Since the nucleophile is not involved in the rate-limiting first step, the nature of the nucleophile does not affect the rate of an  $S_N1$  reaction.

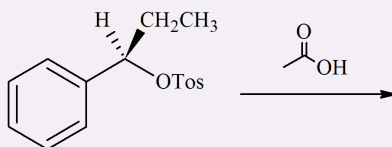


### Exercise

3. Consider two nucleophilic substitutions that occur uncatalyzed in solution. Assume that reaction A is  $S_N2$ , and reaction B is  $S_N1$ . Predict, in each case, what would happen to the rate of the reaction if the concentration of the nucleophile were doubled, while all other conditions remained constant.



4. Give the products of the following  $S_N1$  reaction. Show stereochemistry.

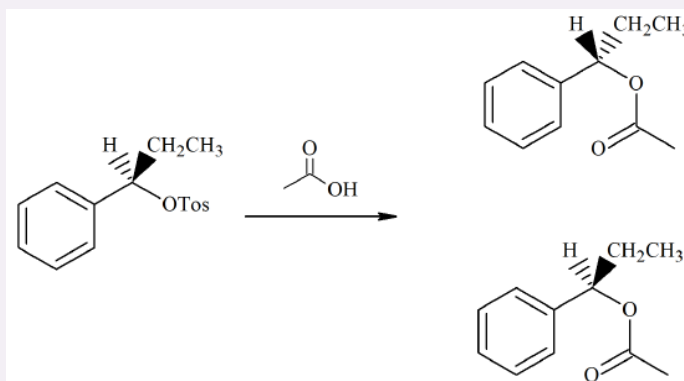


Solution

3. For Reaction A, the rate law is  $\text{rate} = k[\text{CH}_3\text{I}][\text{CH}_3\text{S}^-]$ . Therefore, if the concentration of the nucleophile,  $\text{CH}_3\text{S}^-$ , is doubled and the concentration of the alkyl halide remains the same, then the reaction rate will double.

For Reaction B, the rate law is  $\text{rate} = k[(\text{CH}_3)_3\text{CBr}]$ . Therefore, if the concentration of the nucleophile,  $\text{CH}_3\text{SH}$ , is doubled and the concentration of the alkyl halide remains the same, then reaction rate stays the same.

4.



### CONTRIBUTORS AND ATTRIBUTIONS

- Dr. Dietmar Kennepohl FCIC (Professor of Chemistry, Athabasca University)
- Prof. Steven Farmer (Sonoma State University)
- William Reusch, Professor Emeritus (Michigan State U.), Virtual Textbook of Organic Chemistry

- [Organic Chemistry With a Biological Emphasis](#) by [Tim Soderberg](#) (University of Minnesota, Morris)
- Jim Clark ([Chemguide.co.uk](#))

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