

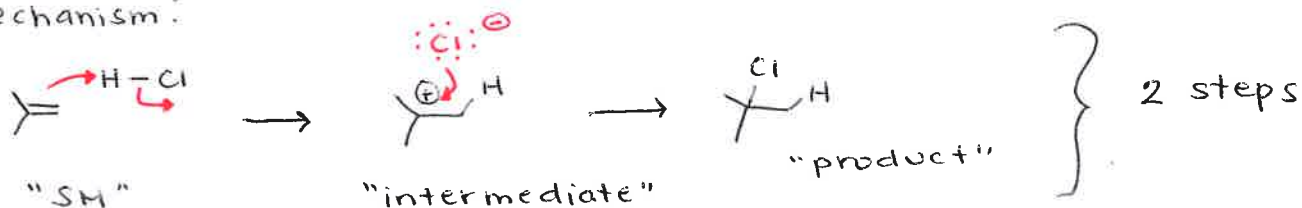
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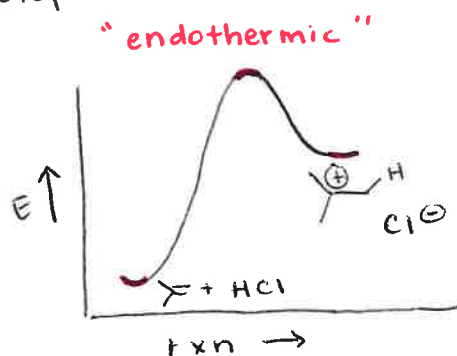
Recall "Rxn Energy Diagram" for:



Mechanism:



Step 1:



Step 2:

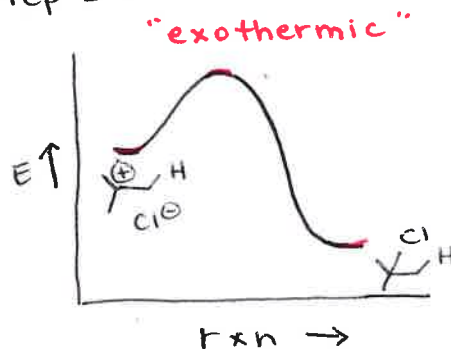
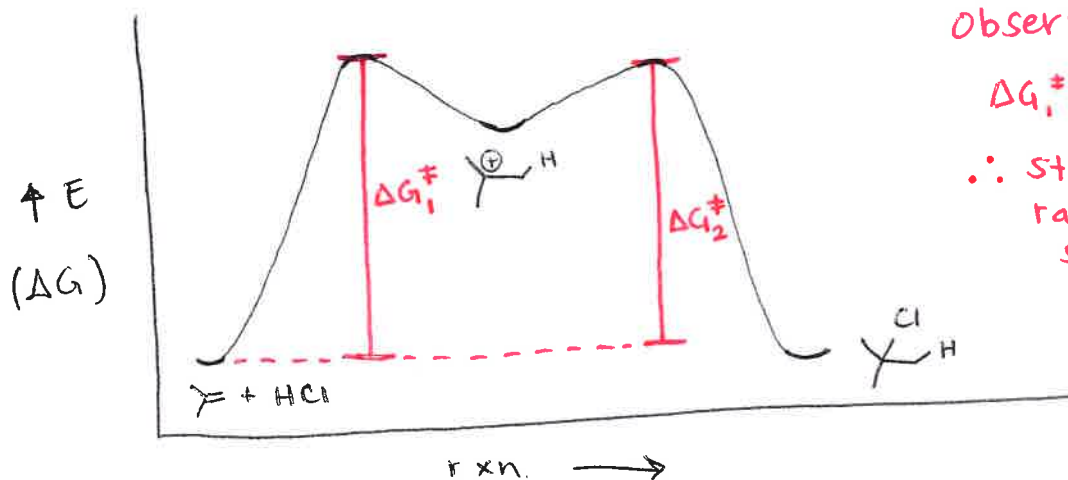


Diagram for entire process:



Observe:

$$\Delta G_1^\ddagger > \Delta G_2^\ddagger$$

\therefore Step 1 is the rate determining step (RDS).

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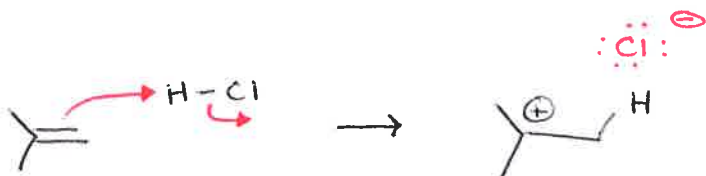
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Hammond postulate:

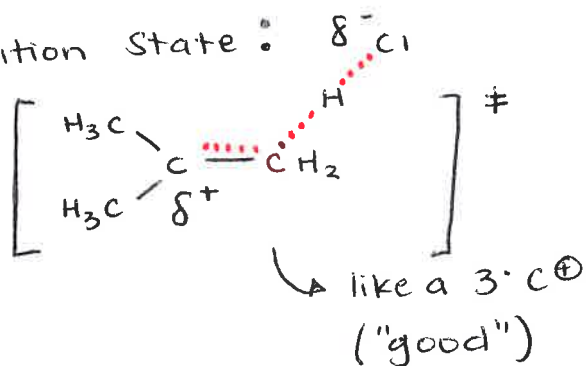
one can extrapolate from knowledge of high energy intermediates (e.g. carbocation) to behavior/stability of transition states that are adjacent in a mechanism.

Apply Hammond postulate to our example of HCl addn.

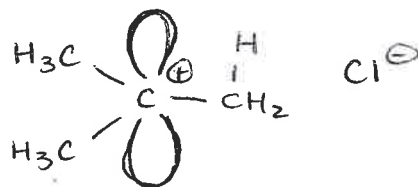
RDS



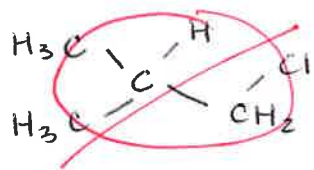
Transition State:



• ok to draw conclusions about TS from knowledge of carbocation.

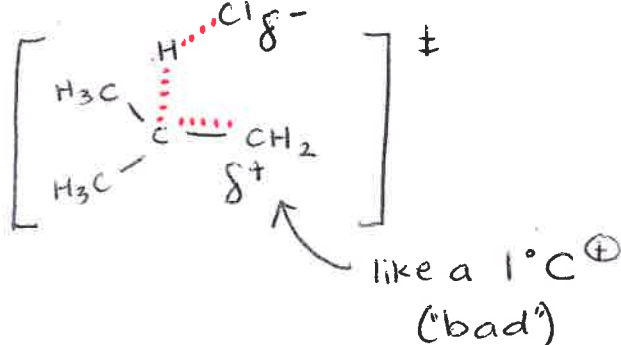


H.P helps us explain isomeric selectivity ... why we do not observe



=>

required T.S

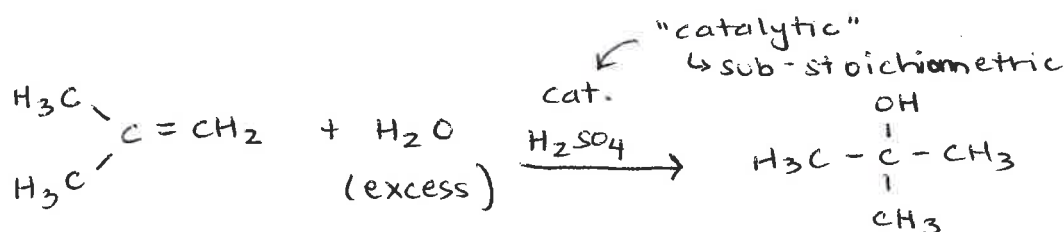
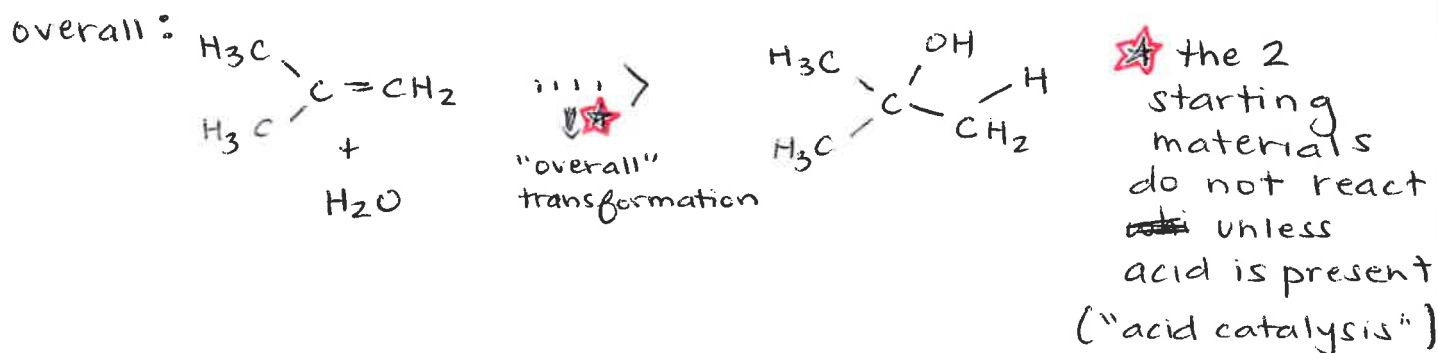


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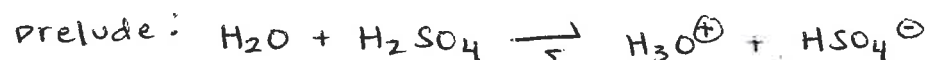
Catalysis

- catalyst is a species that enhances rxn rate w/out being consumed.

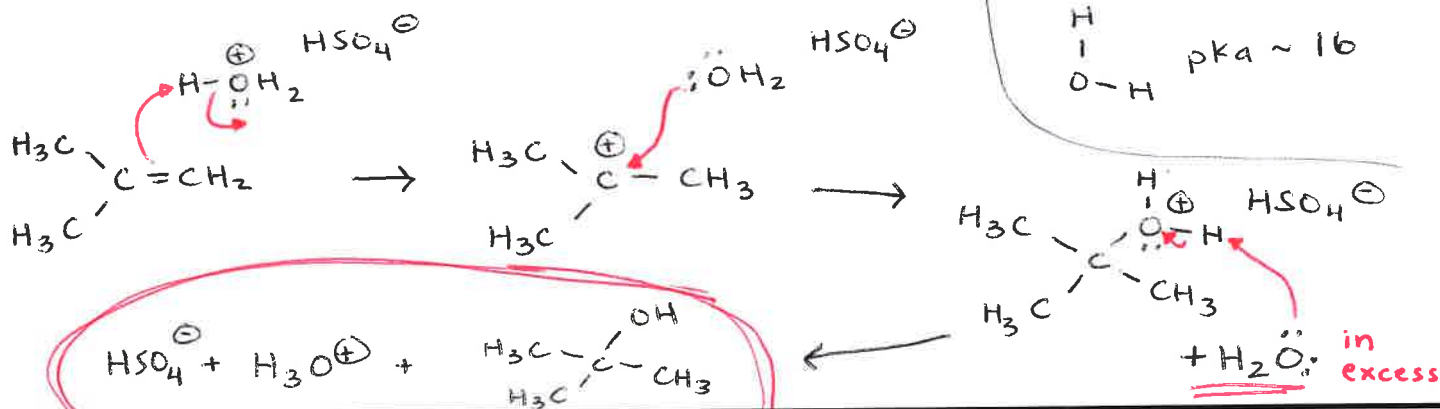
Example: Acid-catalyzed hydration of alkene



MECHANISM:



Note:
 $\text{H}-\text{O}^{\oplus}-\text{H}$ pKa ~ -2
 vs
 $\text{H}-\text{O}-\text{H}$ pKa ~ 16

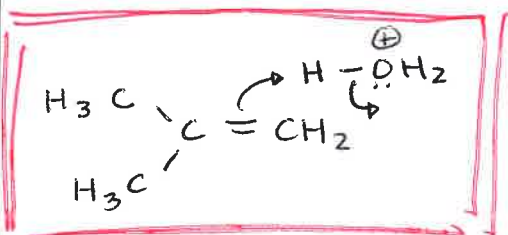


Final products
 - regenerated $\text{H}_3\text{O}^{\oplus} \Rightarrow \therefore$ catalyst

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H_2O is not sufficiently acidic for H^+ to transfer to alkene.
 $\therefore \text{H}_3\text{O}^+$ required



Note:

