

(32) Today

Sections 6.5 – 6.11
Chapter 7.3 - 7.5

Next Class (33)

Chap 7

(34) Second Class from Today

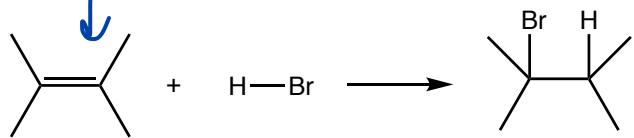
Chap 7

Third Class from Today (35)

Chap 7

not the correct mechanism

Reaction Coordinate Diagrams: Kinetics



Reactant(s) *

Product(s) *

Transition State *

Intermediate *

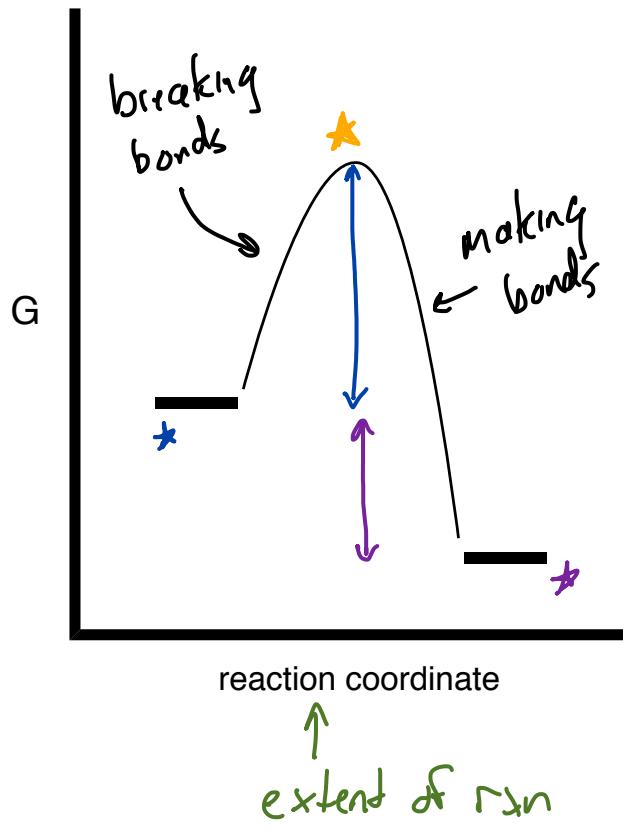
Activation Energy ΔG^\ddagger

rate of a reaction
(kinetics)

ΔG (thermodynamics)
change in Gibbs free E

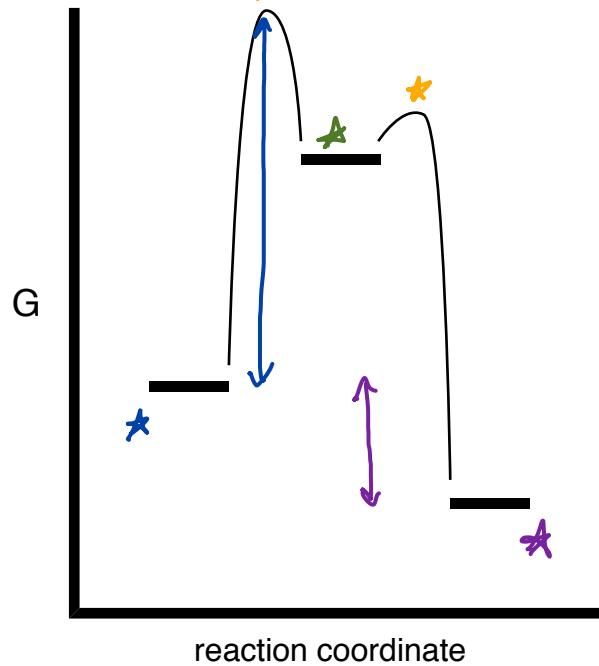
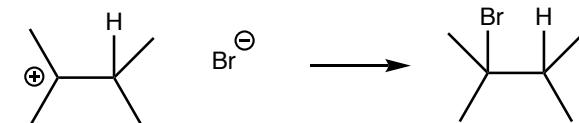
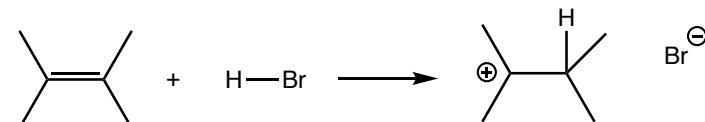
K (thermodynamics)
related to the change in
Gibbs free E

the lower the ΔG^\ddagger
the higher the
intrinsic rate (faster)



this is the correct mechanism

Section 6.9, 6.10



Reactant(s) on the left, combine to make products

Product(s) on the right, formed from reactants

Transition State - the highest energy point between reactants + products
 (reactants + intermediates, intermediates + prod) where the rxn
 switches from mostly breaking bonds (energy in) to mostly making bonds
 (energy out)
 Intermediate
 Molecules of ions that are formed and consumed during a reaction

Activation Energy ΔG^\ddagger relates to the rate of a reaction (kinetics)

the amount of energy that must be overcome for the reaction to proceed. The lower the barrier the faster the reaction

ΔG (thermodynamics)

The change in Gibbs free energy from reactants to products...

Extent of rxn, whether the rxn will go or not

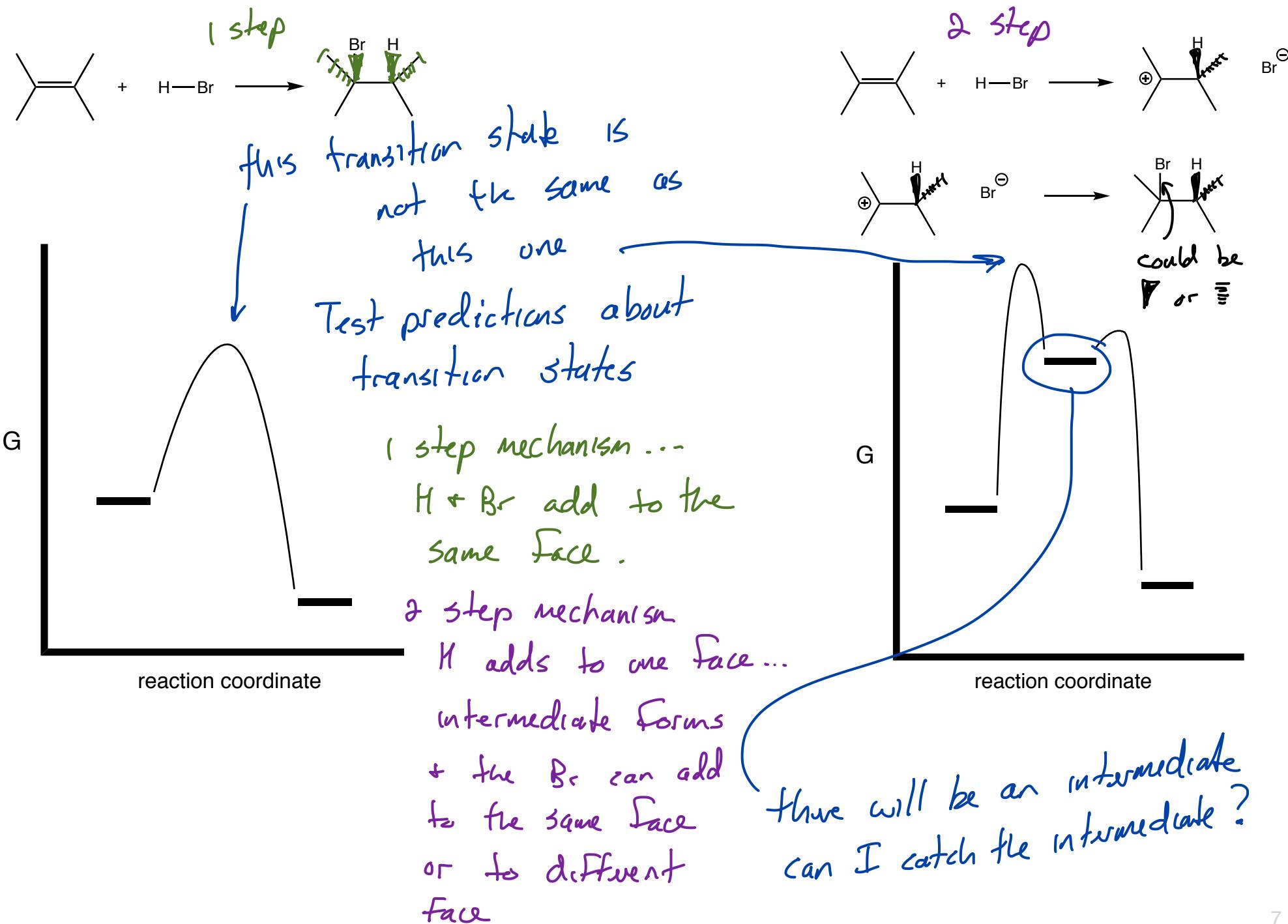
K (thermodynamics)
 Same as above... K is related to ΔG ... extent of rxn

favorable, spontaneous $\Delta G < 0, K > 1$

$$K = \frac{[\text{prod}]}{[\text{react}]}$$

Reaction Coordinate Diagrams: Kinetics

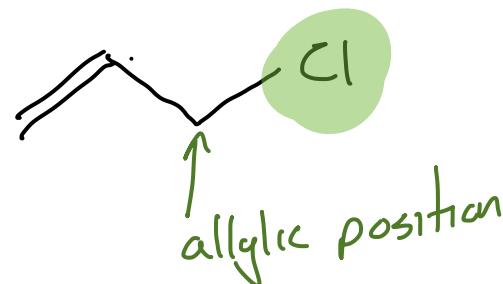
Section 6.9, 6.10



Same rules as alkanes except, alkenes are a functional group, so the position of the double bond gets the lowest number and "ane" ending of parent hydrocarbon is changed to "ene" and the double bond must be contained in the longest carbon chain.

A note on some special names

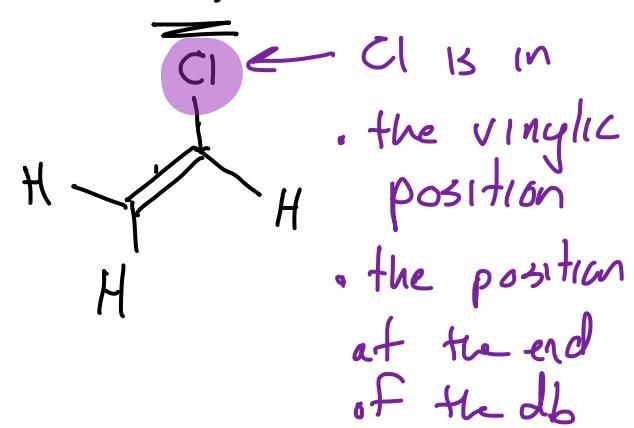
Allyl



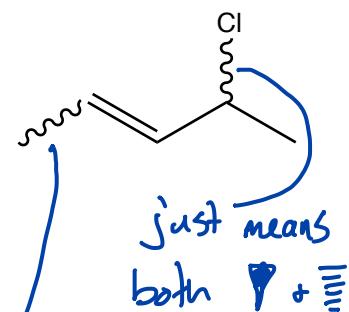
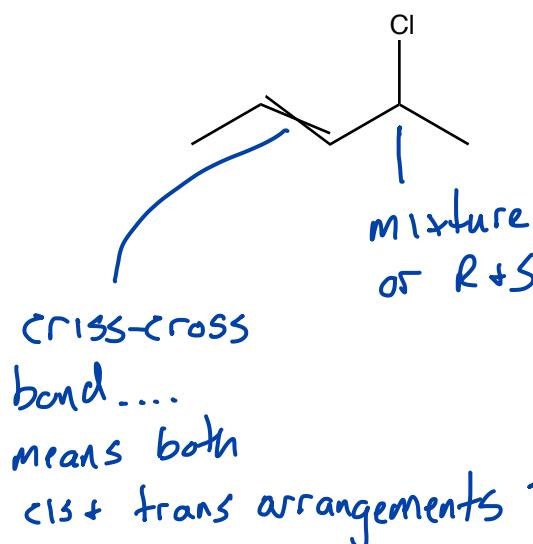
and

PVC polyvinylchloride

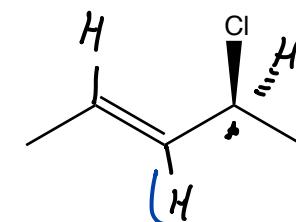
Vinyl



A note on how structures are drawn



Zero attempt
to show
stereochemistry



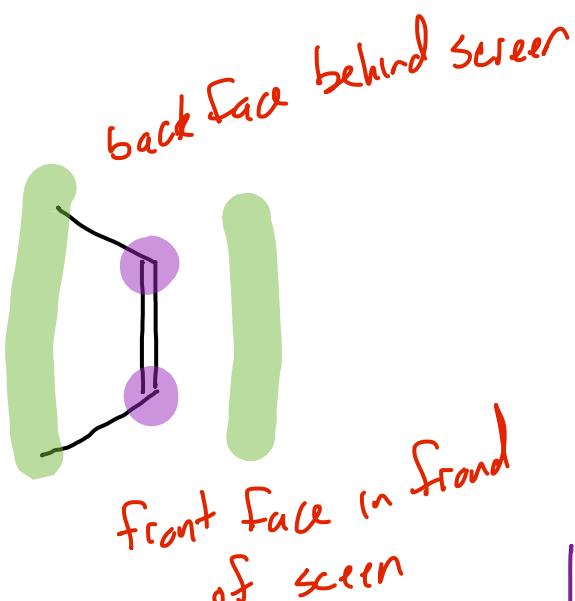
a specific
structure with
stereochemistry
shown

Ends, Sides, and Faces

→ split lengthwise along the node
of the π bond

split lengthwise along π bond

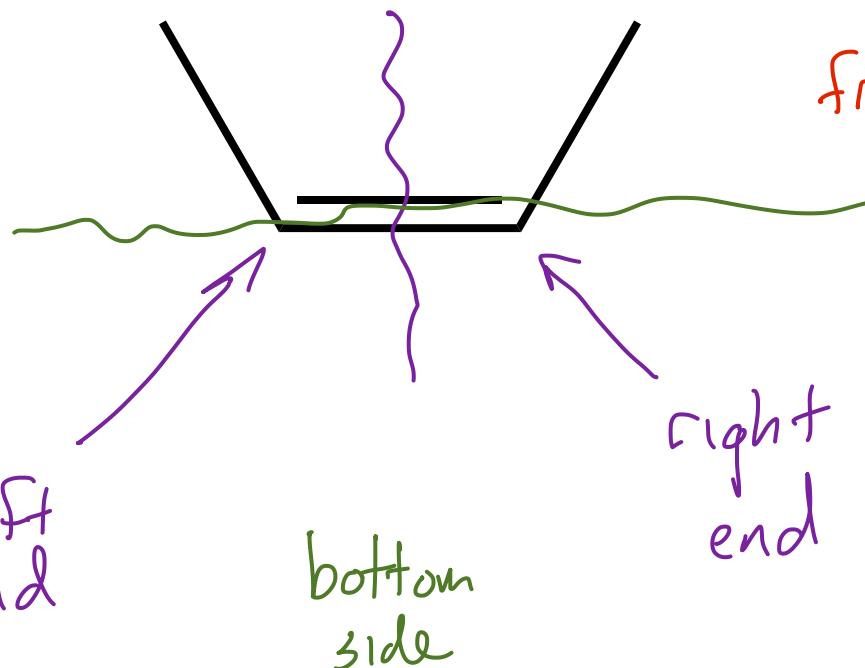
cut between C + C



top side

back face behind the screen

front face in front of the screen



H — top face — H — H 's on the side behind the screen

H — bottom face — methyl groups on the side in front of the screen

cis and *trans* Stereoisomers in alkenes

Section 7.4

