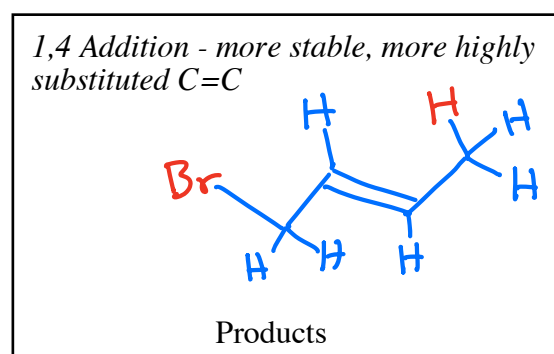
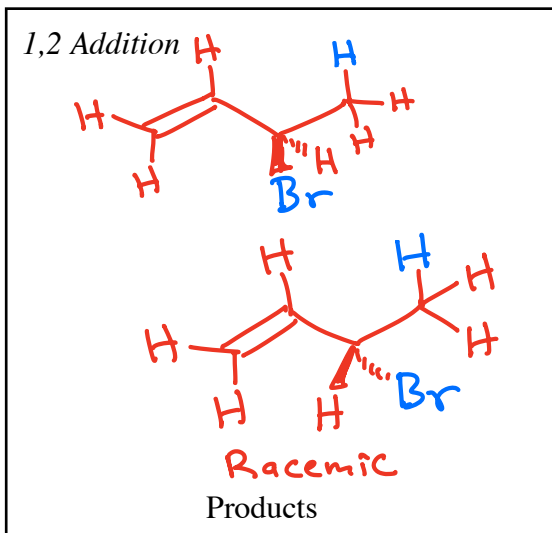
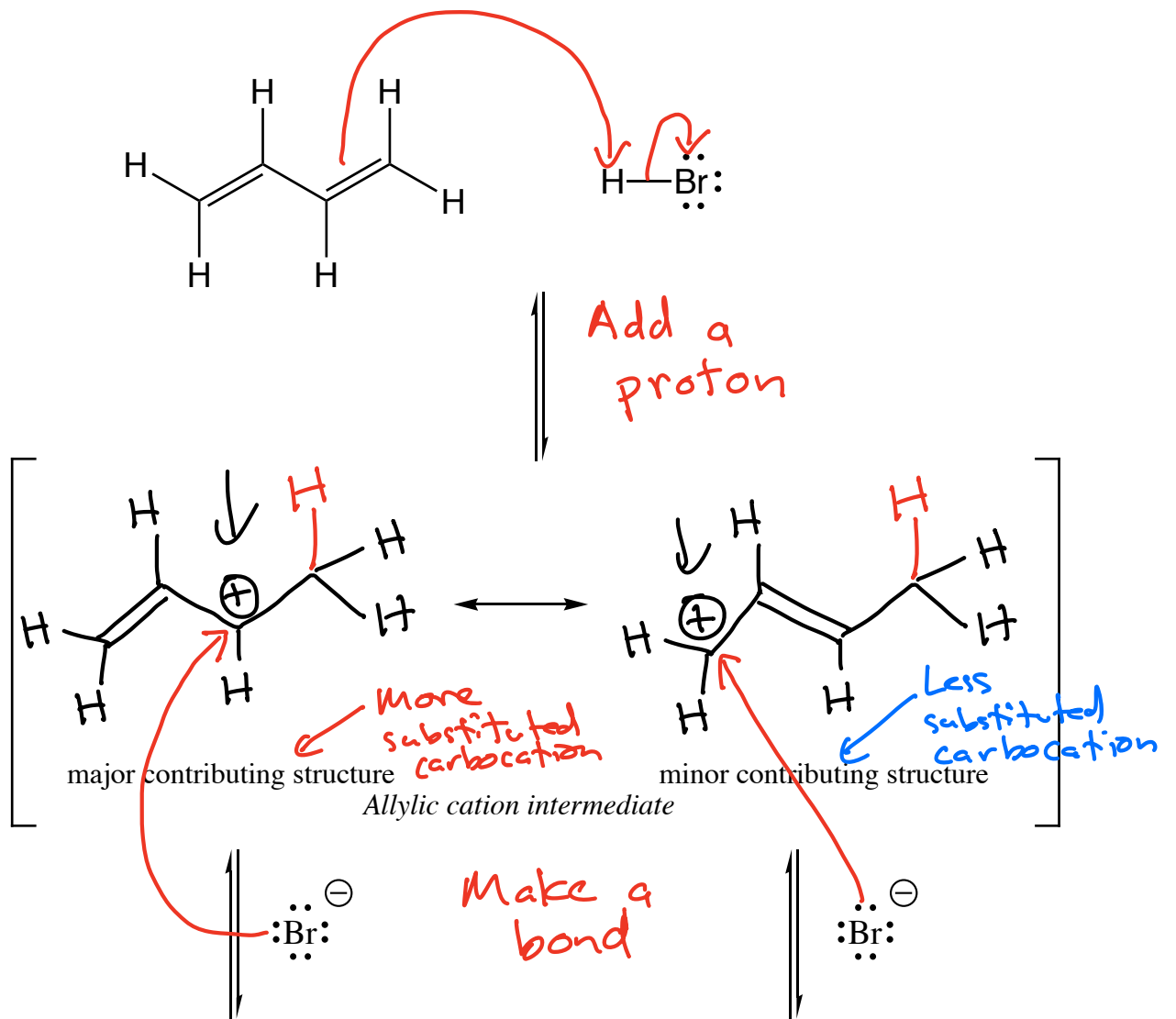
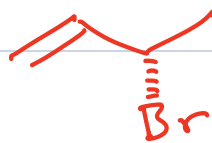
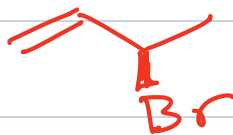
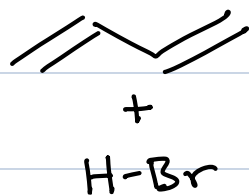


H-X reacting with conjugated dienes



1,2 addition

1,4 addition



Racemic



Temperature of
Reaction

-78°C

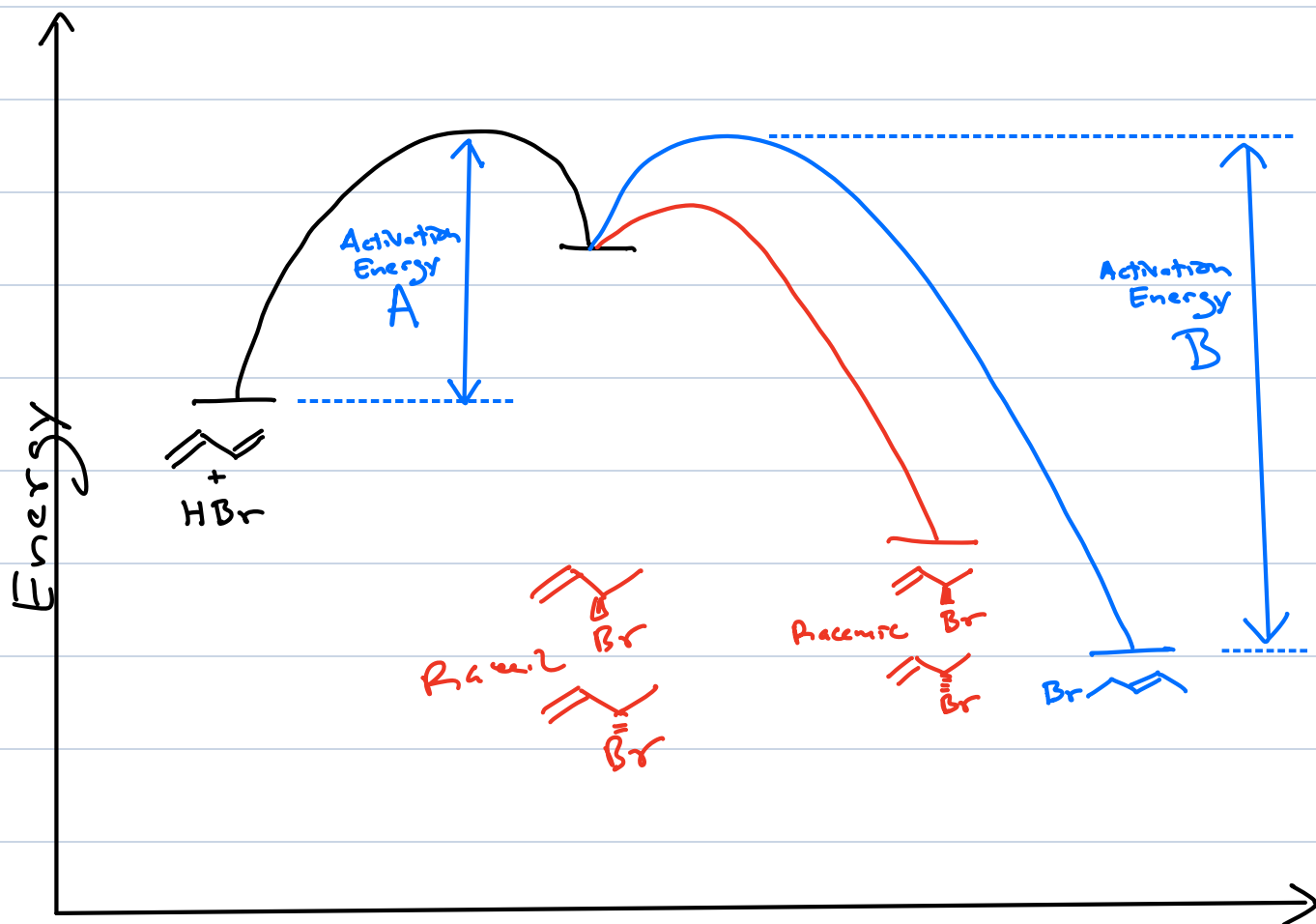
90%

10%

+40°C

15%

85%

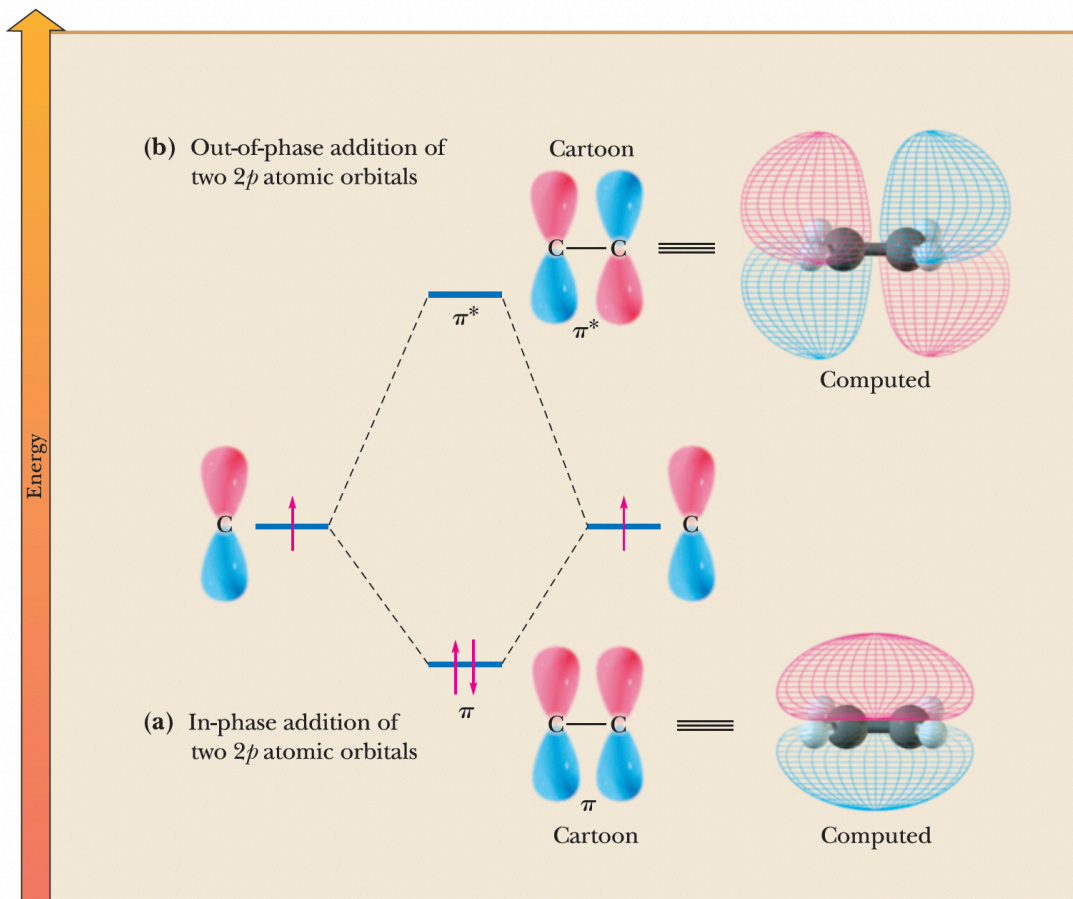


Low temperature \rightarrow Molecules have enough energy to get over activation energy A, but not enough energy to get over activation energy B.

Kinetic Control
 "Fastest" wins

High temperature \rightarrow Molecules have enough energy to get over activation energy A and activation energy B

Thermodynamic Control
 Most stable product wins

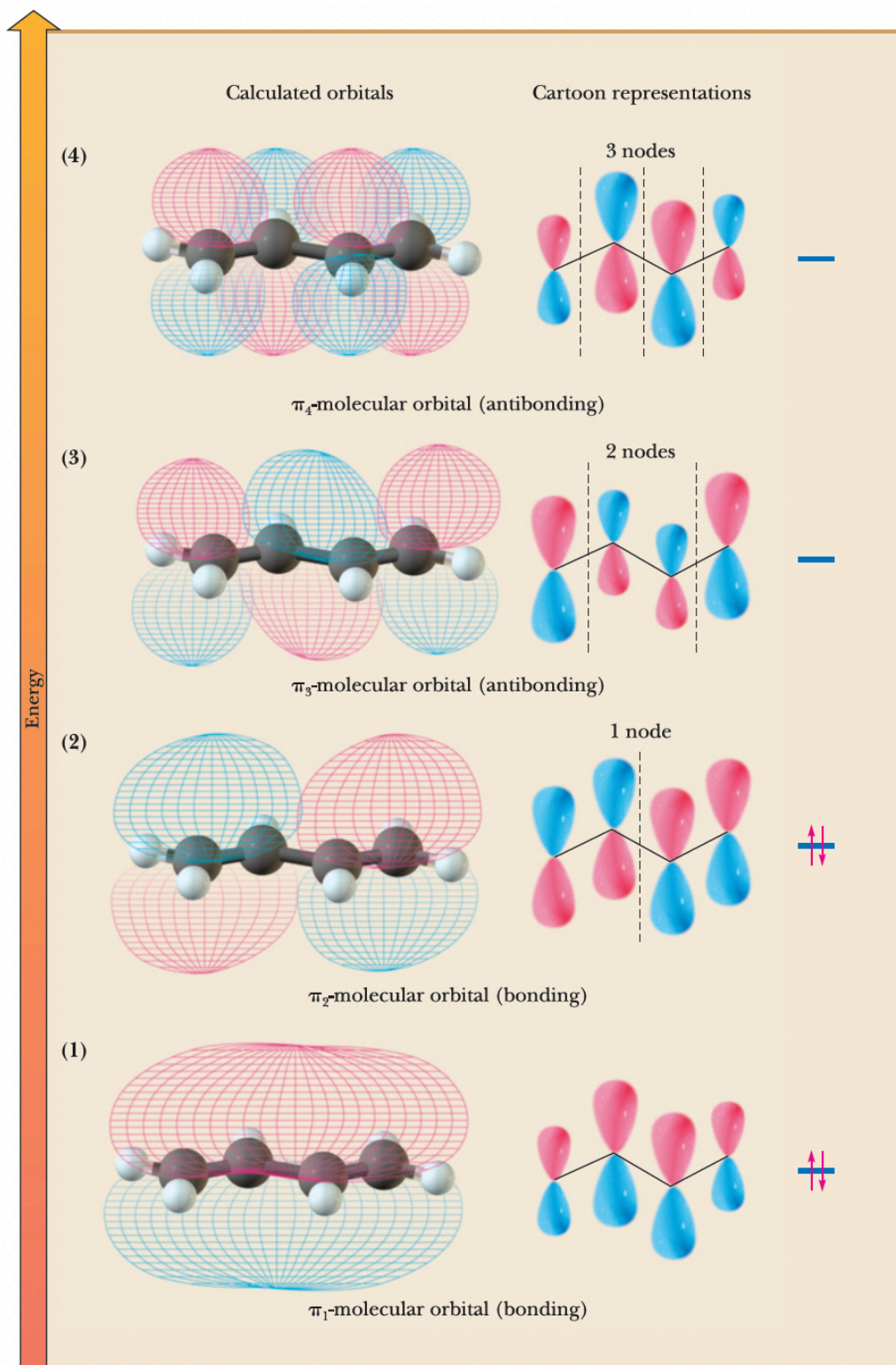


[Watch a video explanation](#)

FIGURE 1.21

Molecular orbital mixing diagram for the creation of any C—C π bond. (a) Addition of two p atomic orbitals in phase leads to a π orbital that is lower in energy than the two separate starting orbitals. When populated with two electrons, the π orbital gives a π bond. (b) Addition of the p orbitals in an out-of-phase manner (meaning a reversal of phasing in one of the starting orbitals) leads to a π^* orbital. Population of this orbital with one or two electrons leads to weakening or cleavage of the π bond, respectively.





[Watch a video explanation](#)

FIGURE 20.2 Structure of 1,3-butadiene—molecular orbital model. Combination of four parallel 2p atomic orbitals gives two π -bonding MOs and two π -antibonding MOs. In the ground state, each π -bonding MO is filled with two spin-paired electrons. The π -antibonding MOs are unoccupied.

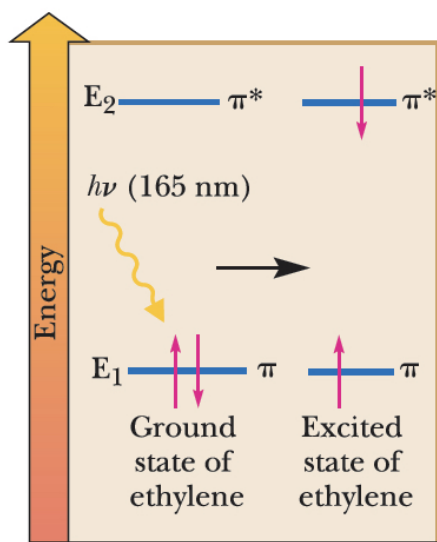


FIGURE 20.6 A $\pi \rightarrow \pi^*$ transition in excitation of ethylene. Absorption of ultraviolet radiation causes a transition of an electron from a π -bonding MO in the ground state to a π -antibonding MO in the excited state. There is no change in electron spin.

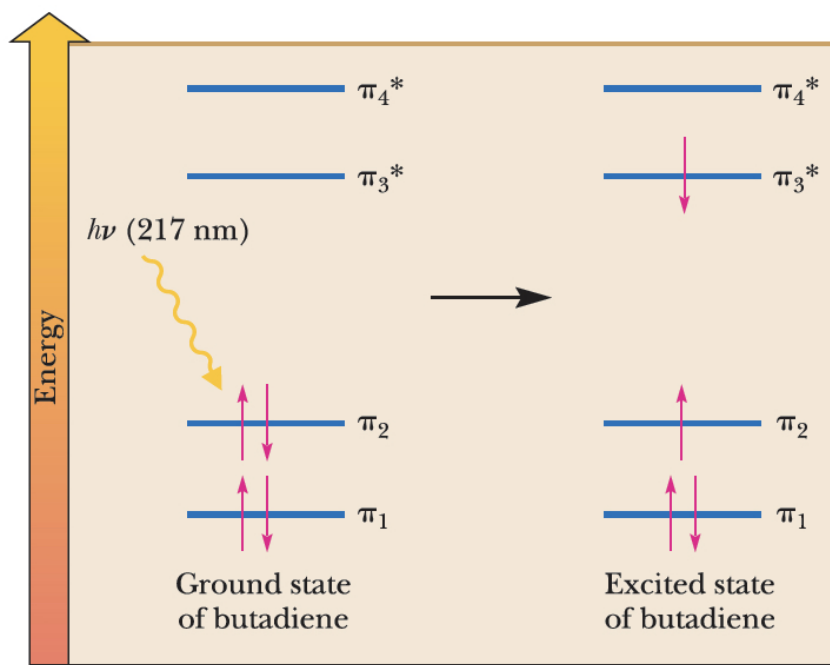


FIGURE 20.7 Electronic excitation of 1,3-butadiene; a $\pi \rightarrow \pi^*$ transition.

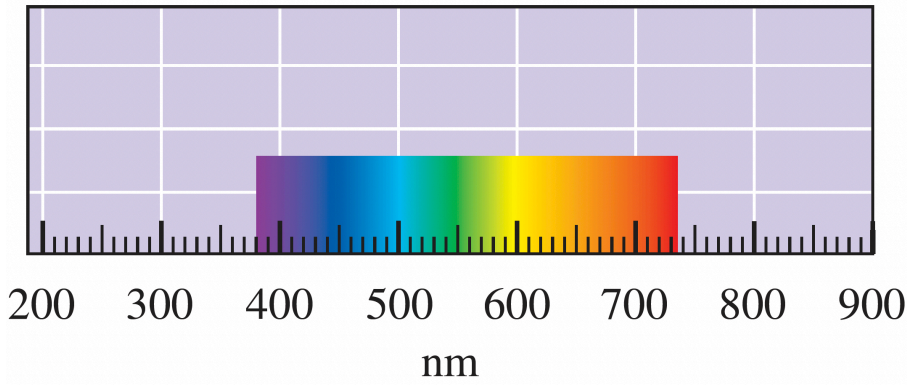
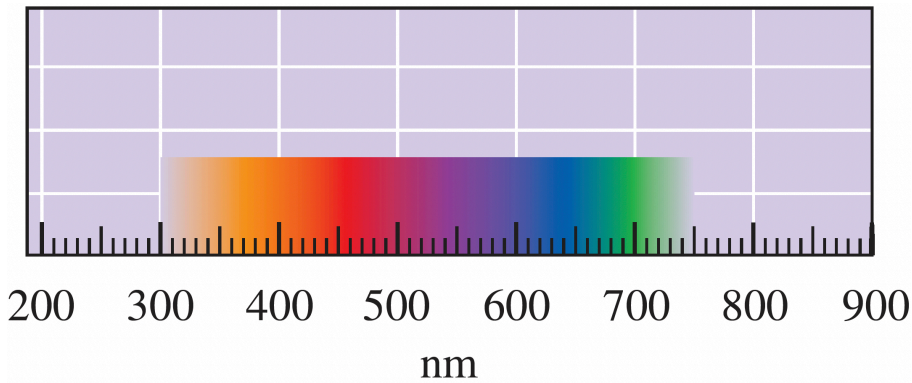
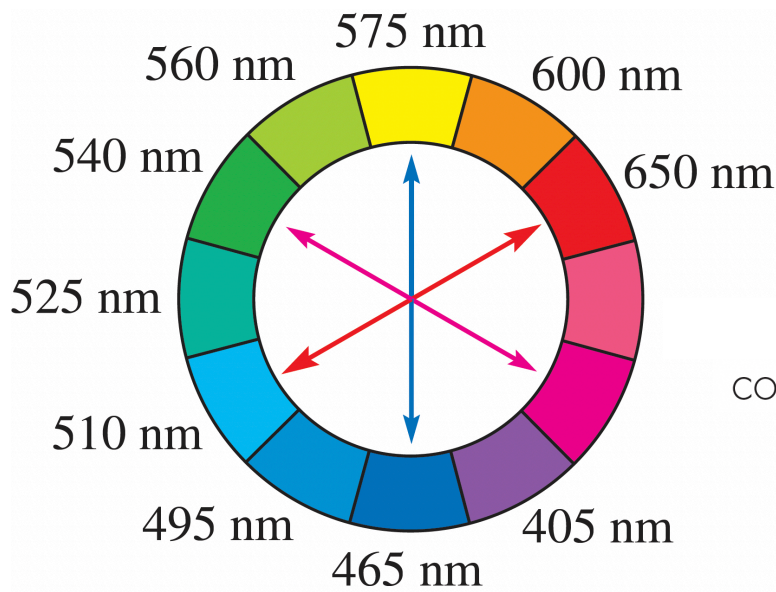


FIGURE 20.5 (a) Visible light color-wavelength correlation.

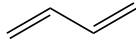


(b) Approximate color of substance (reflected light) if a single wavelength (i.e., the wavelength listed on the numerical scale of the x-axis) is absorbed.



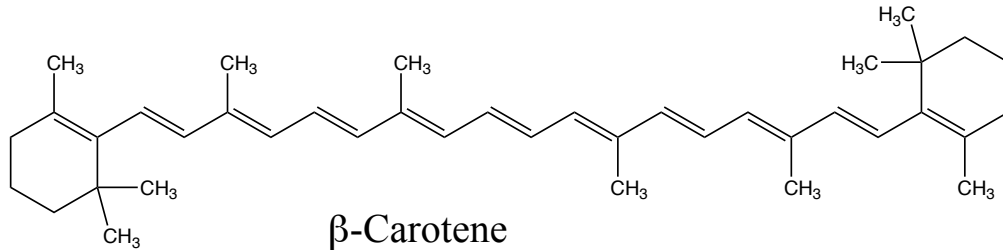
(c) Complementary colors on a color wheel.

Colored arrows are complementary



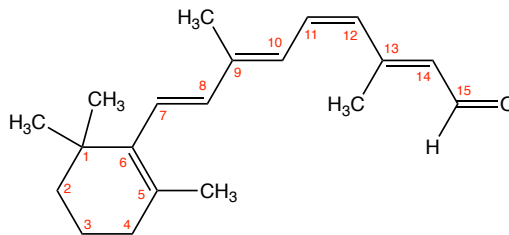
Butadiene

$\lambda_{\max} = 217 \text{ nm}$



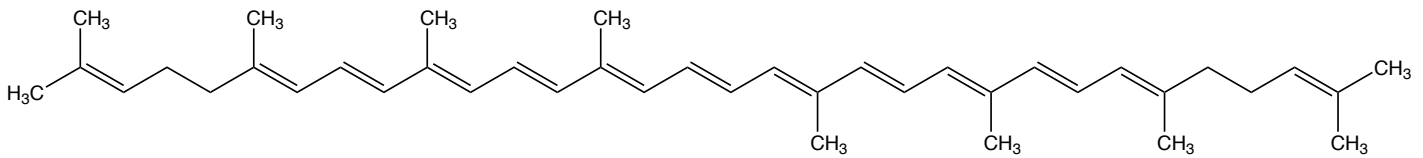
β -Carotene

$\lambda_{\max} = 455 \text{ nm}, 483 \text{ nm}$



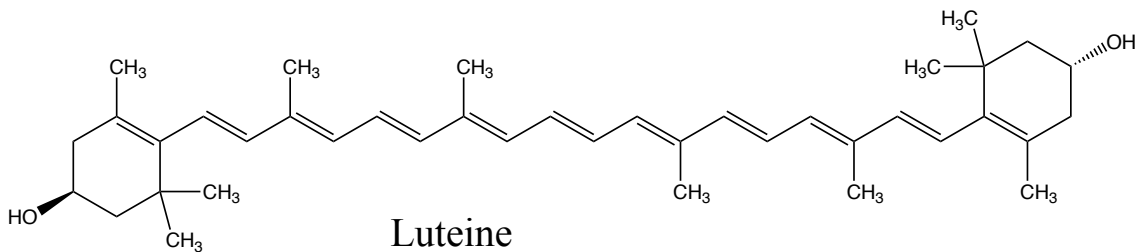
11-*cis*-Retinal

$\lambda_{\max} = 380 \text{ nm}$



Lycopene

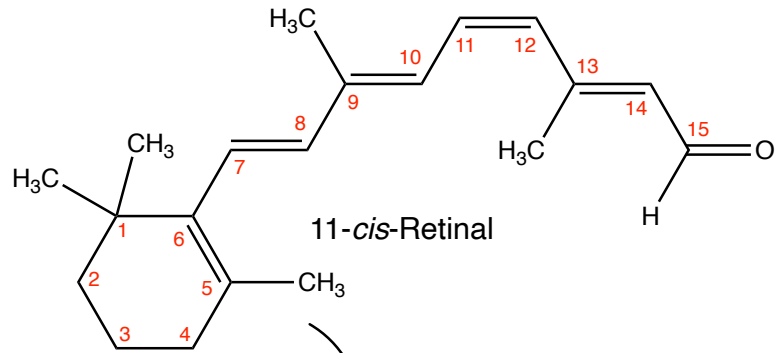
$\lambda_{\max} = 443 \text{ nm}, 471 \text{ nm}, 502 \text{ nm}$



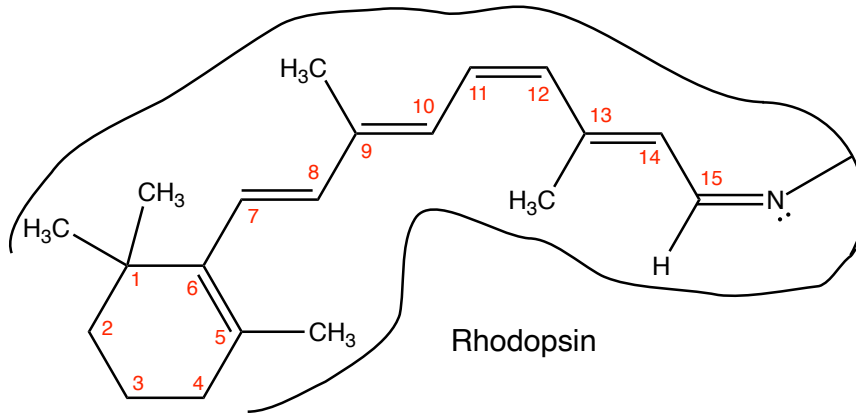
Luteine

$\lambda_{\max} = 445 \text{ nm}, 474 \text{ nm}$

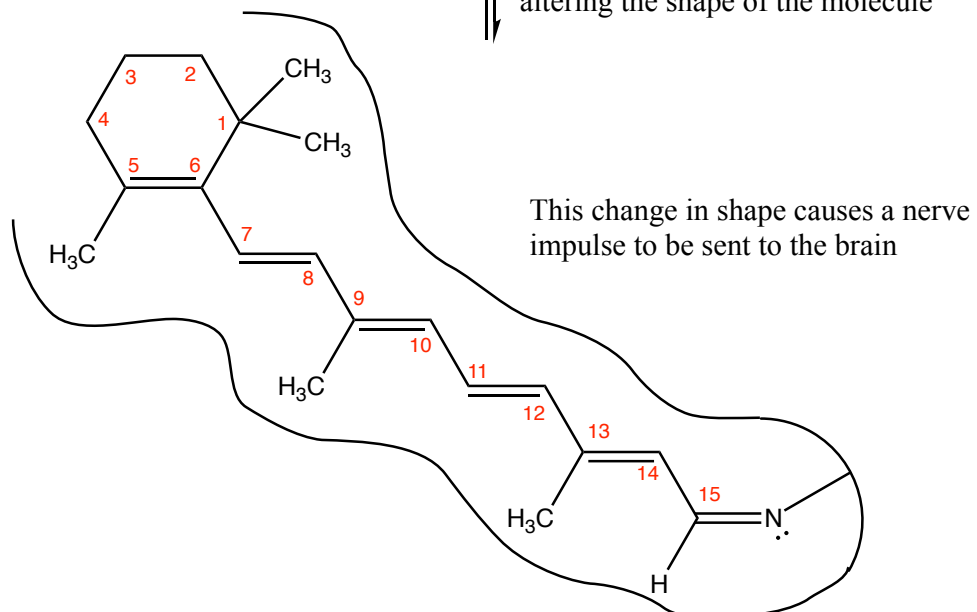
How vision works



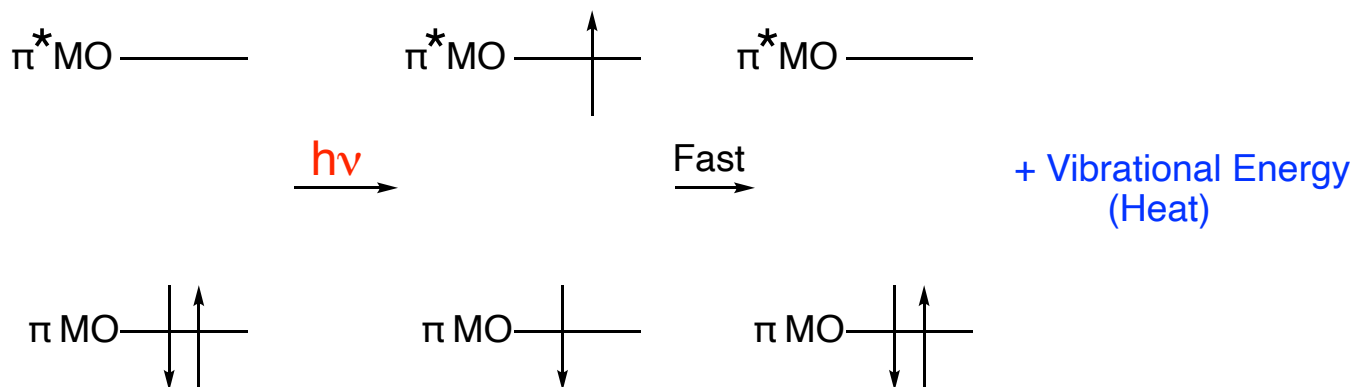
$\text{H}_2\ddot{\text{N}}\text{---}$)
↓
Binds to an -NH_2 group from the amino acid lysine in the protein opsin



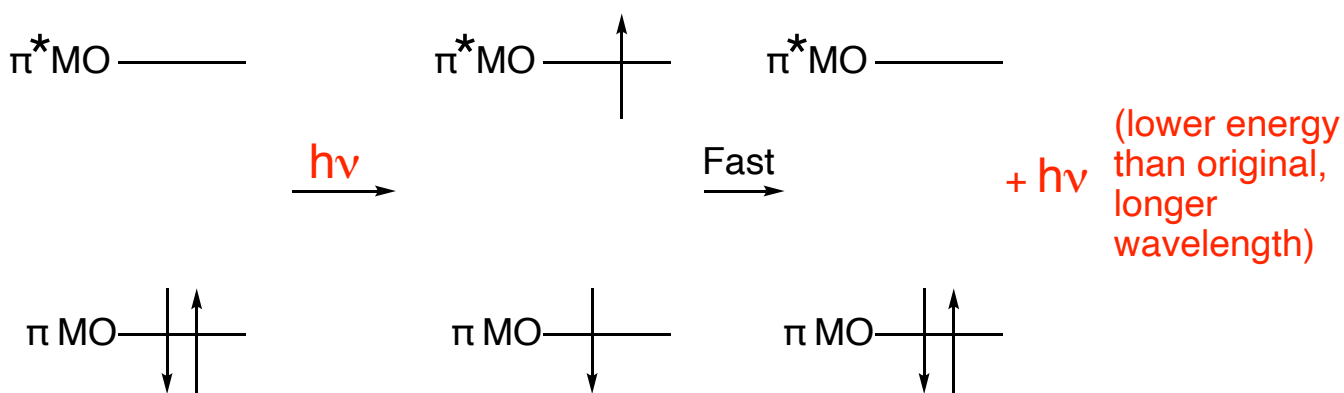
Molecule resets
↕
A photon of visible light is absorbed by the retinal, isomerizing the *cis* bond to *trans*, dramatically altering the shape of the molecule



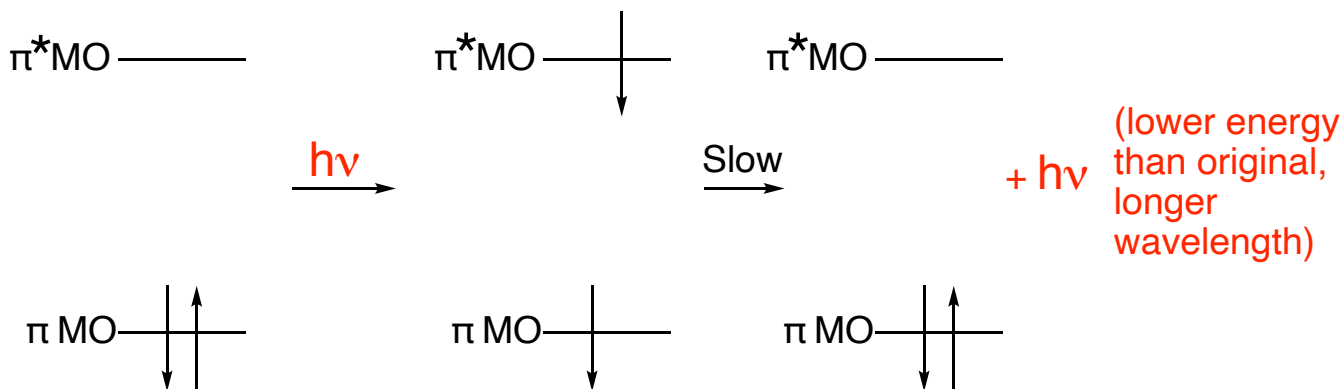
Generation of heat, Most molecules



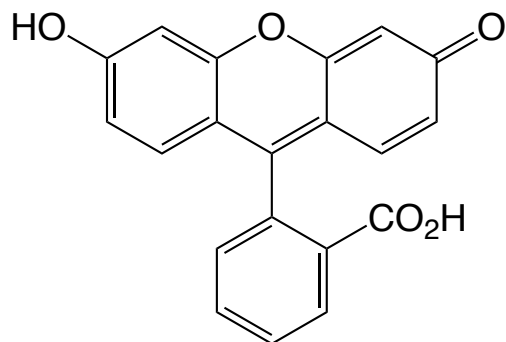
Flourescence - Rigid Molecules, Not uncommon



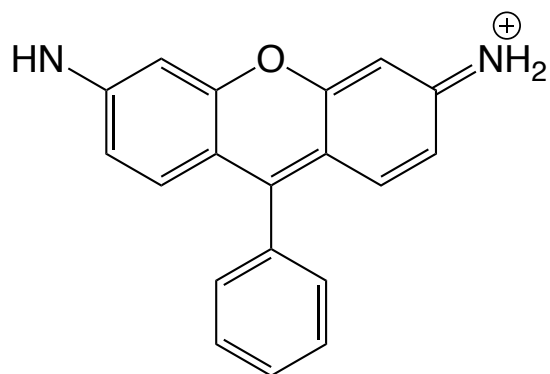
Phosphorescence - "Glow in the Dark", Rare



Flourescence - Rigid Molecules, Not uncommon

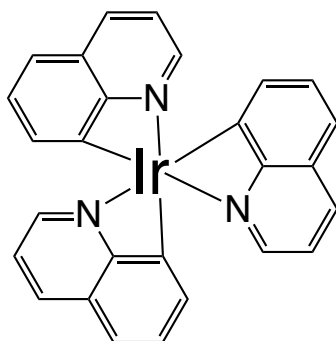


Fluorescein

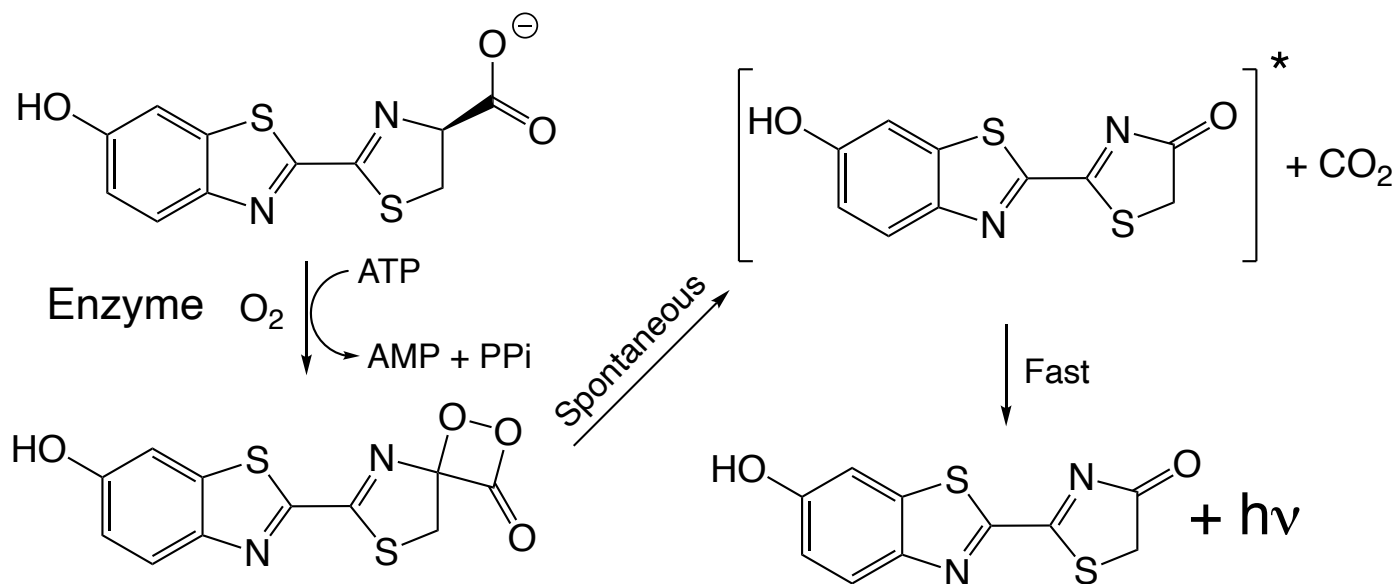


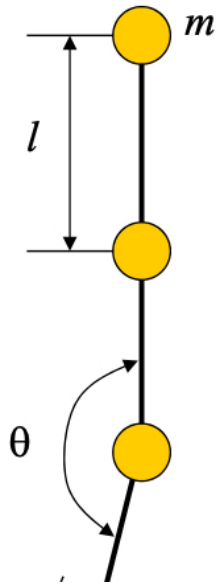
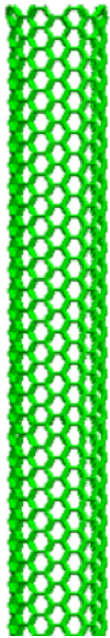
Rhodamine

Phosphorescence - "Glow in the Dark", Rare



Bioluminescence - Fireflies, Deep Sea Creatures - Chemical Reactions





Vertically
aligned CNTs

