

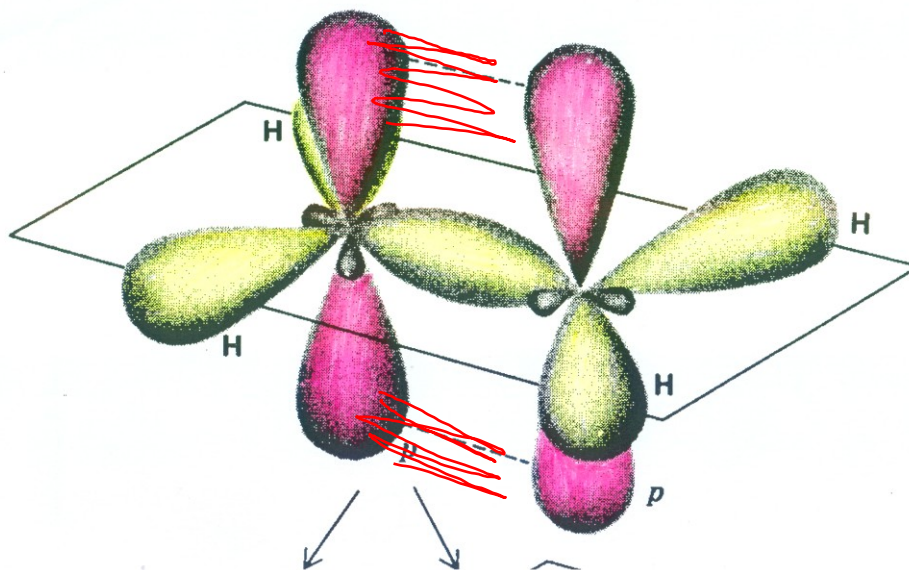
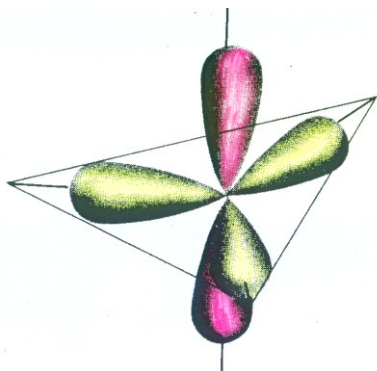
ALKENES AND CYCLOALKENES

Compounds with

C - C 0,154 nm and **C - H** bonds
C = C 0, 134 nm and **=C-H** bonds

Bonding energy

C - C 351 kJ/mol σ - bond
C = C 610 kJ/mol $\Delta = 259$ kJ/mol π - bond



ALKENES AND CYCLOALKENES

Nomenclature

$\text{H}_2\text{C}=\text{CH}_2$ ethene, ethylene

$\text{H}_2\text{C}=\text{CH}-\text{CH}_3$ propene, propylene

$\text{H}_2\text{C}=\text{CH}-\text{CH}_2-\text{CH}_3$ butene

$\text{H}_3\text{C}-\underset{\text{H}}{\text{C}}=\text{CH}-\text{CH}_3$ but-2-ene

$\text{H}_2\text{C}=\text{CH}-$

$\text{H}_2\text{C}=\text{CH}-\text{CH}_2-$

$\text{H}_3\text{C}-\text{CH}=\text{CH}-$

$\text{H}_2\text{C}=\underset{|}{\text{C}}-\text{CH}_3$

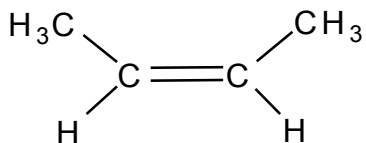
ethenyl-, **vinyl-**

prop-2-en-1-yl, **allyl-**

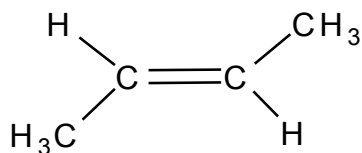
prop-1-en-1-yl

prop-1-en-2-yl

Geometric isomerie



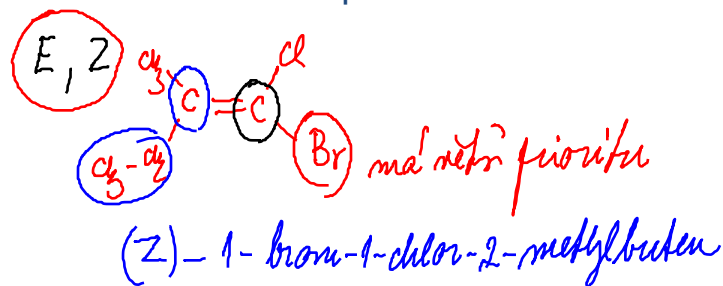
cis-but-2-ene



trans-but-2-ene

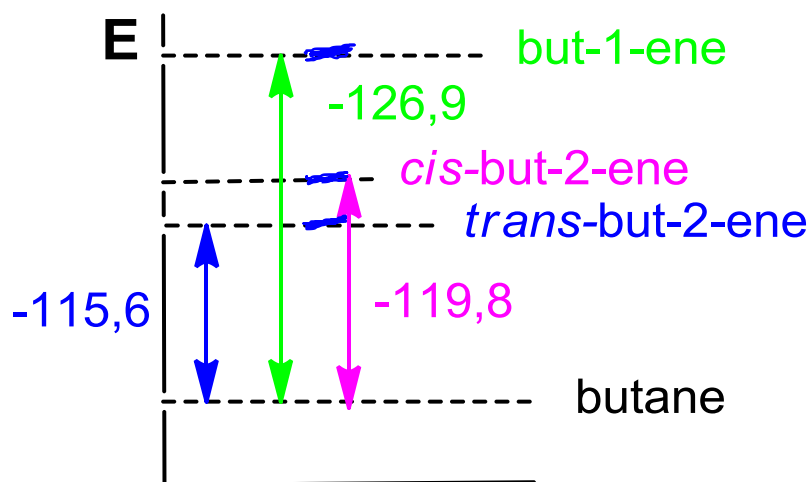
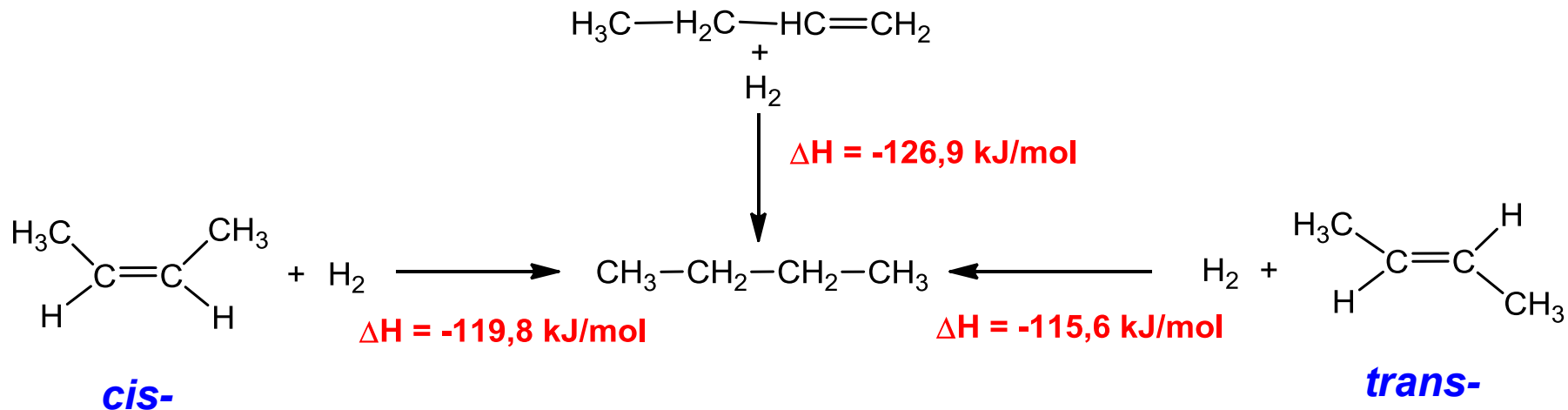
(E) (Z)

application **cis-** and **trans-** is suitable only for simple molecules



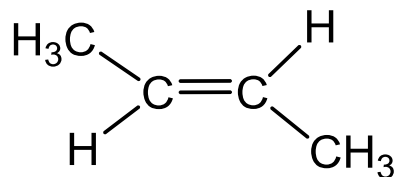
ALKENES AND CYCLOALKENES

STABILITY of alkenes

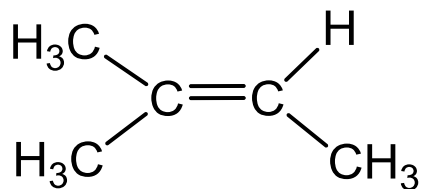


stability of *trans*-isomer is higher

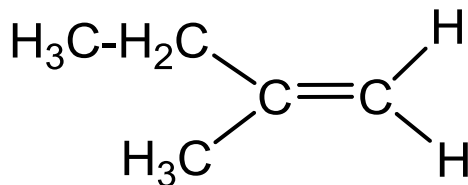
ALKENES AND CYCLOALKENES



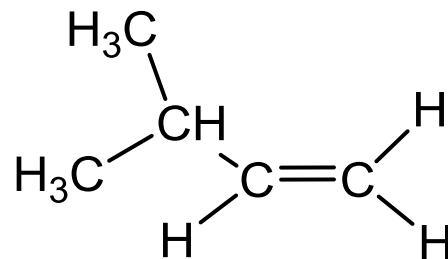
$$\Delta H = -115,6 \text{ kJ/mol}$$



$$\Delta H = -112,7 \text{ kJ/mol}$$



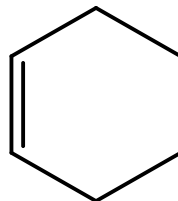
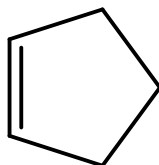
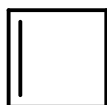
$$\Delta H = -119,4 \text{ kJ/mol}$$



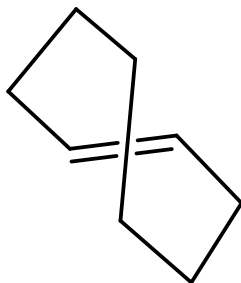
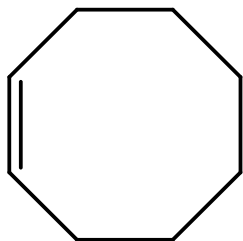
$$\Delta H = -126,9 \text{ kJ/mol}$$

the more the system at the double bond is branched, the more is stable

ALKENES AND CYCLOALKENES



all cycloalkenes with less than 6 atoms in a ring are only known as *cis*- isomers



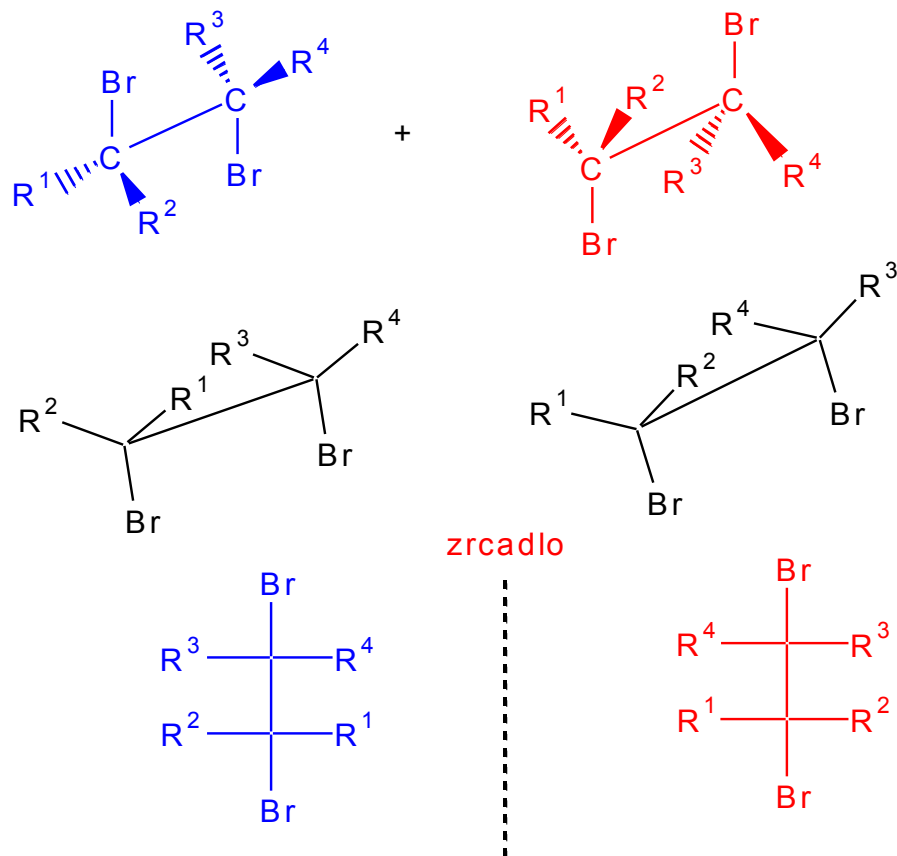
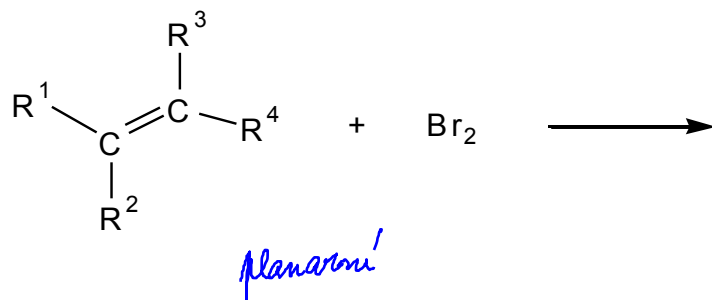
only stable is *trans*-cyclooctene

ALKENES AND CYCLOALKENES

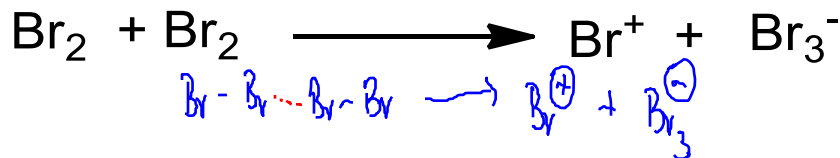
THE MAIN DIRECTION OF ALKENE REACTIVITY

ADDITION REACTION

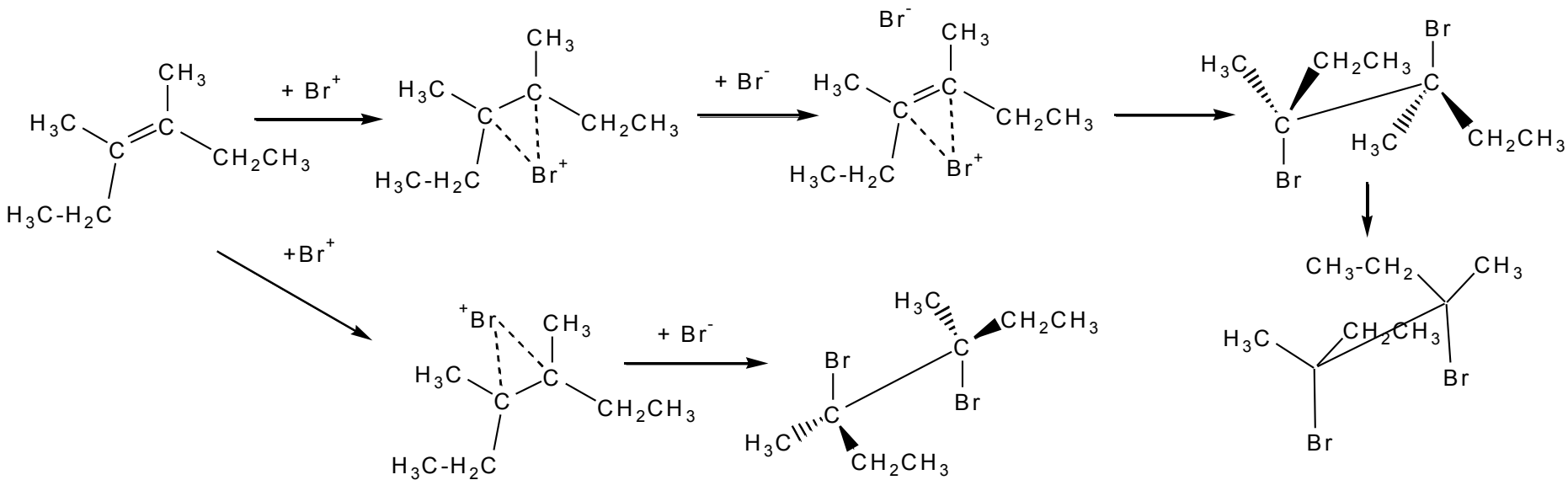
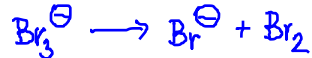
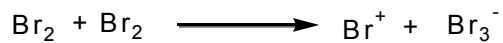
Reactions are stereospecific



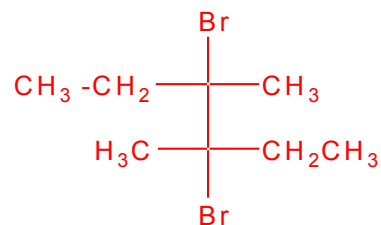
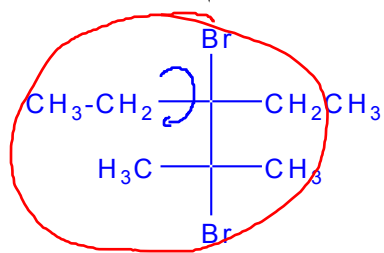
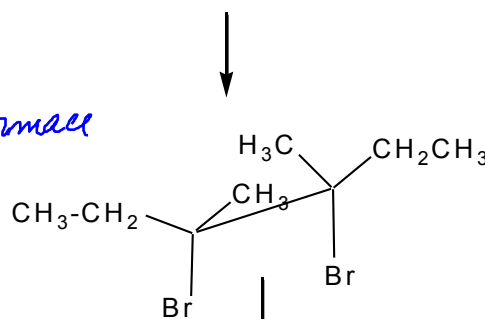
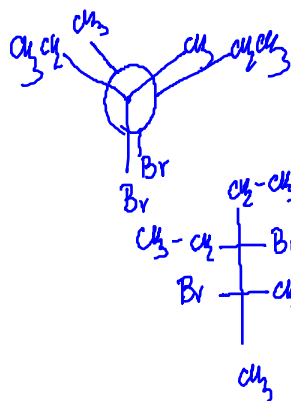
During addition of symmetrical molecule to non-symmetrically substituted olefine enantiomers are created – **reaction is stereospecific- trans- (anti)- addition**



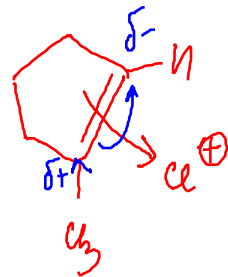
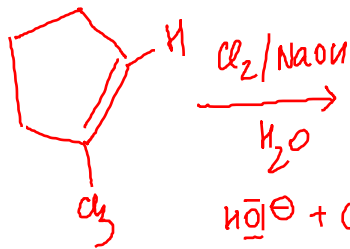
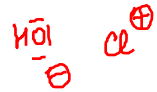
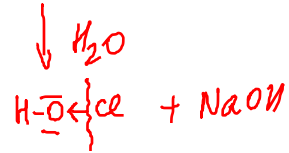
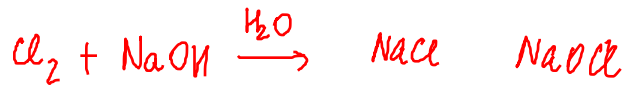
Stereospecific mechanism of additions



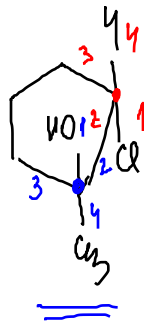
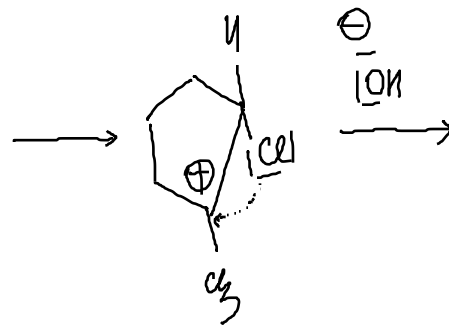
rotacyjna konformacja



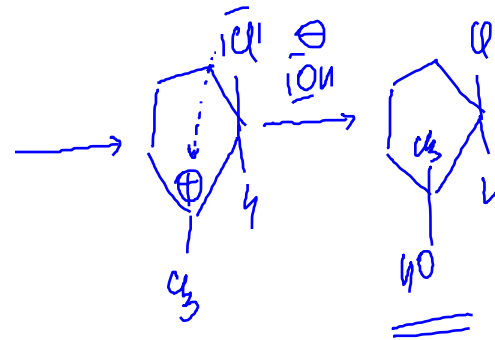
Regiospecific mechanism of additions



π -komplex



(1S, 2S) -
-chlor-1-methylcyclopentanol



Regiospecific mechanism of additions

