Recall:

Nomenclature of Alkene

- Note that there is no free rotation around C=C.

Rules:

- 1. Find longest chain
- 2. Number from end to contain both ends of C=C and give lowest number to 1st C of C=C
- 3. Change "ane" to "ene" precede with number to indicate first double bond position

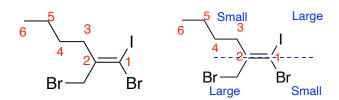
E/Z Nomenclature

E - Entegegen - Opposite

Z - Zusammen - Together

Naming based on atomic number, similar process to identifying S/R stereochemistry

Example:



Substituents: 1-iodo; 1-bromo; 2-(1-bromomethyl)

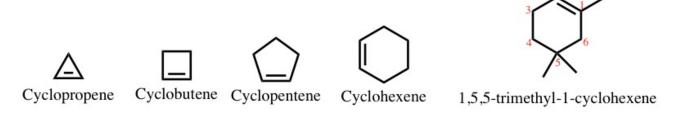
1

Parent: Hex-1-ene or 1-Hexene

Name: *E*-1-bromo-2-(1-bromomethyl)-1-iodo-1-

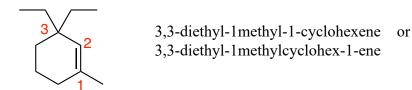
hexene

Nomenclature of Cycloalkenes



Rule: Number the cycloalkene such that the double bond is between C1 and C2 and that the first substituent has the lowest number possible.

Example:

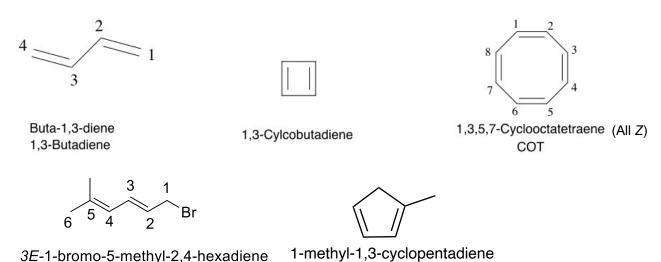


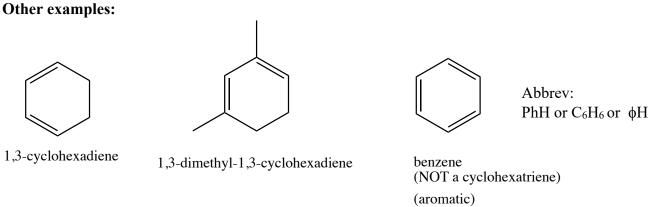
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Nomenclature of alkenes with multiple carbon-carbon double bonds (poly-enes):

Multiple C C 4 2 2 Diene 3 Drop -ne and add "diene", "triene", etc. Buta-1,3-Diene 1,3-Butadiene 1...etc

- 1) Find the longest chain containing the maximum number of double bonds.
- 2) Start numbering such that the first doubly bonded position would have the lowest number possible
- 3) Write out the full name. Number the substituents according to their position in the chain and list them alphabetically.



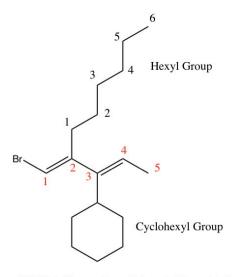


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It is therefore (E)-1-bromo-1,3,5,7-cyclooctatetraene

1,3-pentadiene (trans) = (E)-1,3-pentadiene

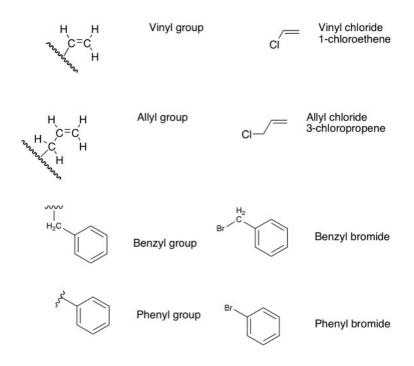
1,3-pentadiene (cis) = (Z)-1,3-pentadiene



1E,3E-1-Bromo-3-cyclohexyl-2-hexyl-1,3-pentadiene

Note: Carbons attached to double and triple bonds are depicted as additional carbon-carbon bonds in the representations above.

Special Nomenclature of Common Groups:

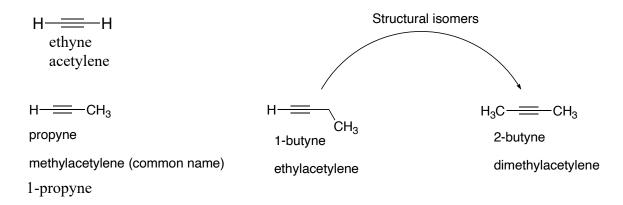


Note: phenyl bromide is commonly called bromobenzene

Nomenclature of Alkynes (also known as acetylenes)

Rules:

- Find longest chain with max number of multiple bonds
- Number from end to give 1st multiply bonded position the lowest number
- Drop "ane" and add "yne"
- For multiple triple bonds, drop "ne" and add "diyne"," triyne", etc.
- Halides and alkyl substituents take lower priority than double or triple bon



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Multiple alkynes end with:

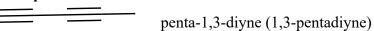
2 C \equiv C diyne

3 C≡C triyne

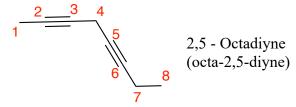
4 C C tetrayne

Mixed double and triple bond containing compounds are "eneynes."

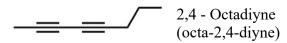
Example 1:



Example 2



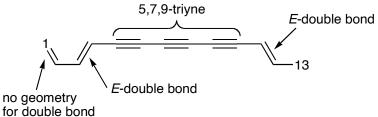
Example 3:



Example 4:

The below example is from canola – defense substance (anti-nematode)

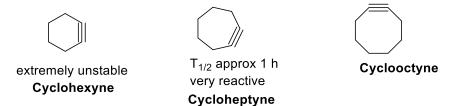
- Parent alkane of 13 carbons is tridecane hence trideca
- Start numbering the chain such that the **first multiply bonded position** gets the lowest number possible.



3*E*,11*E*-trideca-1,3,11-triene-5,7,9-triyne

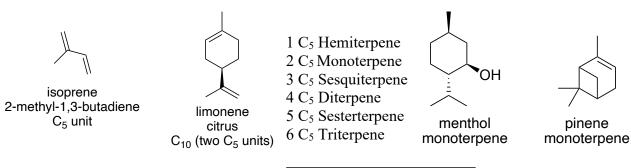
Note: alkene stereochemistry can go right to the numbers indicating positions of double bonds: **trideca-1,3***E*,11*E*-triene-5,7,9-triyn

Cyclic Structures with Triple Bonds

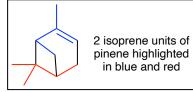


- Cycloalkynes are very reactive as the triple bonds wants to assume a 180° bond angle.
- Cycloheptyne is more stable as the angle is larger ($T_{1/2}=1$ h)
- Cyclooctyne is known to be relatively stable compound used in azide-alkyne [3+2] cycloadditions (used in biorthogonal coupling Nobel Prize 2022 Carolyne Betozzi & Barry Sharpless)

Terpenes or Isoprenoids or Terpenoids

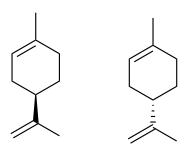


- Steroids are made from triterpenes
- -Pinene is made from two isoprene units



- Leopold Ruzicka: did significant work with terpenes

Examples:



Enantiomers of Limonene

2,6,10-trimethyldodeca-2,6,10-triene

2,6,10-trimethyl-2,6,10-dodecatriene