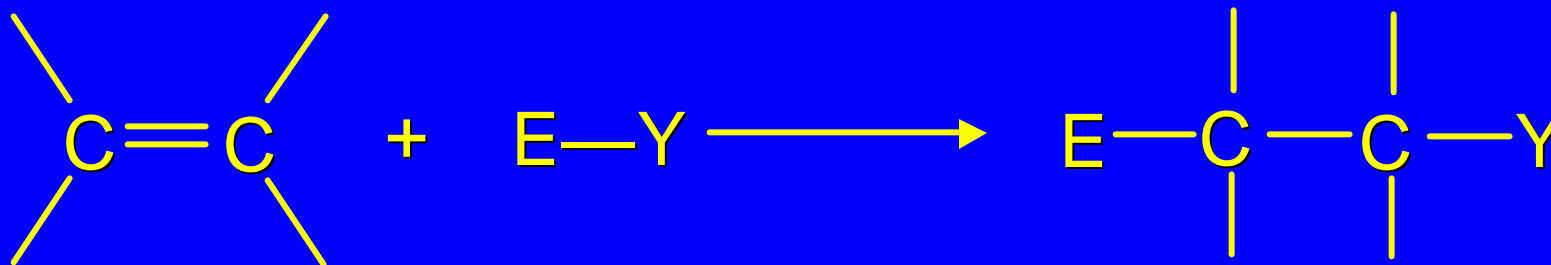


7.13

Chemical Reactions That  
Produce Diastereomers

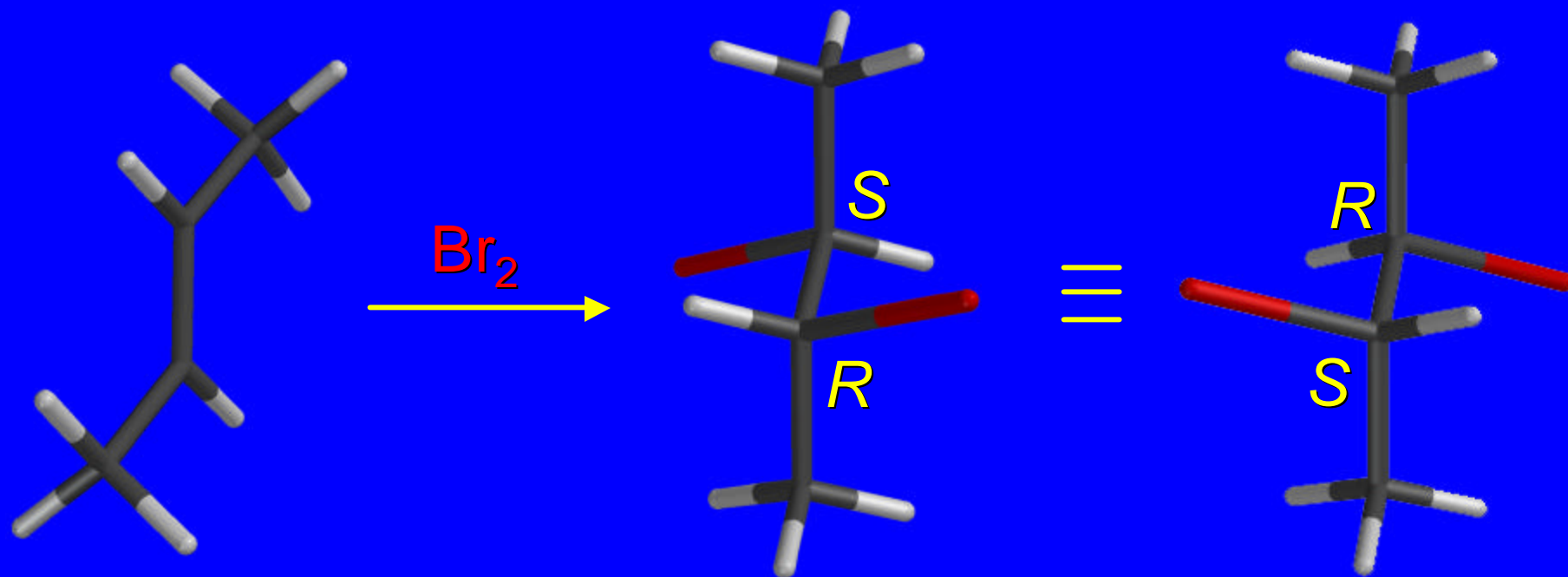
## *Stereochemistry of Addition to Alkenes*



*In order to know understand stereochemistry of product, you need to know two things:*

- (1) stereochemistry of alkene (cis or trans; Z or E)
- (2) stereochemistry of mechanism (syn or anti)

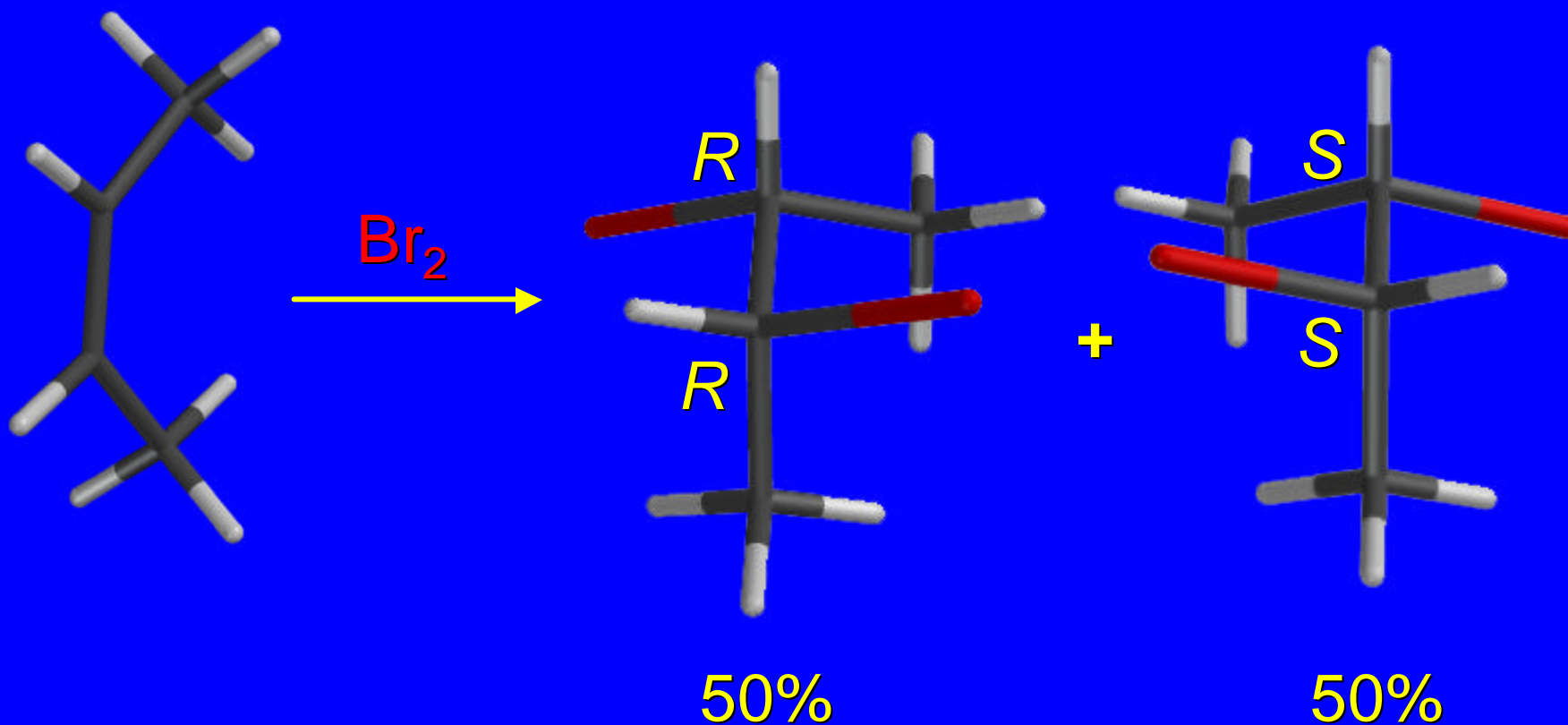
*Bromine Addition to trans-2-Butene*  
Fig. 7.14 (p 284)



meso

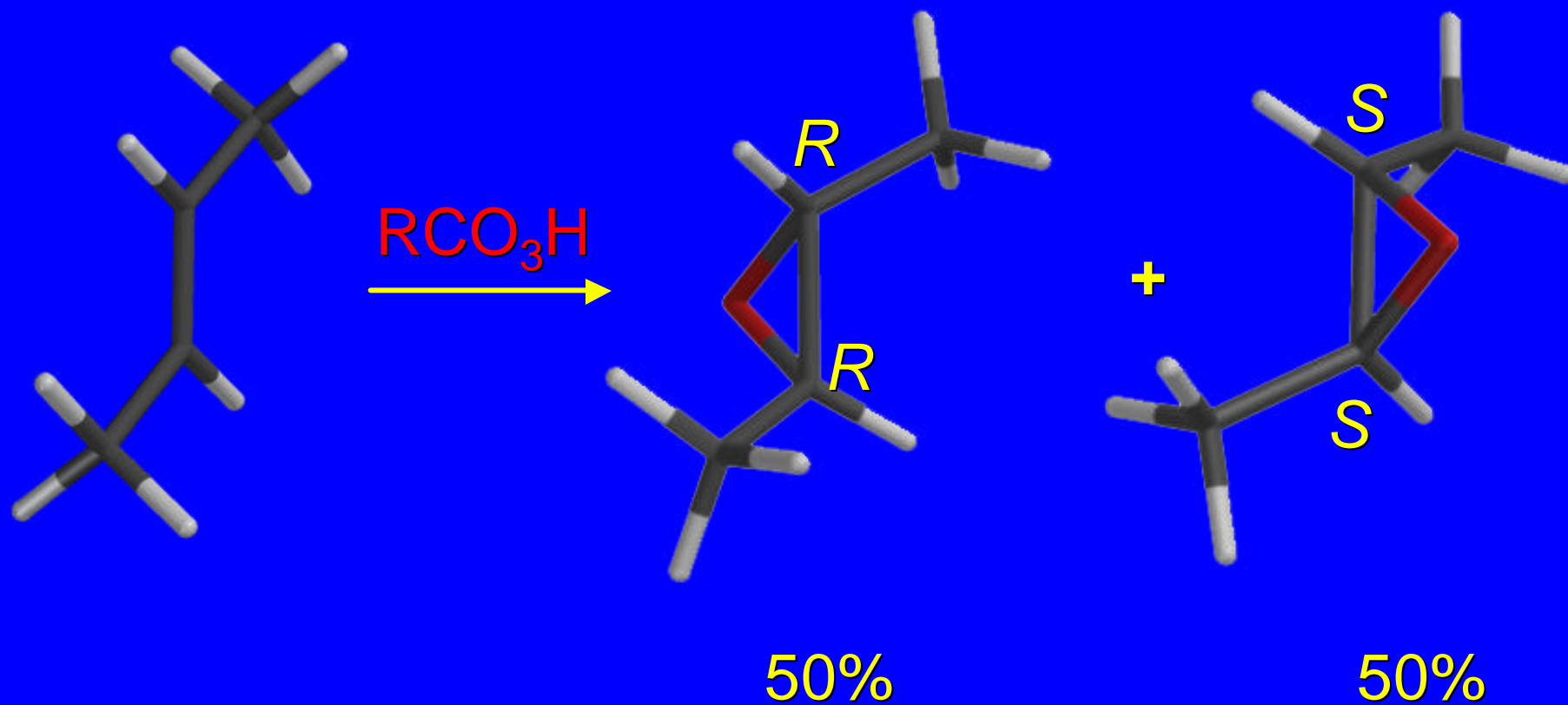
anti addition to *trans*-2-butene gives meso diastereomer

*Bromine Addition to cis-2-Butene*  
*Fig. 7.15 (p 285)*



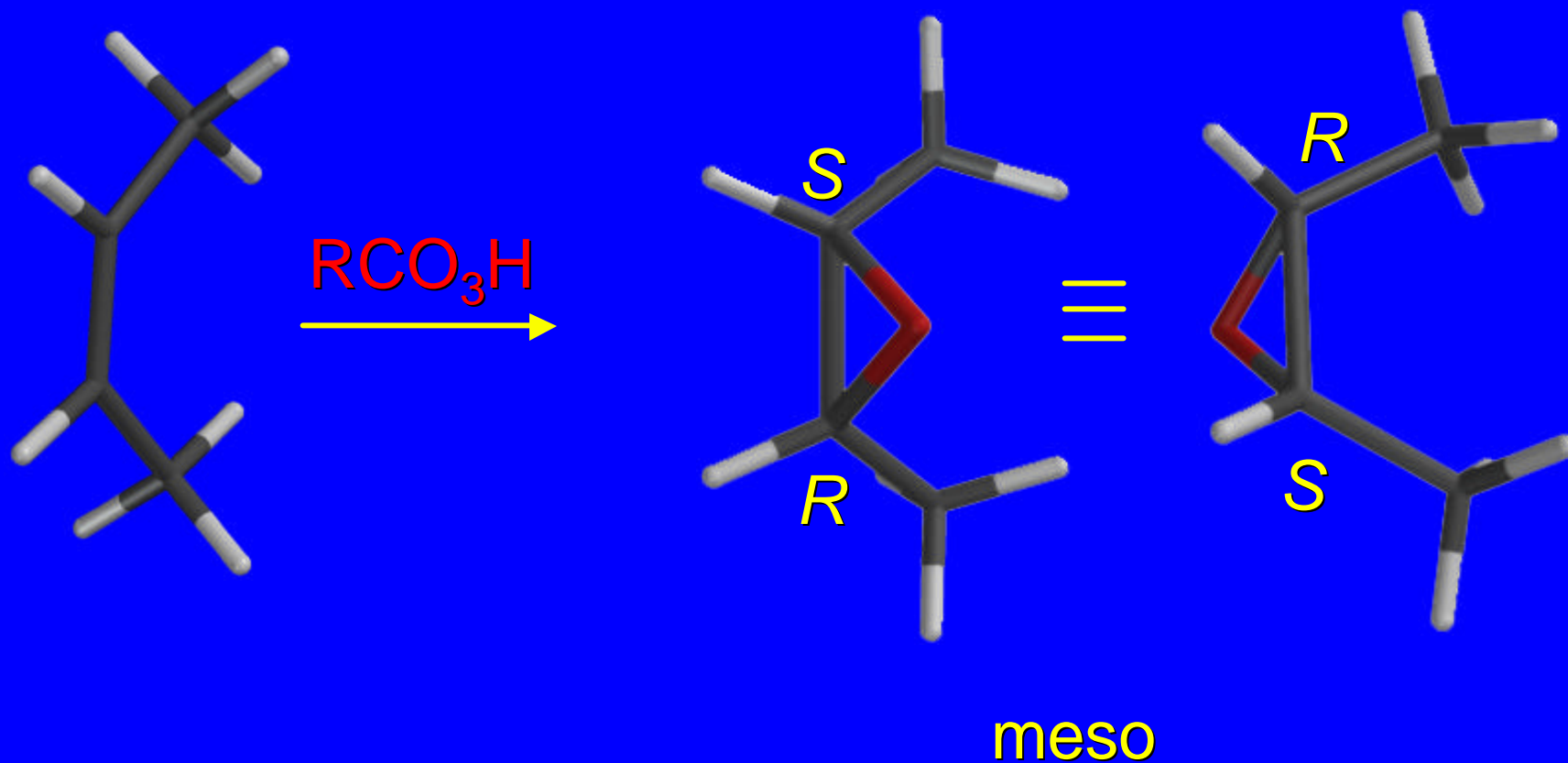
anti addition to *cis*-2-butene gives racemic mixture of chiral diastereomer

*Epoxidation of trans-2-Butene*  
*Problem 7.17 (p 285)*



syn addition to *trans*-2-butene gives racemic mixture of chiral diastereomer

*Epoxidation of cis-2-Butene*  
*Problem 7.17 (p 285)*



syn addition to *cis*-2-butene gives meso diastereomer

## *Stereospecific reaction*

of two stereoisomers of a particular starting material, each one gives different stereoisomeric forms of the product

related to mechanism: terms such as syn addition and anti addition refer to stereospecificity

*cis*-2-butene      bromination      anti      2*R*,3*R* + 2*S*,3*S*

*trans*-2-butene      bromination      anti      meso

*Stereospecific reaction*

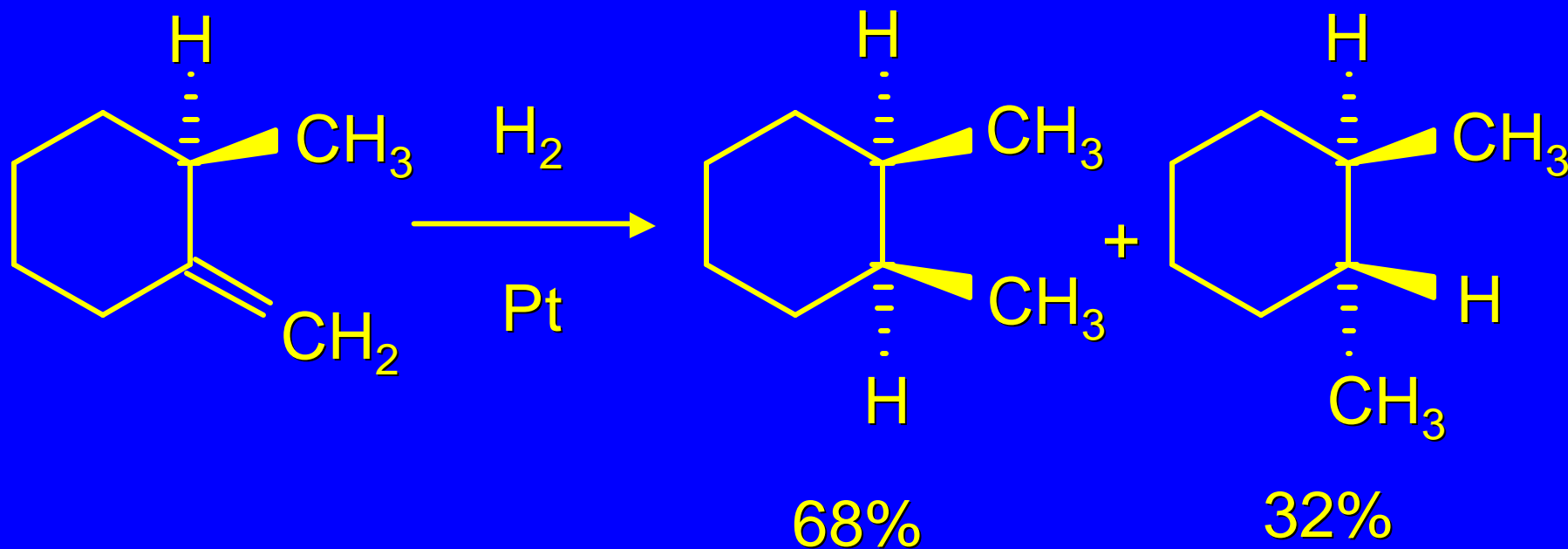
*cis*-2-butene      epoxidation      syn      meso

*trans*-2-butene      epoxidation      syn      2*R*,3*R* + 2*S*,3*S*



## *Stereoselective reaction*

a single starting material can give two or more stereoisomeric products, but gives one of them in greater amounts than any other



7.14

## Resolution of Enantiomers

separation of a racemic mixture into its two enantiomeric forms

# Strategy

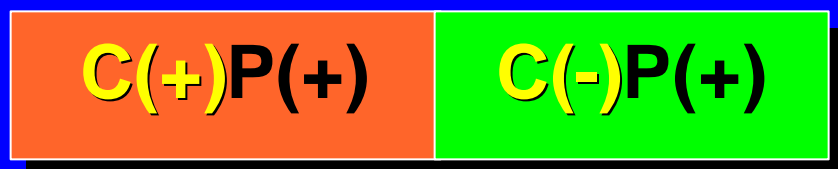
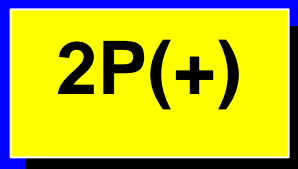
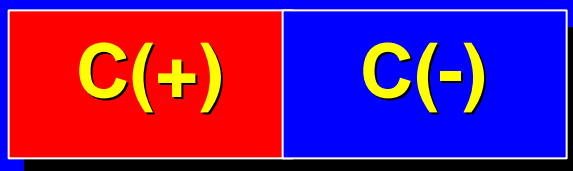
enantiomers

C(+)

C(-)

# Strategy

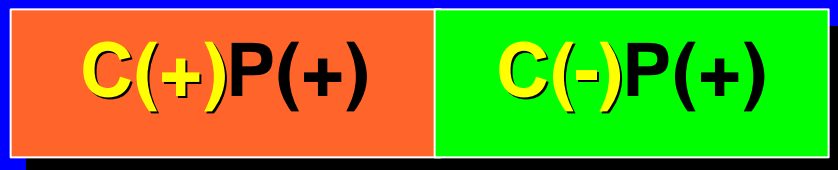
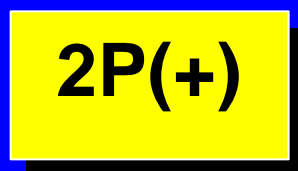
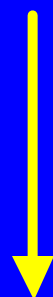
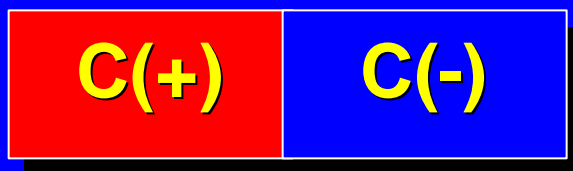
enantiomers



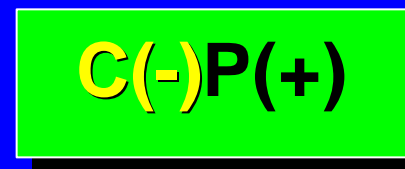
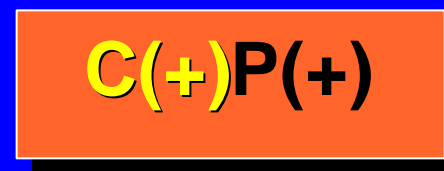
diastereomers

# Strategy

enantiomers

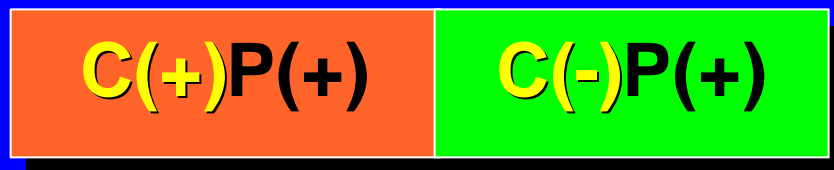
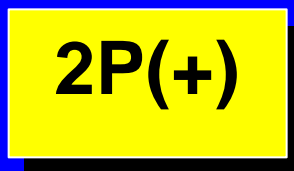
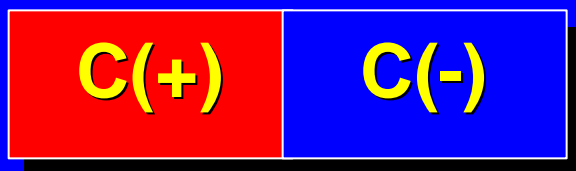


diastereomers



# Strategy

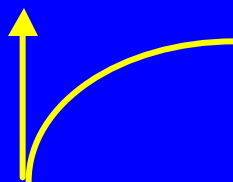
enantiomers



diastereomers

**C(+)**

**P(+)**

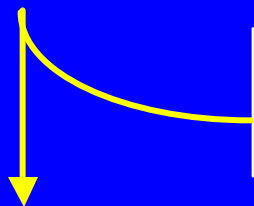
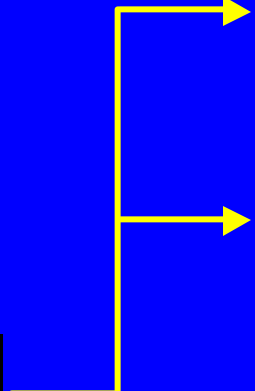


**C(+)****P(+)**

**C(-)****P(+)**

**P(+)**

**C(-)**



7.15

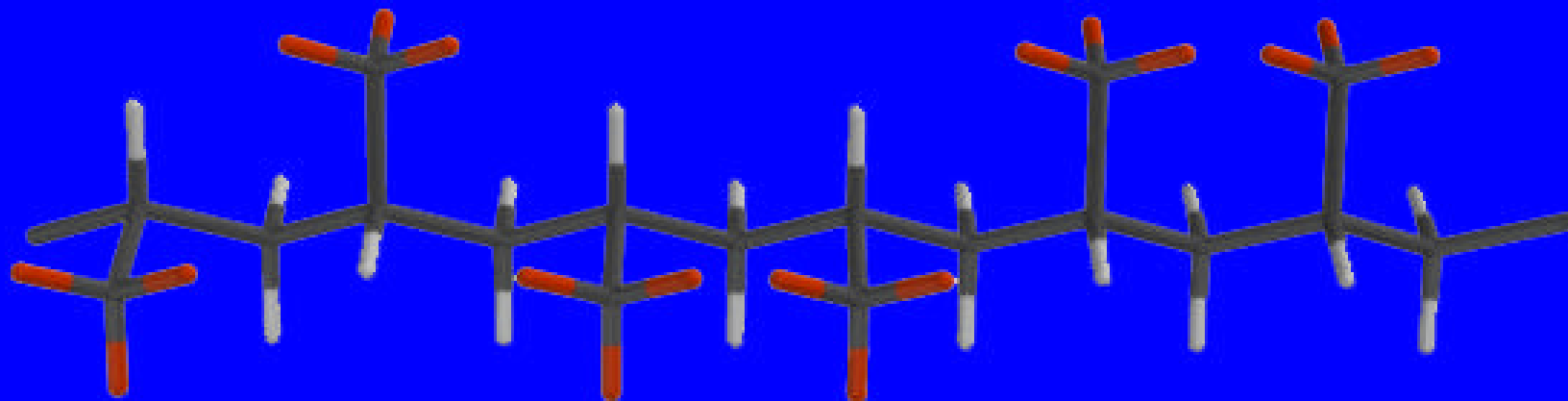
## Stereoregular Polymers

atactic

isotactic

syndiotactic

## *Atactic Polypropylene*



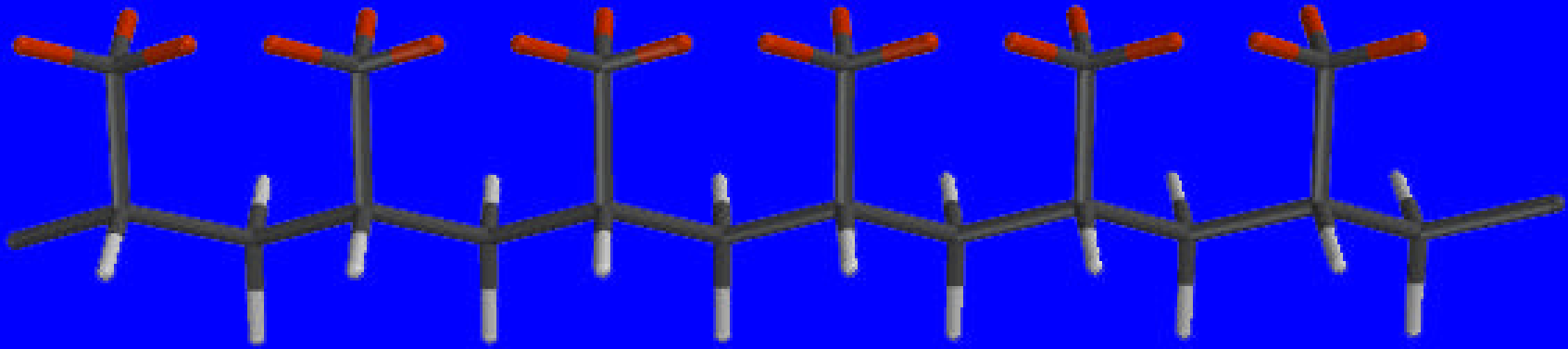
random stereochemistry of methyl groups  
attached to main chain (stereorandom)

properties not very useful for fibers etc.

formed by free-radical polymerization



## *Isotactic Polypropylene*

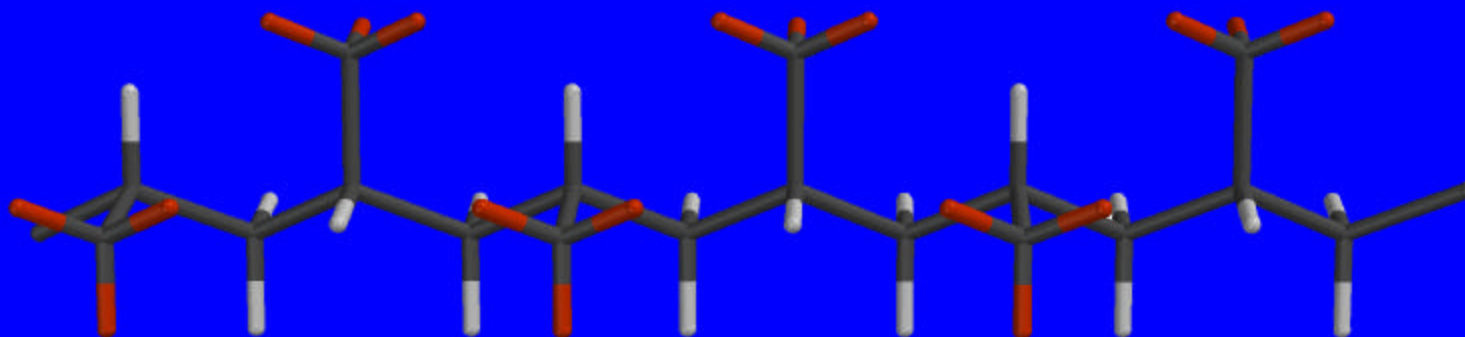


stereoregular polymer; all methyl groups on same side of main chain

useful properties

prepared by coordination polymerization under Ziegler-Natta conditions

## *Syndiotactic Polypropylene*



stereoregular polymer; methyl groups  
alternate side-to-side on main chain

useful properties

prepared by coordination polymerization  
under Ziegler-Natta conditions

7.16

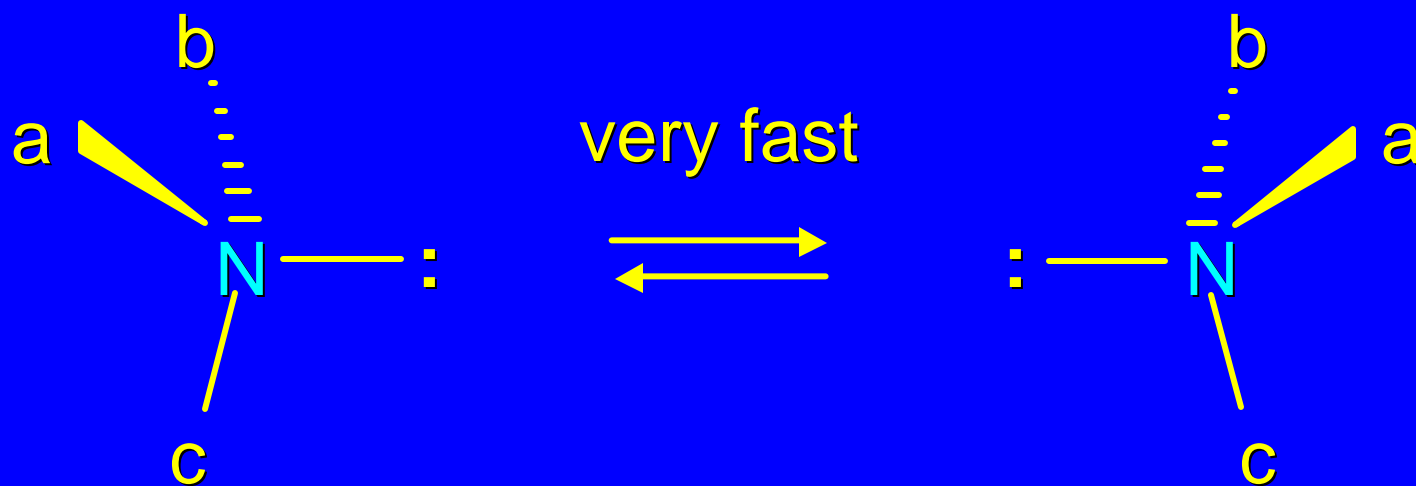
Stereogenic Centers  
Other Than Carbon

# *Silicon*



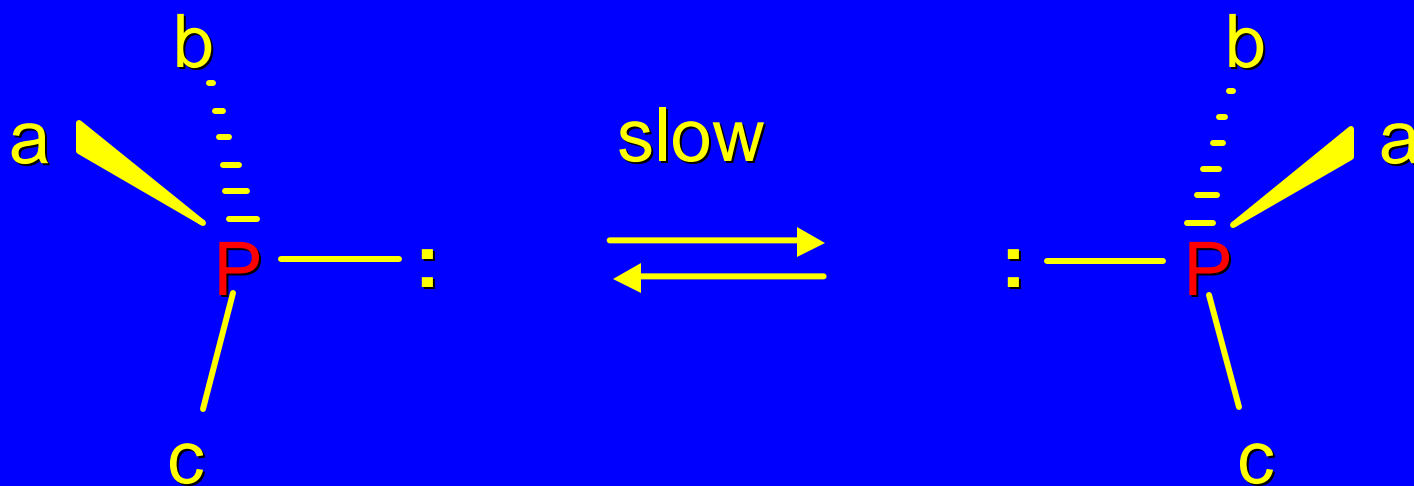
silicon, like carbon, forms four bonds in its stable compounds and many chiral silicon compounds have been resolved

## Nitrogen in amines



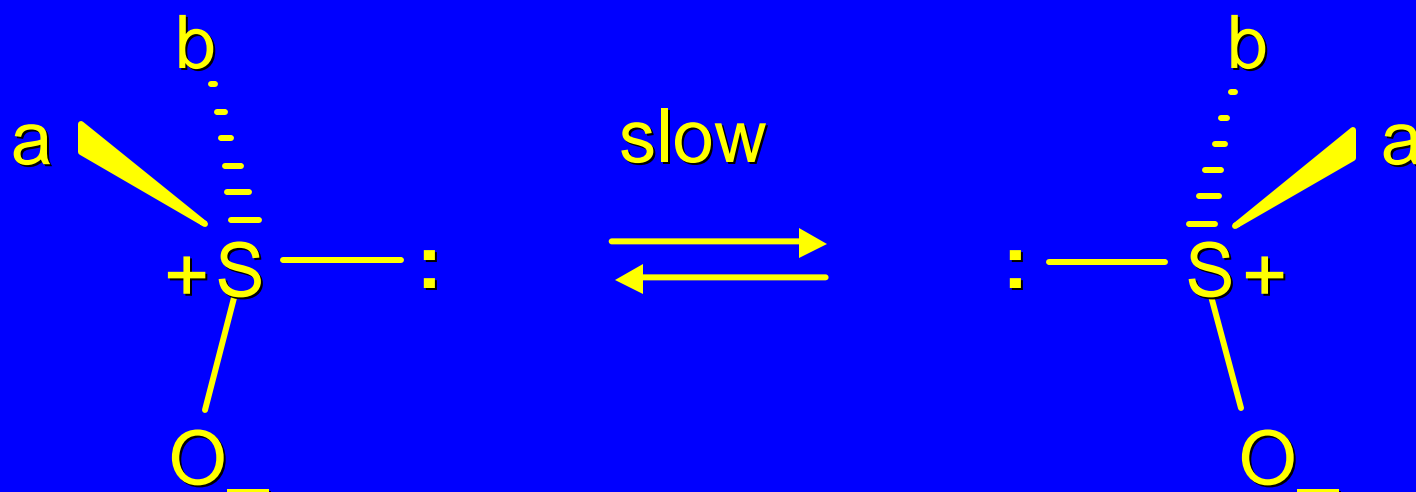
pyramidal geometry at nitrogen can produce a chiral structure, but enantiomers equilibrate too rapidly to be resolved

## *Phosphorus in phosphines*



pyramidal geometry at phosphorus can produce a chiral structure; pyramidal inversion slower than for amines and compounds of the type shown have been resolved

## Sulfur in sulfoxides



pyramidal geometry at sulfur can produce a chiral structure; pyramidal inversion is slow and compounds of the type shown have been resolved