CHEM 261 Nov 27, 2018

### **RECALL: Elimination reactions**

-OH and -OCH<sub>3</sub> are bad leaving groups and so these reactions would not occur spontaneously without an acid catalyst.

To determine whether an elimination can occur, ask yourself three questions:

$$Br \longrightarrow Br \longrightarrow Br \longrightarrow Br + KBr + HO$$

- 1. Is there a good leaving group present? Eg. Yes, Br is a good leaving group
- 2. Is there a hydrogen on the carbon next to the carbon containing the leaving group? Eg. Yes, on the bridge-heads on either side of the carbon containing the Br.
- 3. Is Bredt's Rule being followed?

  Eg. No, if a double bond was being formed, it would be at a bridge-head and Bredt's rule states that a double bond cannot be formed at a bridge-head if the rings are small and all bridges >0. (double bond too strained)

Substitution would likely not occur either as the electrophilic site is hindered (tertiary carbon).

## **Addition Reactions**

For alternate regiochemistry (addition of Br onto the less substituted carbon) need dialkyl peroxide

$$\longrightarrow \xrightarrow{\mathsf{H}-\mathsf{Br}} \xrightarrow{\mathsf{Br}} \equiv \overset{\mathsf{CH}_3 \; \mathsf{Br}}{\overset{\mathsf{CH}_3 \; \mathsf{CH}_3 \; \mathsf{Br}}}{\overset{\mathsf{CH}_3 \; \mathsf{CH}_3 \; \mathsf{$$

# Examples of peroxides

Hydrogen peroxide HO-OH

#### Radical mechanism

## Example without HBr - a polymerization reaction occurs

shorthand notation for polypropylene

Note more stable radical (in this case tetiary) is always formed by addition onto double bond

There is a termination step (not shown) that ends this polymerization. It requires combination of 2 radicals. It could be two growing chain radicals meeting or it could be from peroxide. If less peroxide is used the polymer chain will be longer.

### **Polymers**

Poly = many Meros = parts

### **Examples of Biopolymers**

- 1. Polysaccharides
  - polymers of sugars
- 2. Proteins and peptides
  - polymers of amino acids
- 3. Nucleic acid polymers (DNA and RNA)
  - polymers of nucleotides
- 4. Fats and polyketides
  - polymers of fatty acids
- 5. Polyisoprenoids/ terpenoids
  - polymers of isoprene

### **Polymer formation**

#### **Teflon**

Tetrafluoroethylene Toxic gas Teflon

Many polymers degrade into their components if heated enough, and can further decompose.

### **Polyethylene**

Ethylene

Polyethylene

## Polyacrylonitrile

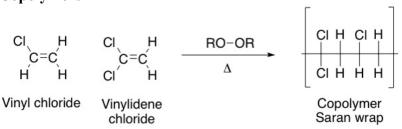
Polyacrylonitrile can form HCN if it is heated to decomposition.

## **Polyvinyl chloride**

$$\begin{array}{c|cccc}
CI & H & RO-OR & CI & H \\
C = C & & \Delta & H & H & H
\end{array}$$
Vinyl chloride
$$\begin{array}{c|cccc}
CI & H & & & \\
\hline
Polyvinyl chloride & (PVC)
\end{array}$$

Such polymers containing chloride can form HCl if decomposed.

### Copolymers



Copolymers are composed of two different subunits.

## Lucite

## **Polystyrene**

$$Ph \equiv C_6H_5 \equiv$$

phenyl group is Ph

# **Example: Mechanism of polystyrene formation**

## Short-hand for mechanism of polystyrene formation

Divinyl benzene can be added as a cross-linker so chains link on both of its double bonds. This make the copolymer more solid (as you encounter in many products) – typically about one part in 100 to one part in 6 of divinylbenzene may be added.

