### **Problem 16.18**

## **Problem 16.20**

Styrene, the simplest alkenylbenzene, is prepared commercially for use in plastics manuature by catalytic dehydrogenation of ethylbenzene. How might you prepare styrene from benzene using reactions you've studied?

Styrene

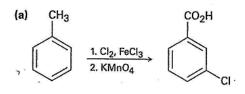
# **Problem 16.47**

Starting with either benzene or toluene, how would you synthesize the follewing substances? Assume that ortho and para isomers can be separated.

- (a) 2-Bromo-4-nitrotoluene
- (b) 1,3,5-Trinitrobenzene
- © 2,4,6-Tribromoaniline
- (d) m-Fluorobenzoic acid

#### **Problem 16.48**

As written, the following syntheses have flaws. What is wrong with east



#### **Problem 16.51**

Addition of HBr to 1-phenylpropene yields only (1-bromopropyl) benzelv. Propose a mechanism for the reaction, and explain why none of the college regioisomer is produced.

$$+ HBr \rightarrow \bigcirc$$

### **Problem 16.58**

Benzenediazonium carboxylate decomposes when heated to yield Name and a reactive substance that can't be isolated. When benzenediazonium boxylate is heated in the presence of furan, the following reaction is observed.

What intermediate is involved in this reaction? Propose a mechanism formation.

# **Problem 16.68**

How would you synthesize the following compounds from benzene? Assume that ortho and para isomers can be separated.

$$(a)$$
  $CH_3$   $(b)$   $CH_3$   $CH_2CHCH_3$   $CH_2CHCH_3$   $CH_3$ 

You know the mechanism of HBr addition to alkenes, and you know the of various substituent groups on aromatic substitution. Use this know predict which of the following two alkenes reacts faster with HBr. Explain answer by drawing resonance structures of the carbocation intermediates.

$$\begin{array}{c|c} \text{CH=CH}_2 & \text{CH=CH}_2 \\ \text{CH}_3\text{O} & \text{O}_2\text{N} \end{array}$$