

Suggested solution

Hydration of the top alkene with aqueous acid goes via a carbocation intermediate, and would therefore give the product on the left. To get the product on the right we need a more sophisticated method, involving hydroboration to place hydroxyl on the less hindered carbon and then oxidation to the alcohol (p. 444 of the textbook).



Using sodium borohydride to add two hydroxyl groups to the second alkene will give the diastereoisomer on the left, because the mechanism ensures both OH groups add to the same side of the alkene (p. 443 of the textbook). To get the trans diol, we can make an epoxide with *m*-CPBA and then open it with water, as full reaction then proceeds with inversion of configuration.



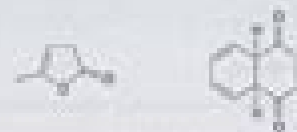
A possible alternative would be to start with the cis alkene the product will be the diol shown, which, if we culture it in an open chain conformation, is the same diol as the one that comes from the trans alkene via the epoxide.

**Suggested solutions for Chapter 20**

20

PROBLEM 1

Draw all possible enol forms of these carbonyl compounds, and comment on your stability.

**Purpose of the problem**

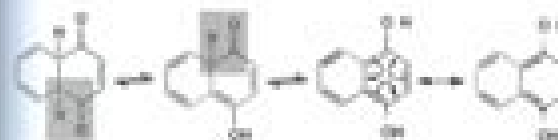
Simple exercise in drawing made with an extra twist.

Suggested solution

There is only one enol for the first compound and it might be stable because it is aromatic (two alkenes and one oxygen giving two electrons each, making six in all).



The second compound has seven possibilities, one of which is very stable indeed as it has two benzene rings. We haven't drawn the mechanism for such enolisation this time but note the different reaction arrows for tautomerisation (equilibrium) and decarboxylation (two ways of drawing the same compound).



■ Aromaticity is discussed in chapter 17 of the textbook (pages 1021–103). Compounds in which lone pairs contribute to the aromatic system are introduced on page 102, but also have two other chapters devoted to them (104 and 105). These chapters have more information than you need at this stage.

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 - The Rise of Digital Reading Clayden Organic Chemistry Solutions Manual
 - Advantages of eBooks Over Traditional Books
2. Identifying Clayden Organic Chemistry Solutions Manual
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Clayden Organic Chemistry Solutions Manual
 - User-Friendly Interface
4. Exploring eBook Recommendations from Clayden Organic Chemistry Solutions Manual

- Personalized Recommendations
 - Clayden Organic Chemistry Solutions Manual User Reviews and Ratings
 - Clayden Organic Chemistry Solutions Manual and Bestseller Lists
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 - Clayden Organic Chemistry Solutions Manual eBook Subscription Services
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 - ePub, PDF, MOBI, and More
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 - Clayden Organic Chemistry Solutions Manual Enhanced eBook Features
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 - Highlighting and Note-Taking Clayden Organic Chemistry Solutions Manual
 - Interactive Elements Clayden Organic Chemistry Solutions Manual
 8. Staying Engaged with Clayden Organic Chemistry Solutions Manual
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 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Clayden Organic Chemistry Solutions Manual
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 - Benefits of a Digital Library
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 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
 11. Cultivating a Reading Routine Clayden Organic Chemistry Solutions Manual
 - Setting Reading Goals Clayden Organic Chemistry Solutions Manual
 - Carving Out Dedicated Reading Time
 12. Sourcing Reliable Information of Clayden Organic Chemistry Solutions Manual

- Fact-Checking eBook Content of Clayden Organic Chemistry Solutions Manual
- Distinguishing Credible Sources

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- Utilizing eBooks for Skill Development
- Exploring Educational eBooks

14. Embracing eBook Trends

- Integration of Multimedia Elements
- Interactive and Gamified eBooks

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