Over reaction of Aniline and Phenol

The strongest activating and ortho/para-directing substituents are the amino (-NH$_2$) and hydroxyl (-OH) groups. Direct nitration of phenol (hydroxybenzene) by dilute nitric acid gives modest yields of nitrated phenols and considerable oxidative decomposition to tarry materials; aniline (aminobenzene) is largely destroyed. Bromination of both phenol and aniline is difficult to control, with di- and tri-bromo products forming readily. Because of their high nucleophilic reactivity, aniline and phenol undergo substitution reactions with iodine, a halogen that is normally unreactive with benzene derivatives. The mixed halogen iodine chloride (ICl) provides a more electrophilic iodine moiety, and is effective in iodinating aromatic rings having less powerful activating substituents.

\[
C_6H_5–NH_2 + I_2 + NaHCO_3 \rightarrow p-I-C_6H_4–NH_2 + NaI + CO_2 + H_2O
\]

By acetylating the heteroatom substituent on phenol and aniline, its activating influence can be substantially attenuated. For example, acetylation of aniline gives acetanilide (first step in the following equation), which undergoes nitration at low temperature, yielding the para-nitro product in high yield. The modifying acetyl group can then be removed by acid-catalyzed hydrolysis (last step), to yield para-nitroaniline. Although the activating influence of the amino group has been reduced by this procedure, the acetyl derivative remains an ortho/para-directing and activating substituent.

\[
C_6H_5–NH_2 + (CH_3CO)_2O \xrightarrow{\text{pyridine (a base)}} C_6H_5–NHCOCH_3 \xrightarrow{\text{HNO}_3, 5^\circ C} p-O_2N–C_6H_4–NHCOCH_3 \xrightarrow{\text{H}_3O^+ & \text{heat}} p-O_2N–C_6H_4
\]

The following diagram illustrates how the acetyl group acts to attenuate the overall electron donating character of oxygen and nitrogen. The non-bonding valence electron pairs that are responsible for the high reactivity of these compounds (blue arrows) are diverted to the adjacent carbonyl group (green arrows). However, the overall influence of the modified substituent is still activating and ortho/para-directing.
Objectives

After completing this section, you should be able to

1. explain why phenols and phenoxide ions are very reactive towards electrophilic aromatic substitution (see Section 16.4 of the textbook).
2. write an equation to illustrate the oxidation of a phenol or an arylamine to a quinone, and identify the reagents used to oxidize phenols.
3. write an equation to illustrate the reduction of a quinone to a hydroquinone, and identify the reagents used to reduce quinones.
4. describe, briefly, the biological importance of the redox properties of quinones.

Key Terms

Make certain that you can define, and use in context, the key terms below.

- hydroquinone
- quinone
- ubiquinone

Study Notes

“Quinone” is a term used to describe cyclohexadienones in general, and \( p \)-benzoquinone in particular. In addition to benzene, other aromatic systems also give rise to quinones; for example, 1,4-naphthoquinone
“Hydroquinones” are produced by the reduction of quinones according to the following half-reaction:

\[
2H^+ + 2e^- + \text{quinone} \rightleftharpoons \text{hydroquinone}
\]

“Ubiquinones” are naturally occurring quinones whose role is to transfer a pair of electrons from one substance to another in enzyme-catalyzed reactions. Ubiquinones are also called coenzymes Q.

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**Electrophilic Aromatic Substitution Reactions**

The facility with which the aromatic ring of phenols and phenol ethers undergoes electrophilic substitution has been noted. Two examples are shown in the following diagram. The first shows the Friedel-Crafts synthesis of the food preservative BHT from para-cresol. The second reaction is interesting in that it further demonstrates the delocalization of charge that occurs in the phenolate anion. Carbon dioxide is a weak electrophile and normally does not react with aromatic compounds; however, the negative charge concentration on the phenolate ring enables the carboxylation reaction shown in the second step. The sodium salt of salicylic acid is the major product, and the preference for ortho substitution may reflect the influence of the sodium cation. This is called the Kolbe-Schmidt reaction, and it has served in the preparation of aspirin, as the last step illustrates.

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**Oxidation of Phenols: Quinones**

Phenols are rather easily oxidized despite the absence of a hydrogen atom on the hydroxyl bearing carbon. Among the colored products from the oxidation of phenol by chromic acid is the dicarbonyl compound para-benzoquinone (also known as 1,4-benzoquinone or simply quinone); an ortho isomer is also known. These compounds are easily reduced to their dihydroxybenzene analogs, and it is from these compounds that quinones are best prepared. Note that meta-quinones having similar structures do not exist. The redox equilibria between the dihydroxybenzenes hydroquinone and catechol and their quinone oxidation states are so facile that milder oxidants than chromate (Jones reagent) are generally preferred.
One such oxidant is **Fremy's salt**, shown on the right. Reducing agents other than stannous chloride (e.g. NaBH$_4$) may be used for the reverse reaction. The position of the quinone-hydroquinone redox equilibrium is proportional to the square of the hydrogen ion concentration, as shown by the following half-reactions (electrons are colored blue). The electrode potential for this interconversion may therefore be used to measure the pH of solutions.

\[
\text{Quinone} + 2\text{H}^+ \leftrightarrow 2\text{e}^- \quad \text{Hydroquinone}
\]

**Exercises**

**Questions**

**Q17.10.1**

Predict the major product if the following reagents/reagents were used. No reaction is also a possible answer.

(a) 1 equivalent of PBr$_3$
(b) 1 equivalent of SOCl$_2$
(c) Dess–Martin periodinane
(d) 3 equivalents of acetyl chloride and AlCl$_3$ as a catalyst

(e) Heat and H$_2$SO$_4$ (assume the phenol does not act as a nucleophile in this case)

**Q17.10.2**
Predict the major product if the following reagents/conditions were used. No reaction is also a possible answer.

(a) 2 equivalents of RMgBr and H$_3$O$^+$ work-up
(b) LiAlH$_4$ and H$_3$O$^+$ work-up
(c) NaBH$_4$ and H$_3$O$^+$ work-up

**Solutions**

S17.10.1

(a)  
(b)  
(c)  
(d)  
(e)  
(a)

(b)

(c) No reaction. NaBH$_4$ is milder oxidant than LiAlH$_4$. It typically only reduces ketones and aldehydes.

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**Contributors**

- Dr. Dietmar Kennepohl, FCIC (Professor of Chemistry, Athabasca University)
- Prof. Steven Farmer (Sonoma State University)