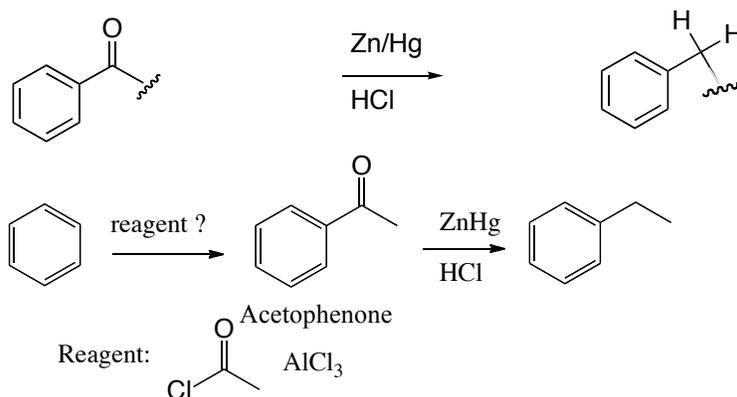
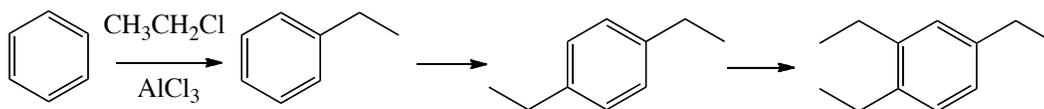


Clemmensen Side Chain Reduction Reaction

The Clemmensen reduction is an example of a reduction of a conjugated ketone to the alkyl chain using a zinc/mercury amalgam and an acid as shown below.

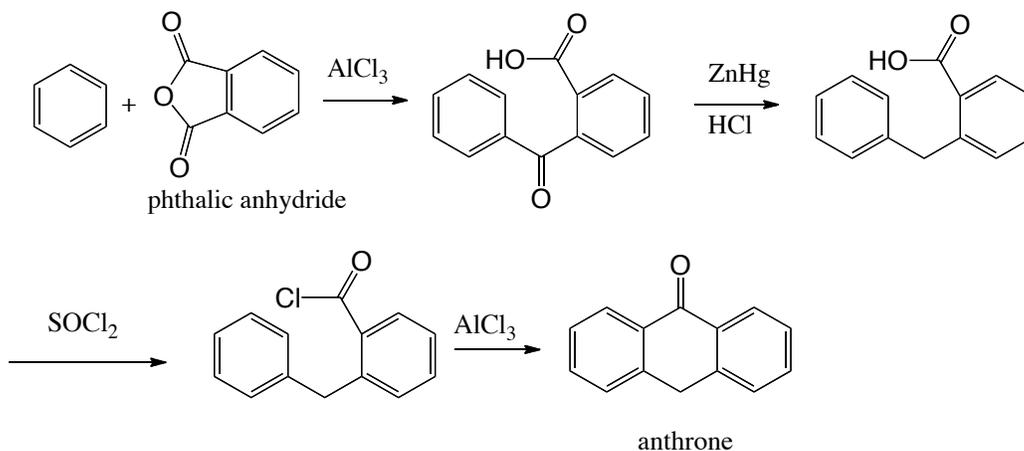


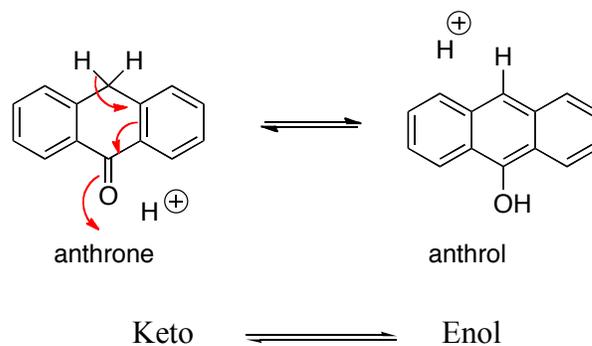
It may seem simpler to make ethyl benzene with ethyl chloride, BUT....



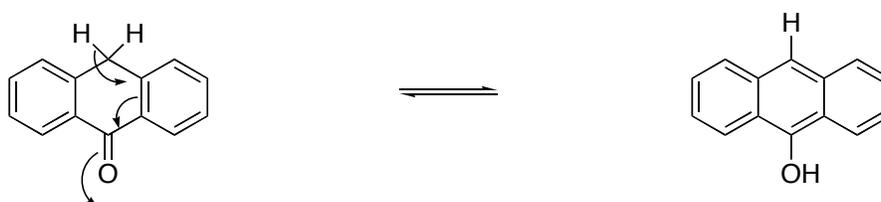
ethyl benzene is more reactive than benzene (inductive effect): reacts **faster**
acetophenone is less reactive than benzene (resonance effect): reacts **slower**

More examples:





Anthrone undergoes a keto to enol isomerization to anthrol, which is a fluorescent dye that absorbs UV light and emits in the visible (remember the demo in class).

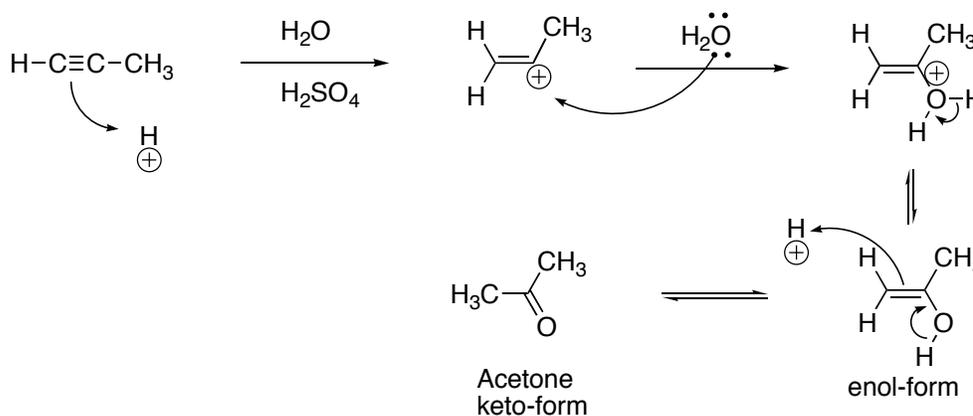


Anthrone (keto form)

Anthrol (enol form)

Anthrone and Anthrol are tautomers (interconverting structural isomers).

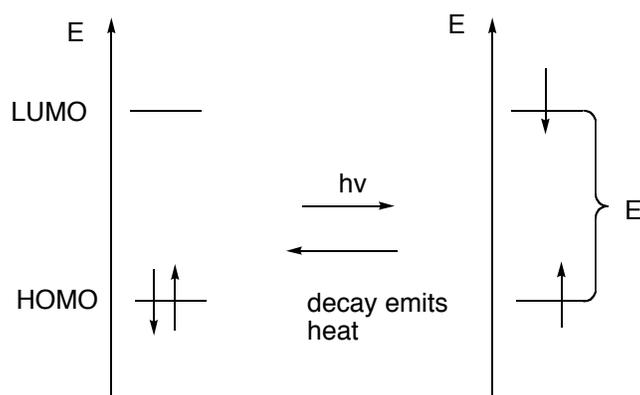
Review tautomerization from CHEM 261



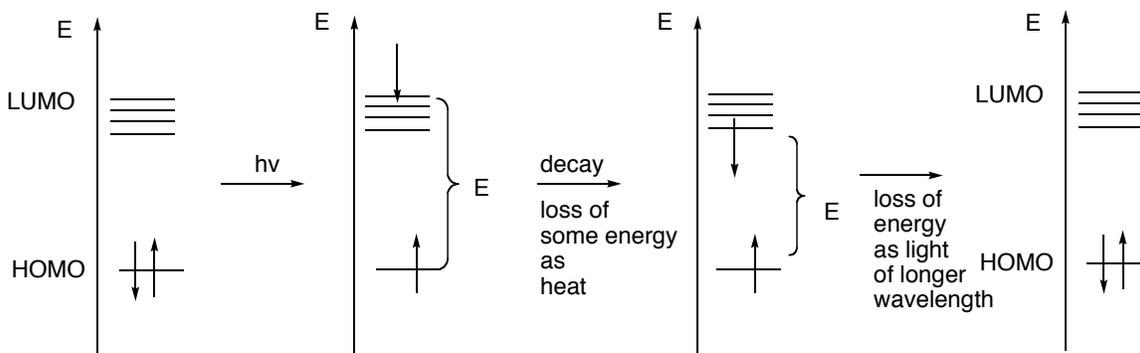
Fluorescence digression

Fluorescence is the emission of electromagnetic energy from an excited state (obtained by irradiation, normally in the ultraviolet) as light (usually visible wavelengths).

Commonly, electromagnetic radiation (light) is absorbed if it matches the difference between the highest occupied molecular orbital (HOMO) and lowest unoccupied molecular orbital (LUMO). The promoted electron in the LUMO then decays back to the HOMO with loss of the energy as heat (molecular motion, collision with other molecules).



However, if a conjugated molecule absorbs a high-energy photon into a higher vibrational state of the LUMO or a higher unoccupied MO and the electron goes in to the lowest vibrational excited state of the LUMO (with the loss of energy as heat), the remaining energy to return to the ground state can be emitted as a lower-energy photon with a longer wavelength (visible).



There are many natural conjugated molecules that exhibit fluorescence, for example, green fluorescence protein (GFP).

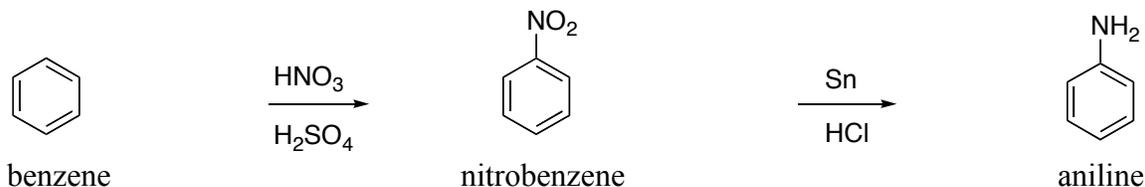
Phosphorescence also occurs for some natural compounds if the electrons become unpaired. Phosphorescence decay is slower than fluorescence, so it lasts longer than fluorescence; it can be seen in novelty items such as 'glow in the dark' stars that can be stuck to the wall.

Chemiluminescence is the result of a reaction that emits energy as light

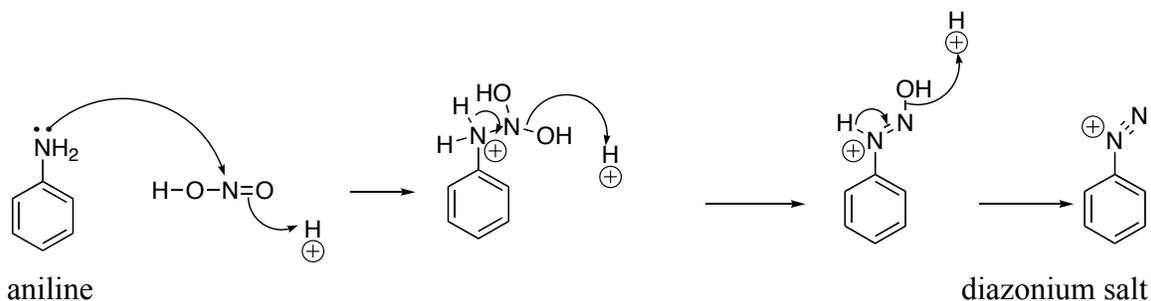
Nitroreduction and Diazotization

Example:

For example, aniline can be formed from benzene by nitration followed by reduction of the nitro group. The first step involves the nitration of benzene to form nitrobenzene. The next step uses tin (Sn) under acidic conditions to reduce the nitro group to an amine and form aniline.

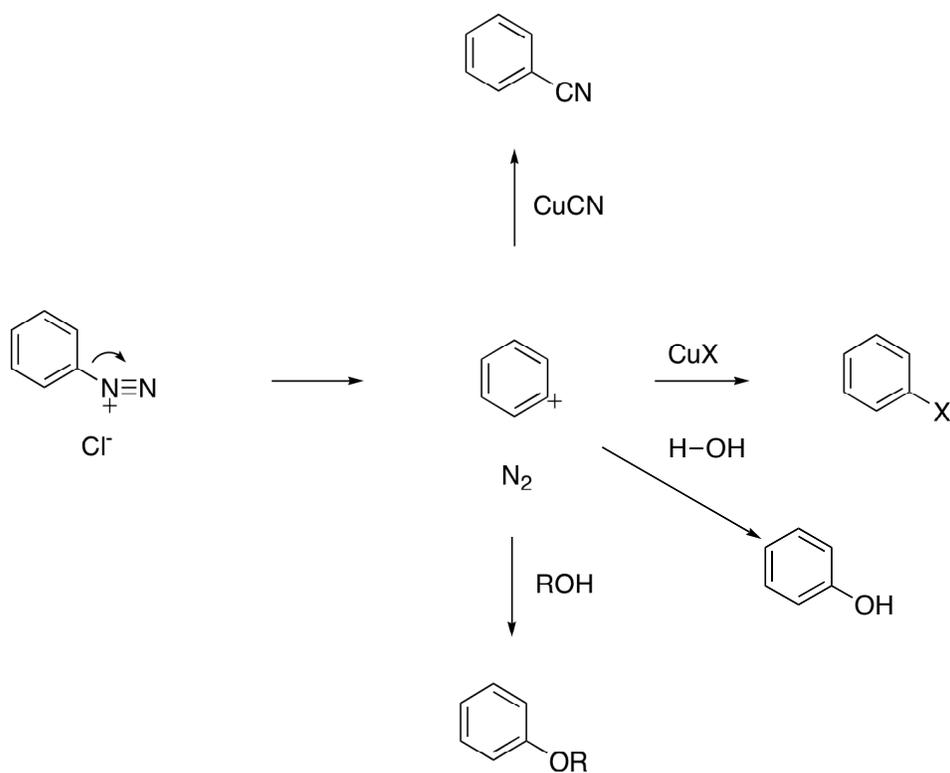


The aniline can then be reacted with nitrous acid to give an explosive diazonium salt. Nitrous acid (HONO) is formed from the addition of sodium nitrite (NaNO_2) and hydrochloric acid (HCl). In a very abbreviated mechanism, the first step of the mechanism is the attack of aniline on nitrous acid to form the new nitrogen-nitrogen bond. This is followed by deprotonation and reprotonation to make the nitrogen-nitrogen double bond. The loss of water then forms the diazonium species. The chloride counterion is from the initial formation of nitrous acid with HCl.

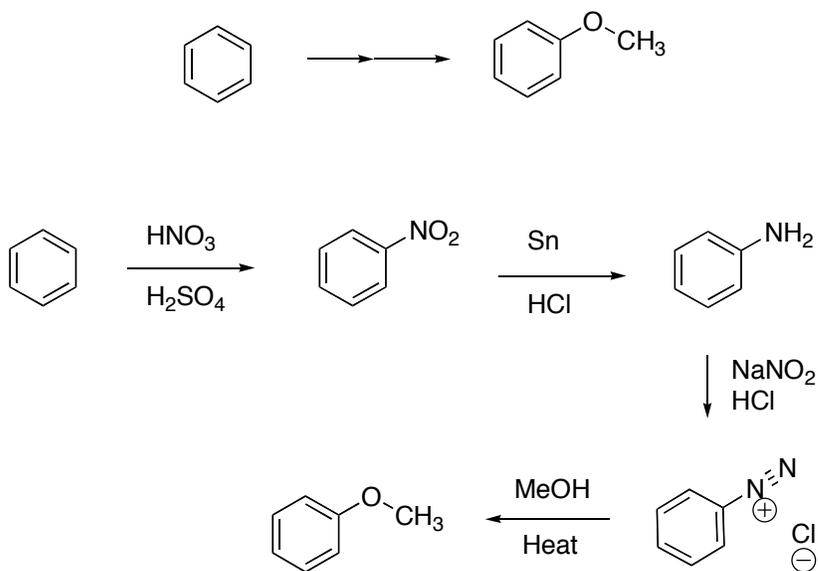


Replacement of N_2 of Diazonium salts

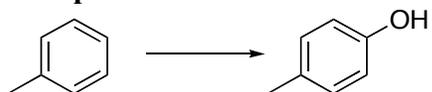
Diazonium salts are extremely useful in organic synthesis, because the diazonium group ($-\text{N}_2$) can be replaced by nucleophiles. The diazonium salt behaves like a phenyl cation and can undergo a variety of nucleophilic aromatic substitutions such as the addition of copper cyanide (CuCN) to give an aromatic cyanide, addition of CuX (where X is a halogen) to give a halogenated benzene, or addition of alcohols to give ethers. Formally these reactions proceed via $\text{S}_{\text{N}}1$ mechanism.

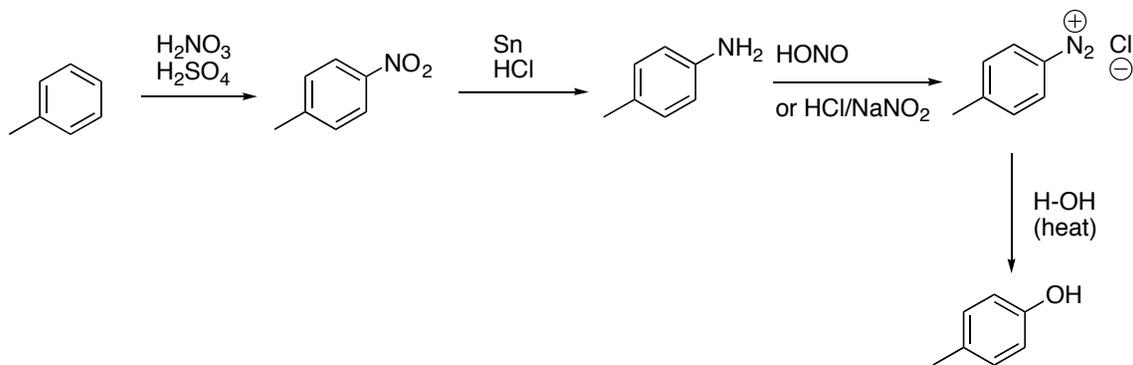


Example: make anisole (methoxybenzene)



Example:

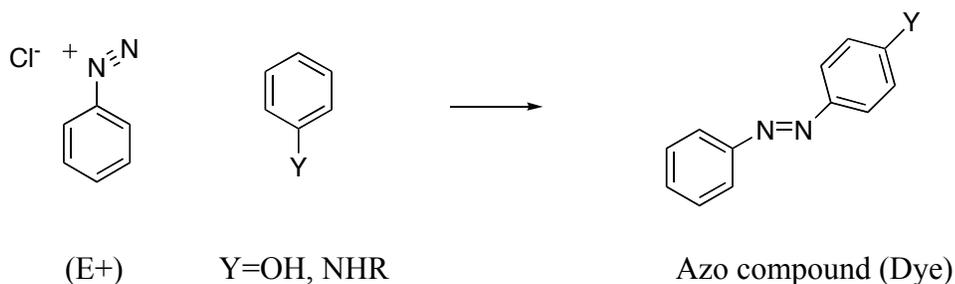




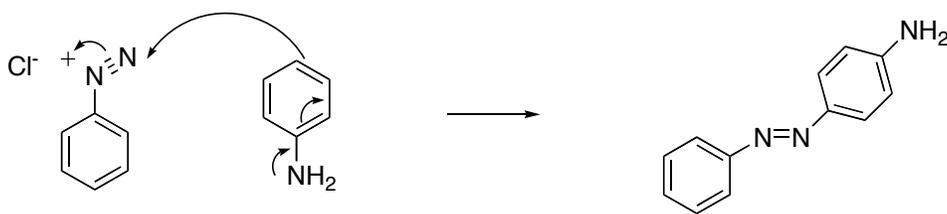
Diazonium Salts in Electrophilic Ar Substitution

Diazonium Coupling: requires phenol or aniline as a partner

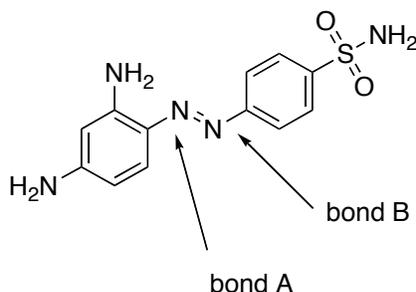
Diazonium salts undergo a coupling reaction with activated aromatic rings to yield very highly colored azo compounds with the general formula Ar-N=N-Ar as shown below. These azo coupled products are widely used as dyes because their extended conjugated π (π) electron systems cause them to absorb in the visible region of the electromagnetic spectrum. As shown below, Y must be OH, NH_2 or NHR or the reaction will generally not work.



Diazonium couplings are typically electrophilic aromatic substitutions in which the positively charged diazonium ion is the electrophile that reacts with the electron rich ring of a phenol or arylamine. Diazonium coupling often takes place at the *para* position, although *ortho* attack can take place if the *para* position is blocked. An example is the formation of an azo aniline derivative. Attack of the aniline from *para* position on the diazonium salt, followed by loss of a proton forms the product.

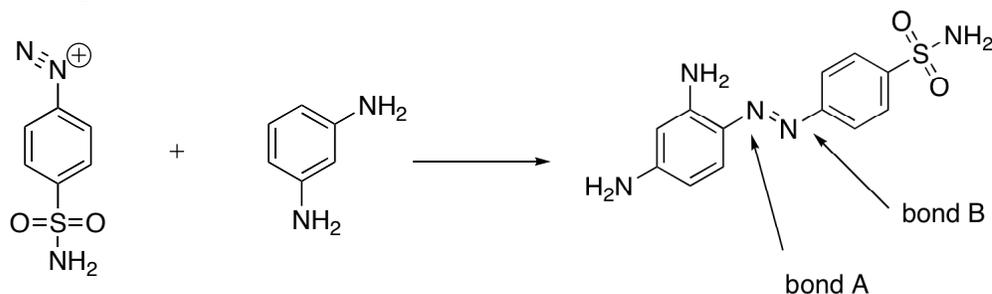


Aromatic azo compounds are valuable intermediates in the preparation of dyes and pharmaceuticals. For example, the sulfa drugs, such as Prontosil, were the first useful antibiotics known and were found to have a broad range spectrum. Prontosil is prepared commercially by a process that uses a diazonium ion. The diazonium coupling requires the use of a diazonium salt and a highly activated aromatic compound. For this reason, bond A (not bond B) is the bond formed in the diazonium coupling reaction. The aniline is highly reactive and would make a good coupling partner to the diazonium salt.

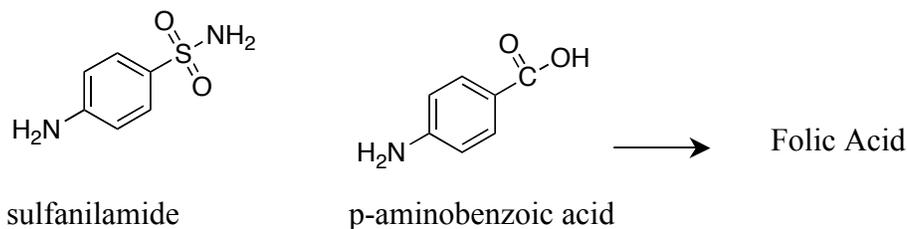


Prontosil

Bond A is formed last due to the nature of the groups on the aromatic rings. Amines are ortho/para directors whereas sulfonamides are meta directors.



Prontosil is an effective antibacterial agent because the human body readily converts it to sulfanilamide, which is the active drug. Sulfanilamide resembles the structure of para-aminobenzoic acid, which bacteria use to make folic acid. The bacteria mistakenly tries to use sulfanilamide (an enzyme inhibitor) and eventually dies.



Enzyme tries to use sulfanilamide to produce folic acid but no reaction takes place. The bacteria is not able to produce the folic acid needed to survive and eventually dies.