

Aromatics III

Electrophilic Substitution Rxns

Ref 15: 1 – 9 (both ed^{ns})

Prob 15: 1, 3, 5, 7 (both ed^{ns})

Adv Rdg 15: 10 – 14 (both ed^{ns})

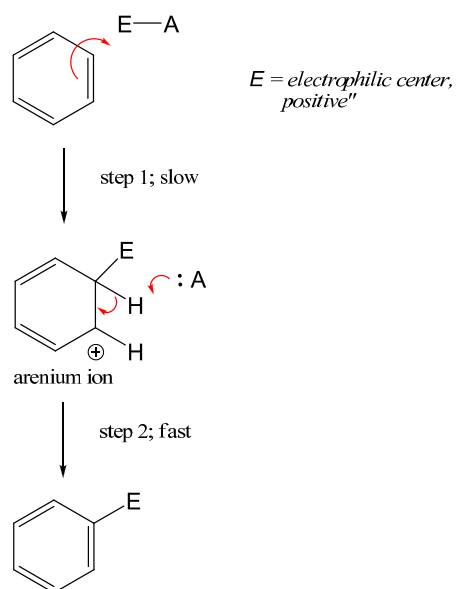
General

- π e⁻'s dominant
- but less reactive than in alkenes
(b/c of aromatic stabilization)
- **strong** electrophiles needed
- end result:

substitution

(not addition)

General Mechanism



N.B.

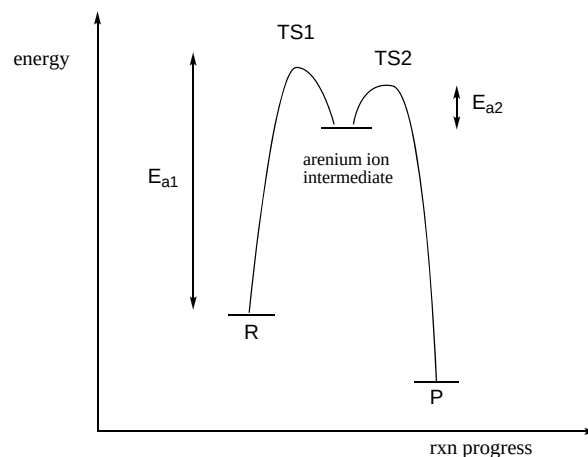
H replaced by E

step 1: similar to alkene rxn

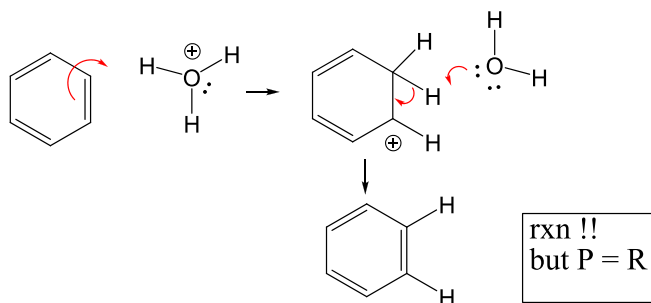
step 2: restoration of aromaticity

Reaction Profile

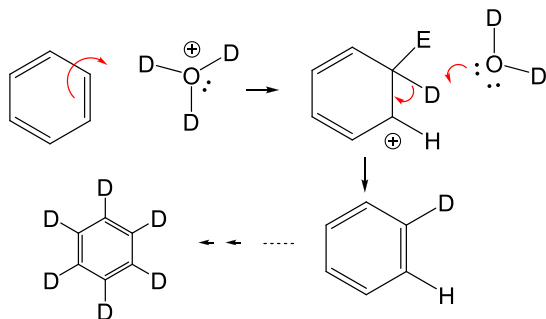
simple 2-step energy diagram; (like S_N1 , ...)



Ex. :Rxn w/ Strong Acid, $C_6H_6 + HCl(aq)$

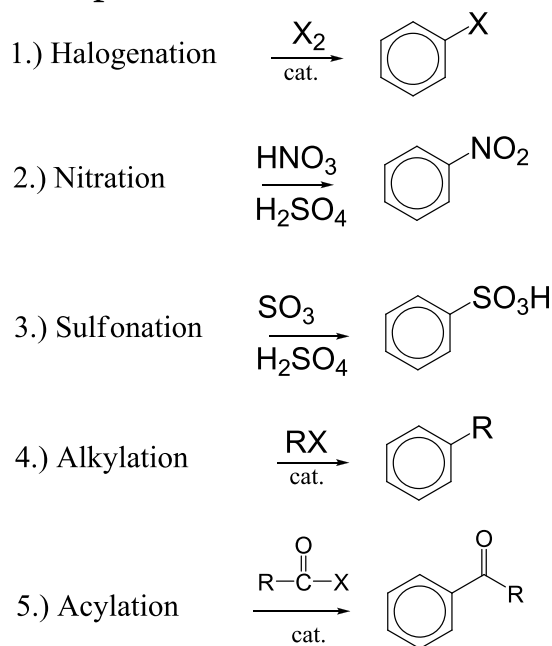


Proof: use HCl/D_2O



has no H signal in NMR !!

5 Important Rxns



1.), 4.), 5.) know detailed mech.

2.), 3.) know reagents & products

1.) Halogenation

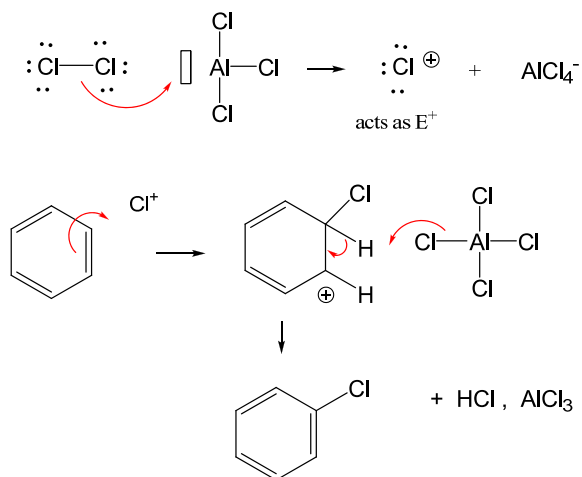
(w/ Cl_2 , Br_2 , I_2)

catalyst needed:

"Friedel-Crafts" = Lewis acid

common cat.: $AlCl_3$, $FeCl_3$ (have e^- gap)

Ex. Chlorination of benzene

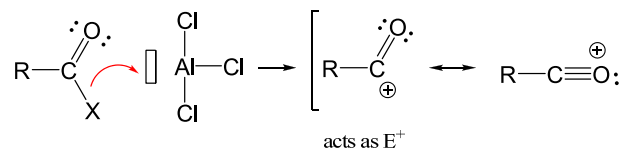


5.) Acylation

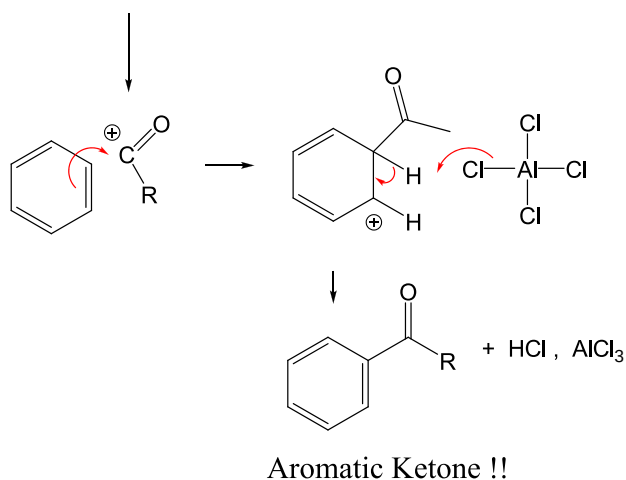


acid chloride =
"acyl chloride"

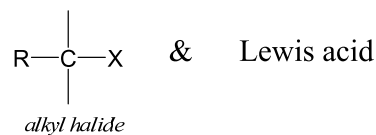
Formation of " E^+ ":



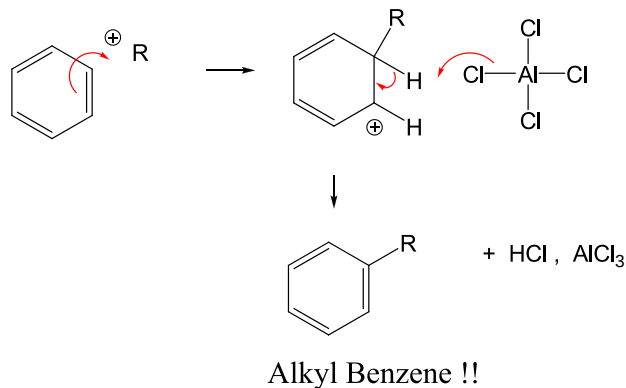
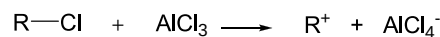
Example:



4.) Alkylation



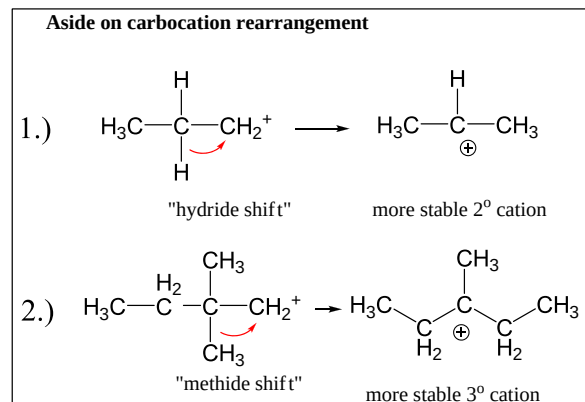
Details:



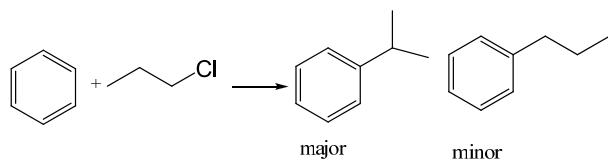
Complication w/ Alkylations

1.) rearrangement of R^+ cation

can occur if more stable cation
can be formed by hydride or alkyl shift



Ex.:



2.) multi - substitution

initial product may be more reactive
than original substrate

Ex.

