

Test 3 May 3

Test on everything since last test up until section 18.11 (Wednesday's class). Up through review sheet 6.

The Effect of Substituents on Reactivity Towards EAS

Section 18.12 & 18.13

Substituents can activate or deactivate the ring

18.12

(donate e^-)
made more e^- rich
(withdraw e^-)
made less e^- rich

for ring to be more reactive ... more attractive to an electrophile E^+

Directing the E^+ to a specific position

stabilize or destabilizing the intermediate

18.13

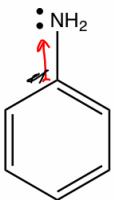
More e⁻ rich

The Effect of Substituents on Reactivity Towards EAS: Strongly Activating

Section 18.12 & 18.13

✓ σ bonds

electronegativity
+ inductive effects

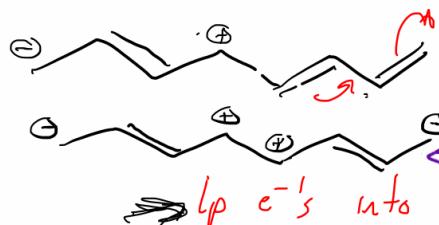


✗ π bonds

resonance/
extended π systems

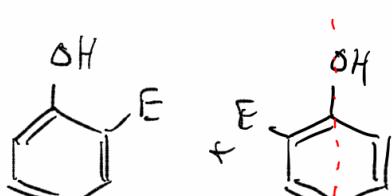
✗ σ withdrawing

π donating



don't do
this

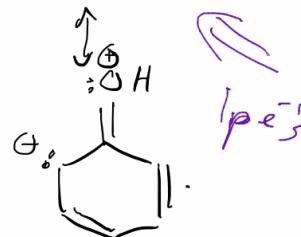
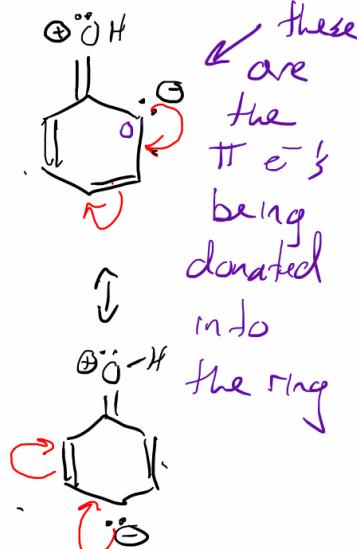
lp e⁻'s into π systems
make certain to walk
lp e⁻'s through entire π
system



✗ products

turn over
rotate 180°

→ empty p adjacent to π ,
→ π adjacent to π



when drawing
resonance contrib
3 is better than
2 so we draw 3

Resonance \rightarrow σ, p vs $m \leftarrow$ resonance skips More e^- rich

The Effect of Substituents on Reactivity Towards EAS: Strongly Activating

Section 18.12 & 18.13

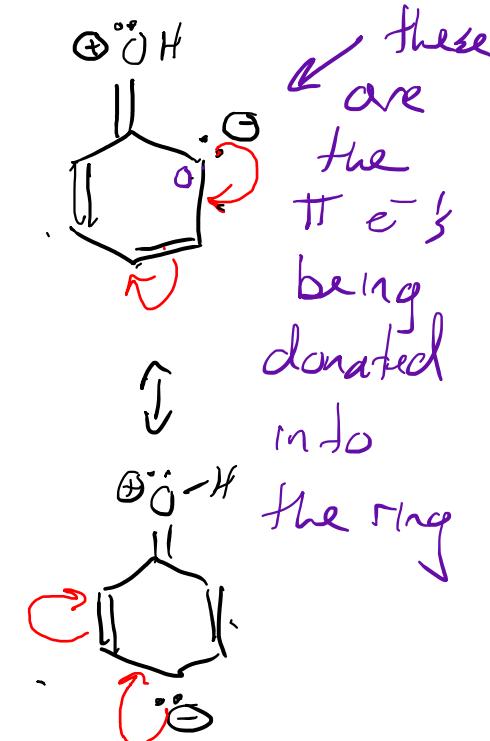
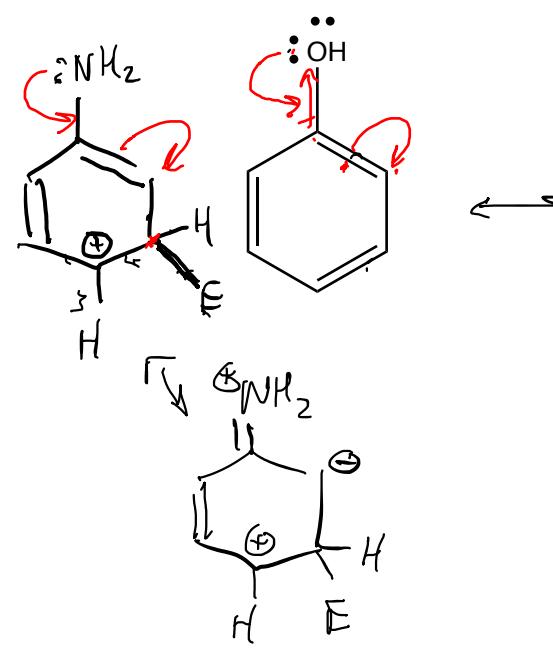
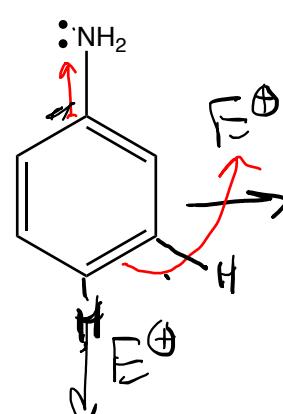
σ bonds

electronegativity
+ inductive effects

π bonds

resonance /
extended π systems

σ withdrawing
 π donating

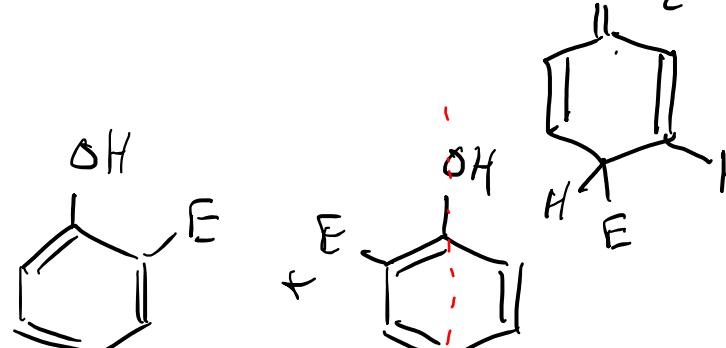
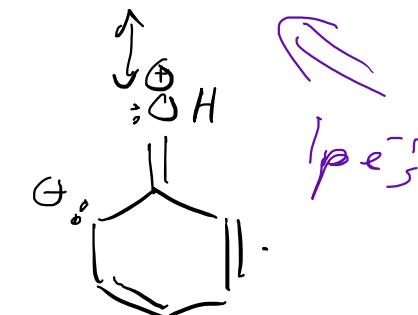


When the E^+ adds

to p or σ the extended π system stabilizes the \oplus .

$lp e^-$'s into π systems make certain to walk $lp e^-$'s through entire π system

\rightarrow empty p adjacent to π
 \rightarrow π adjacent to π



~~X~~ products

turn over
rotate 180°

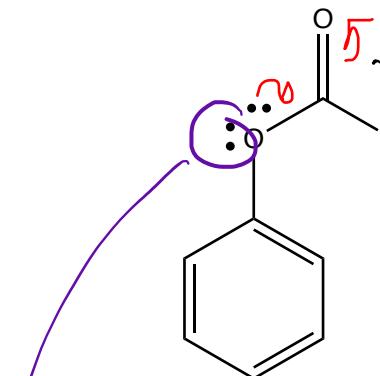
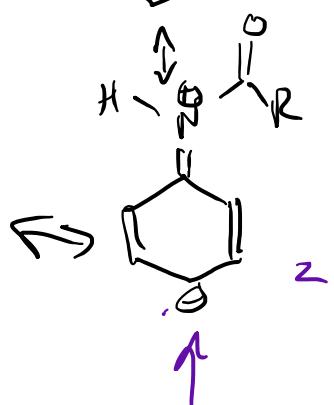
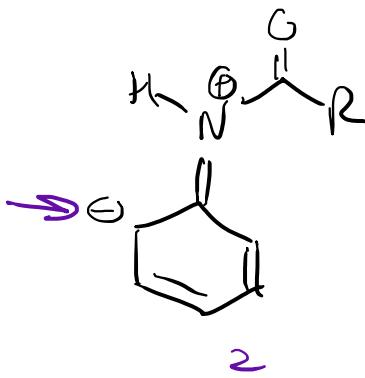
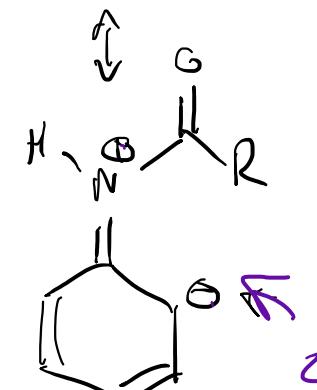
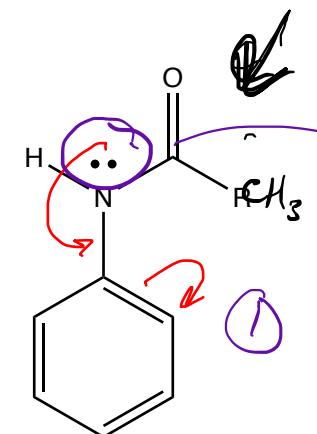
When drawing resonance contrib
3 is better than 2 so we draw 3

The Effect of Substituents on Reactivity Towards EAS: Moderately Activating

Section 18.12 & 18.13

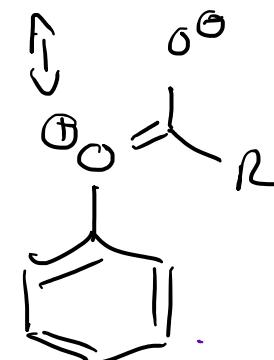
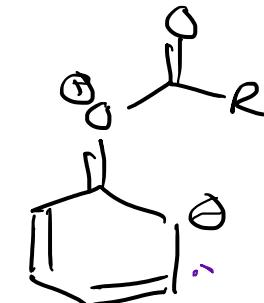
σ -withdrawing

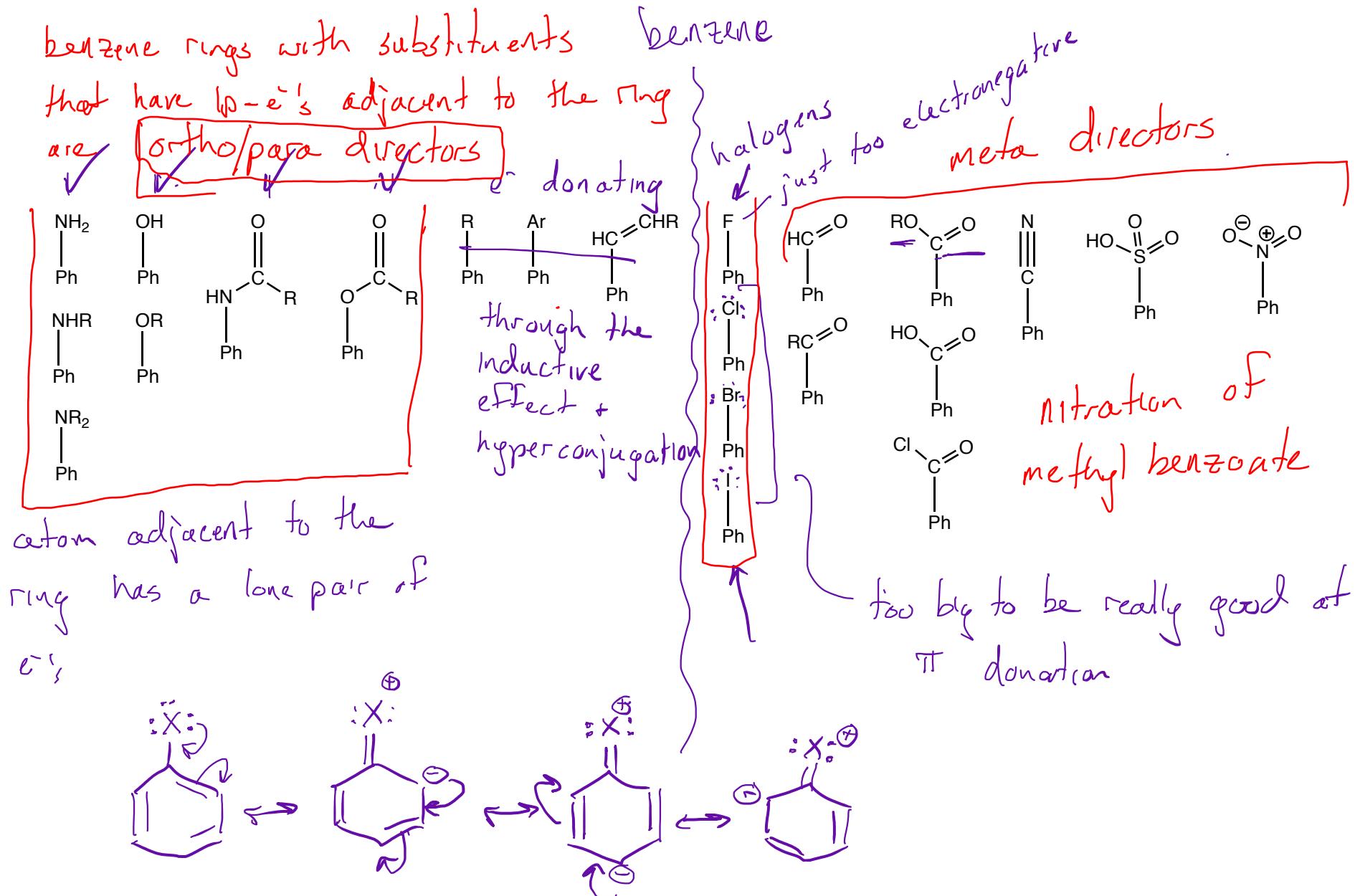
π -donating
enough to be
activators
weaker than
 $NR_2 + \delta R$
because of
 $C=O$



lone pair
 e^- is
means

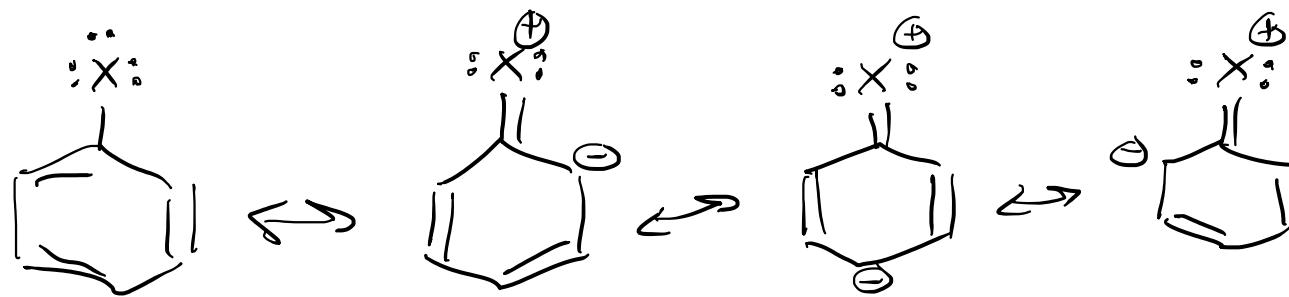
these are going
to be O, p directors
like the last page





resonance effects always shows up at ortho + para positions

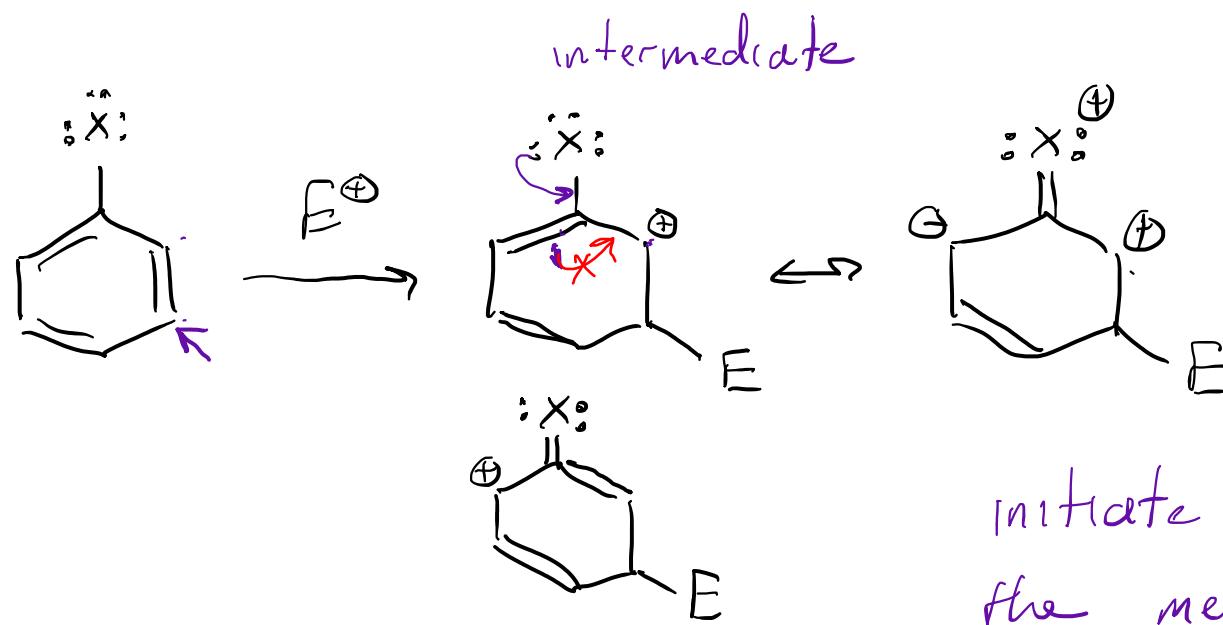
ortho/para director



ortho + para positions

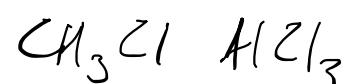
are more e^-

rich, more
attractive to
electrophiles

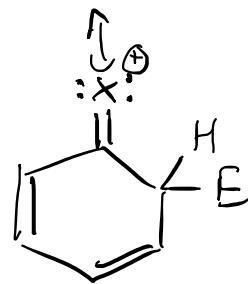
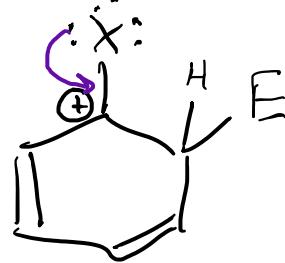
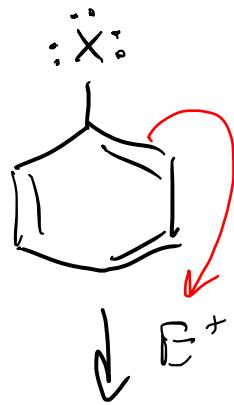


initiate the reaction at
the meta position

lone-pair e^- 's on substituents
don't help



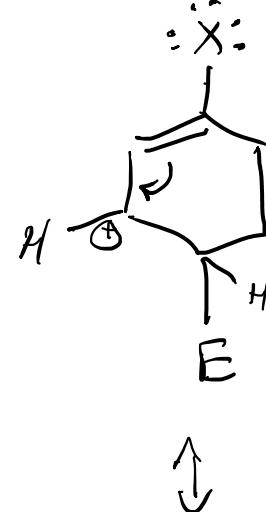
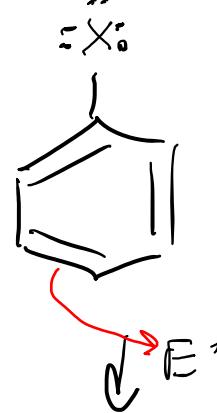
ortho



- \times helps to stabilize
- (+) charge

1. finish drawing intermediate
2. draw resonance contributors to show how \ddot{X} affects stability of intermediate

para



lp e^- 's on the atom adjacent to the benzene ring can stabilize the intermediate when E^+ adds to ortho + para positions

