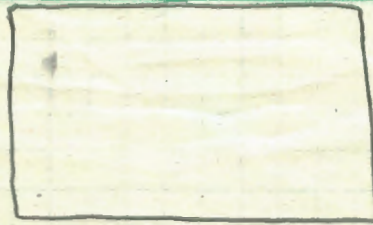
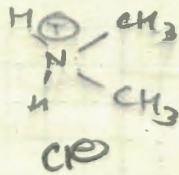
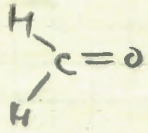


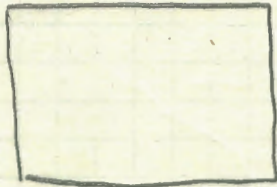
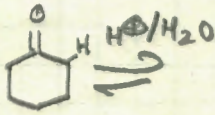
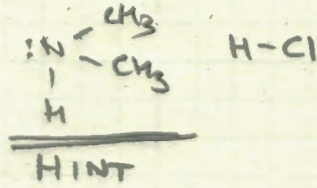
Review



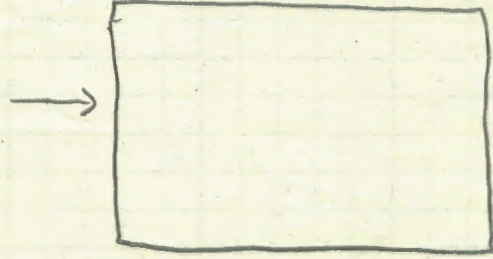
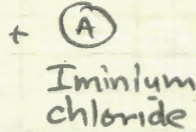
(A)

RECALL
1L equilibrium

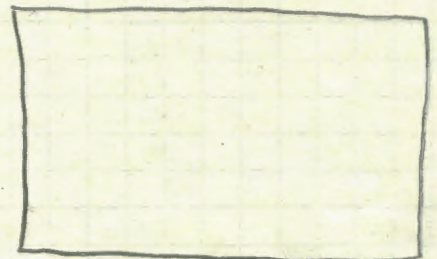
Mannich



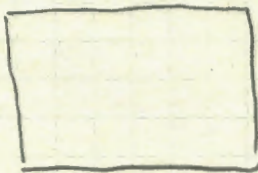
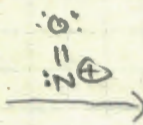
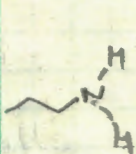
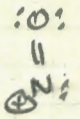
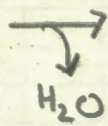
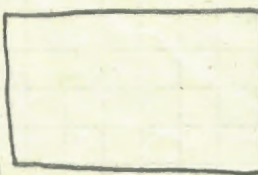
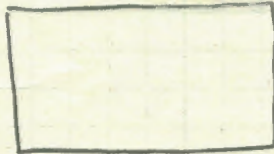
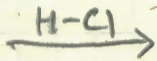
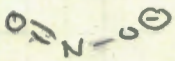
enol



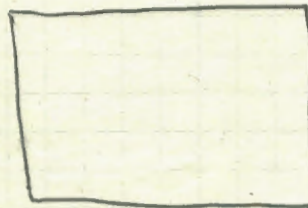
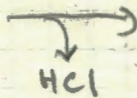
acid
↓ base (neutralizer)



neutral compound



ammonium
cation
chloride salt



a nitrosoamine

(B)

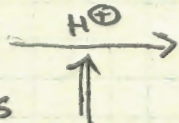
a 2° amine would stop here

From previous page

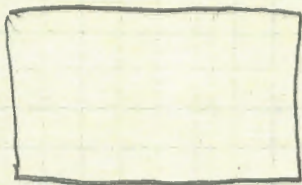
Nitrosamine

(B)

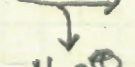
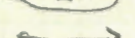
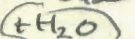
1° amine goes further



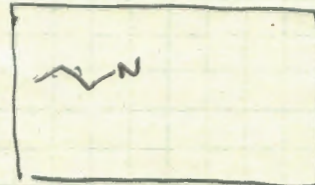
think of where this goes if $\text{R}_2\text{C}=\text{O}$



added as base



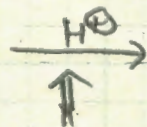
complete structure



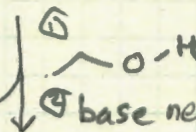
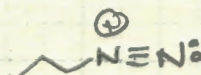
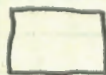
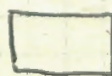
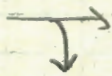
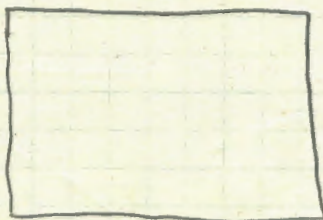
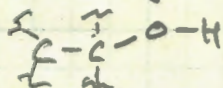
neutral diazohydroxide

(C)

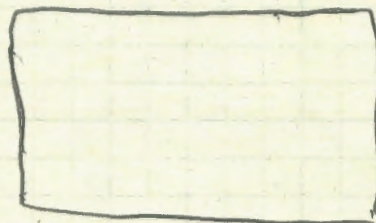
(C)



think where this goes if

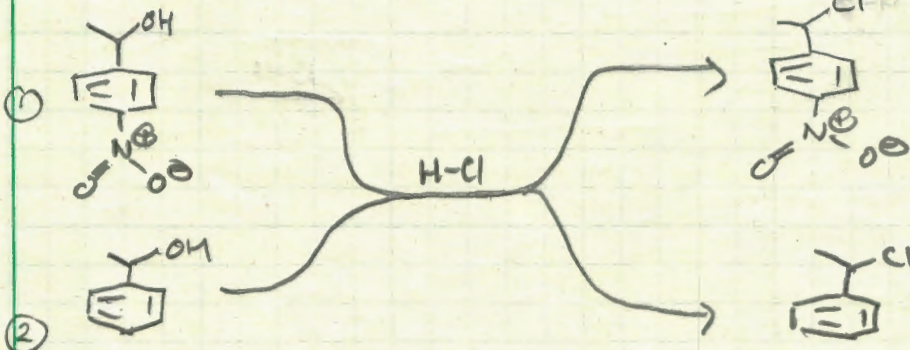


base neutralizer

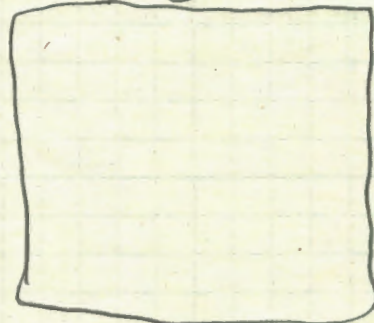


neutral compound

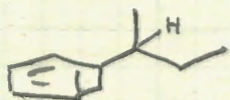
which Rxn is faster and why?



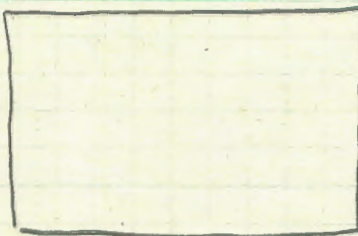
why? be specific



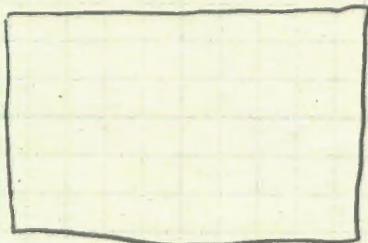
describe the specific electronic property that supports your answer



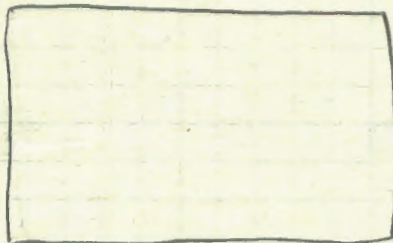
① $KMnO_4, H^+$
 ② H^+/H_2O
 proton source
 in 2nd step



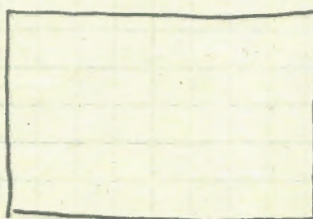
$Br_2, h\nu$



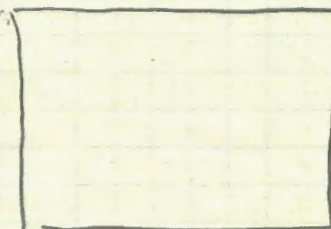
H^+/H_2O
 S_N2



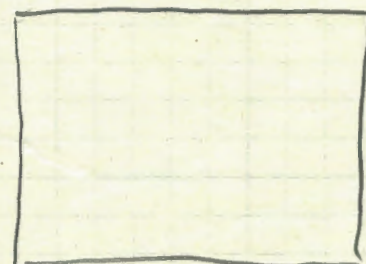
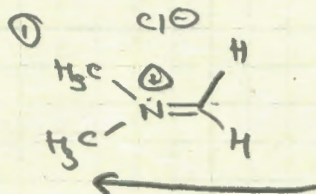
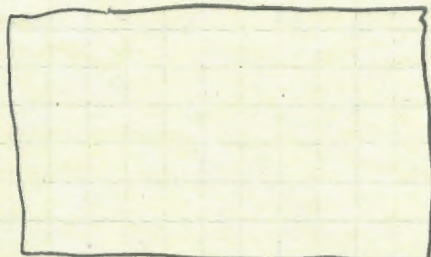
Cl_2
 Δ or
 $h\nu$



H^+
 S_N2

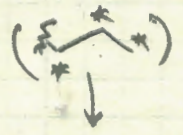
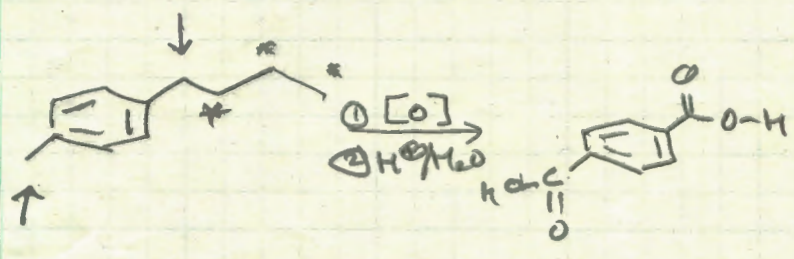
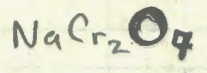
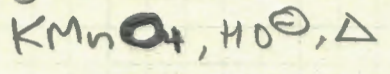


MnO_2 in solvent



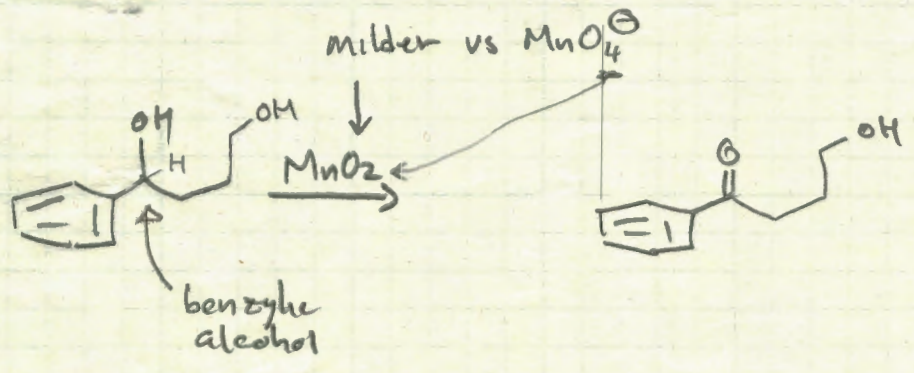
an Ammonium chloride salt

oxidation at benzylic sites with strong oxidants

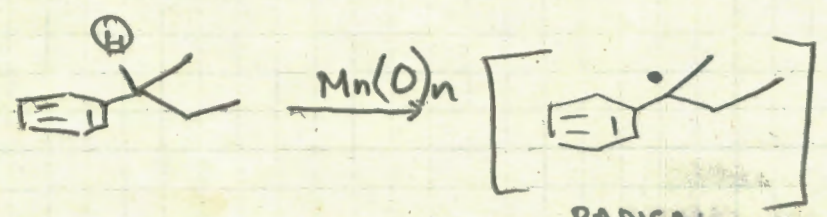
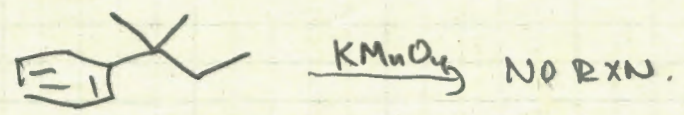
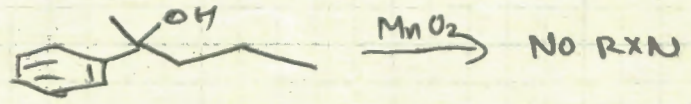


3 x CO₂ ↑

via radical oxidatⁿ steps and $[MnO]_n$ addⁿ.

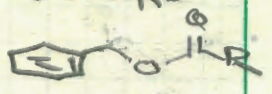
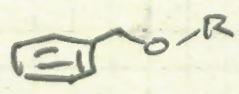
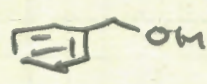


NOTE: these oxidatⁿ Rxns need AT LEAST 1 H ON THE BENZYLIC C.

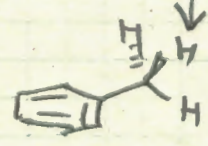


RADICAL
 INTERMEDIATE STARTS
 REDⁿ SEQUENCE
 \downarrow
 Reduction/Oxidatⁿ

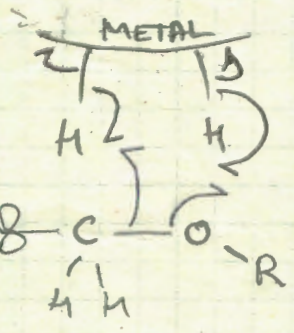
Reduction of BENZYLIC ALCOHOL ETHERS ESTERS



H₂, Metal



via



Convert



① What is lost?
one carbon

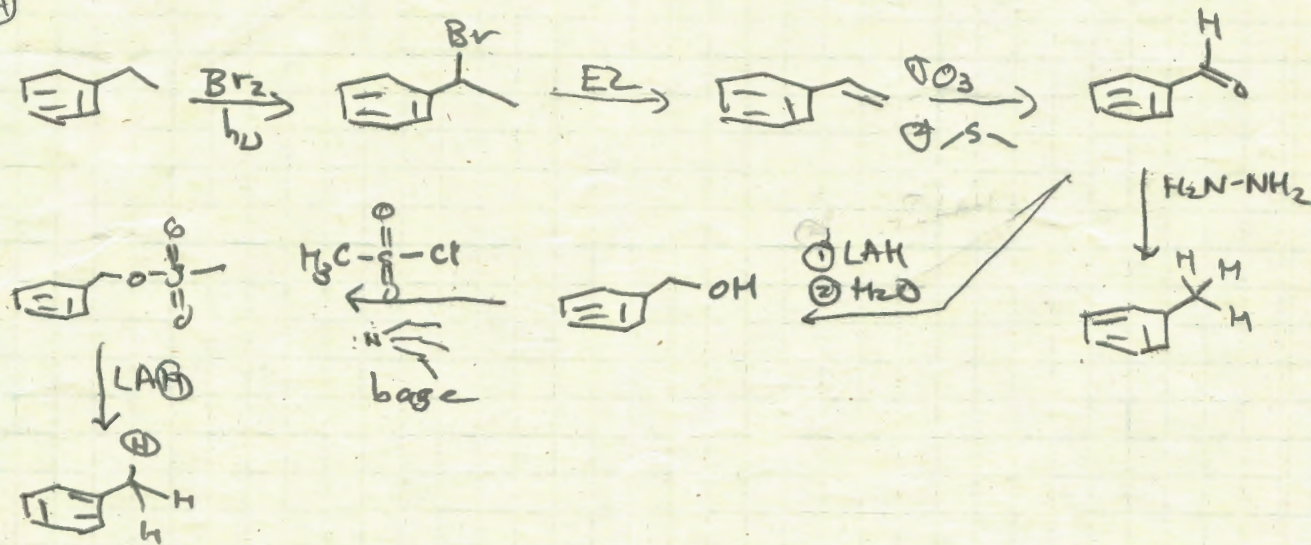
② What are some ways to do
C-C bond cleavage?

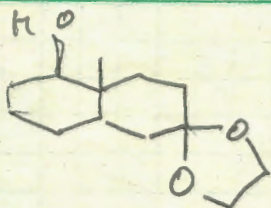
(A) O_3, Zn (B) -S-

OR

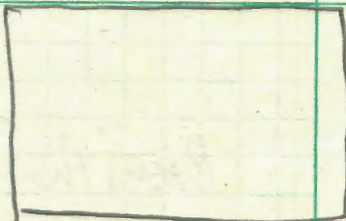
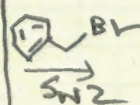
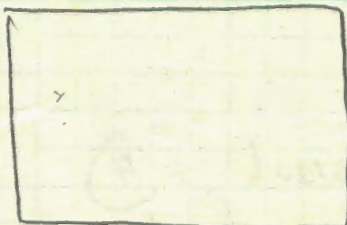
(B) $\text{KMnO}_4, \text{HO}^-$

④



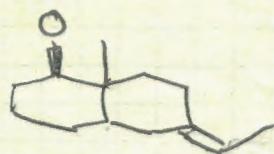
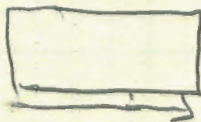
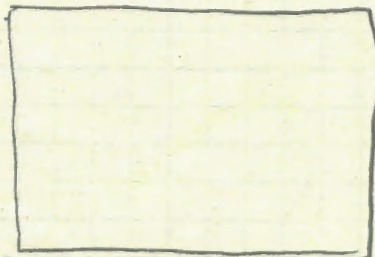


(base)
NaH



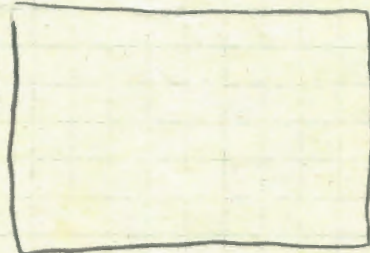
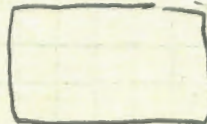
(A)

(A) $\xrightarrow[\Delta]{H^+/H_2O}$

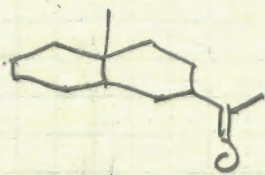


Complete the structure above

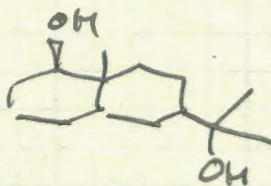
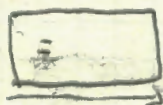
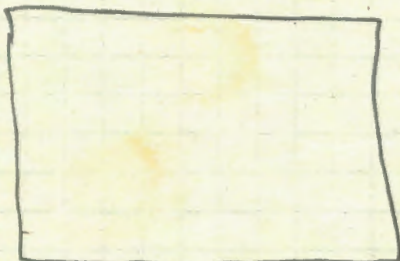
$\xrightarrow[1]{BH_3}$ $\xrightarrow[2]{H_2O_2, H^+}$



(B) $\xrightarrow[2]{H^+}$ $\xrightarrow[1]{CH_3Li}$
(neutralizer)



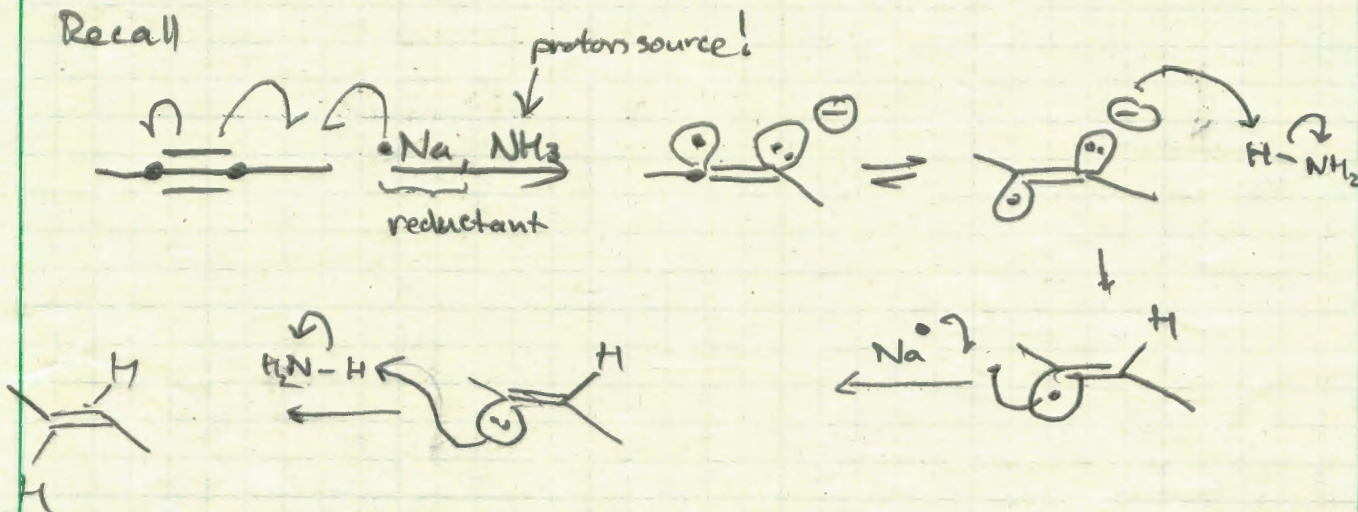
complete the str above



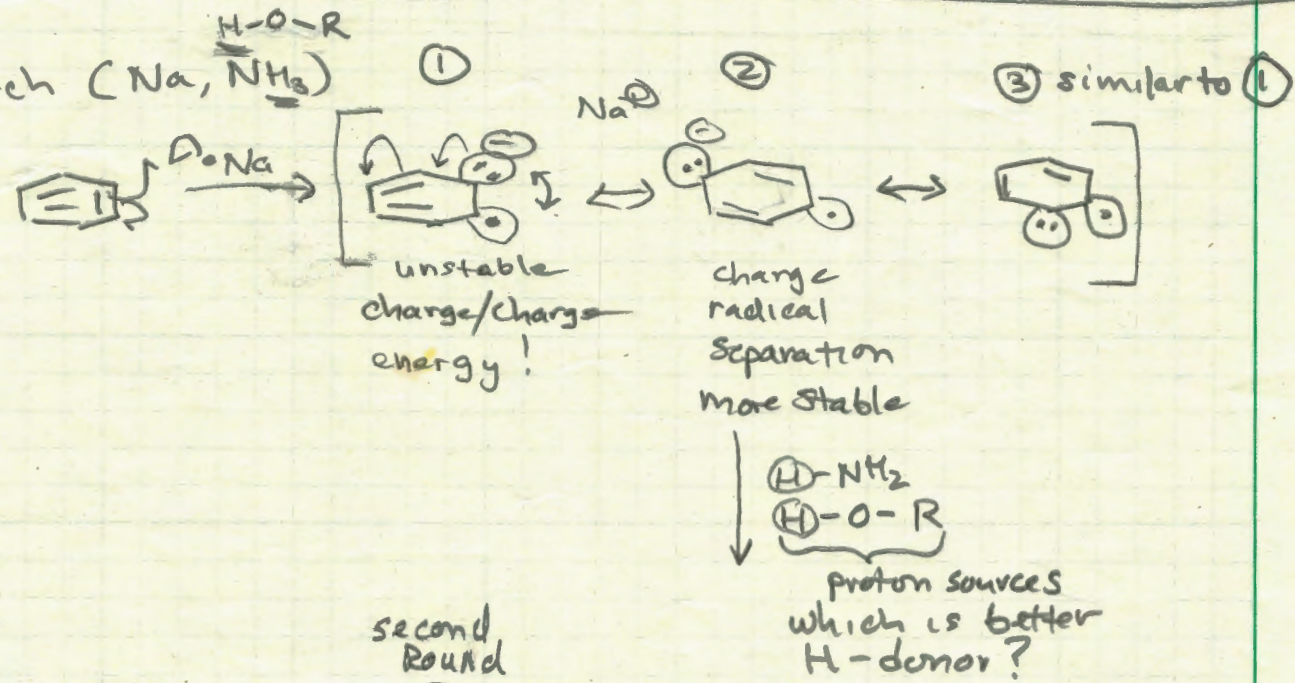
(B) \rightarrow

Birch Reduction of an Aromatic Ring

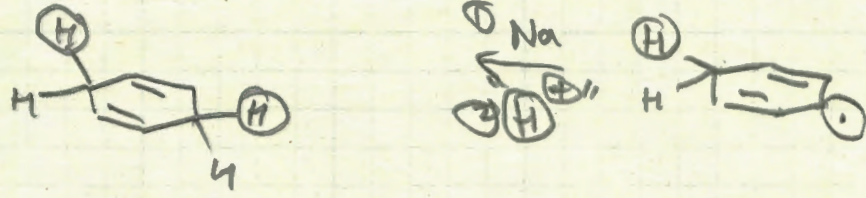
Recall

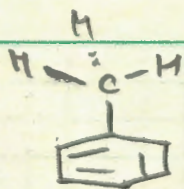


Birch (Na, NH₃)

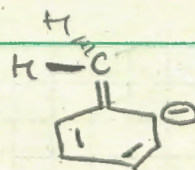
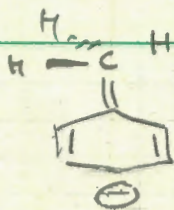


second round

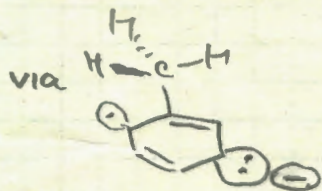




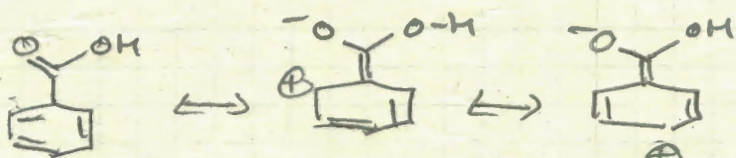
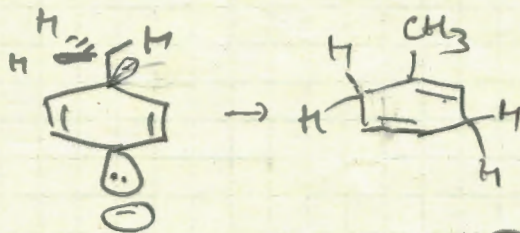
hyperconjugation



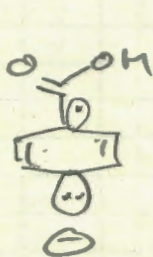
Birch anion will NOT form here because ξ CH₃ is an ortho/para director placing (-) chrg here



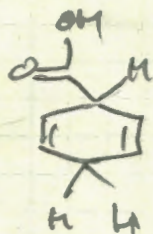
preferred over



(-) chrg favorable posits here!



preferred ∴



90% Birch product