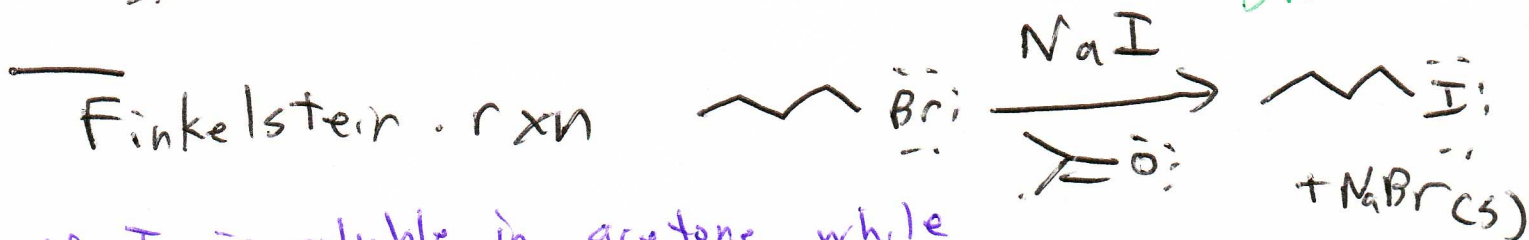
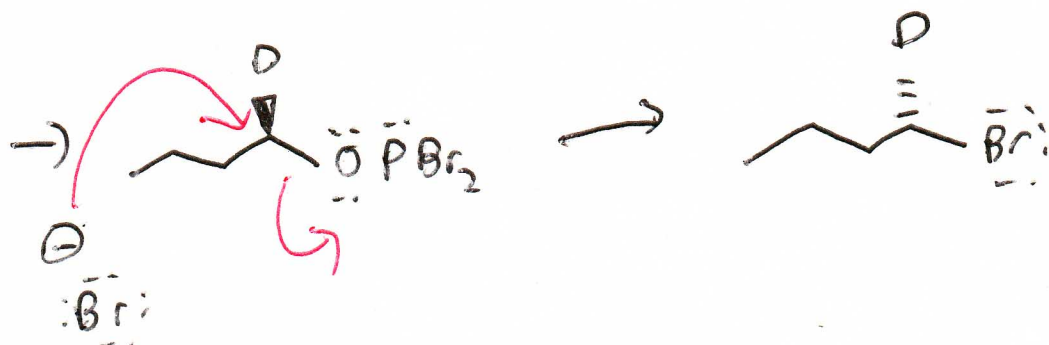
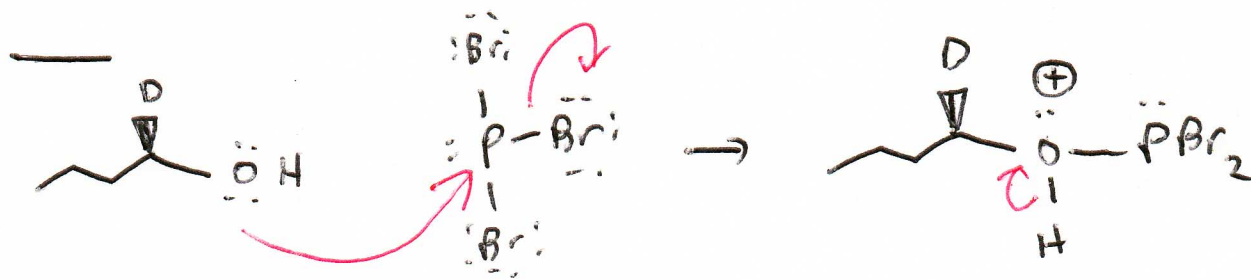
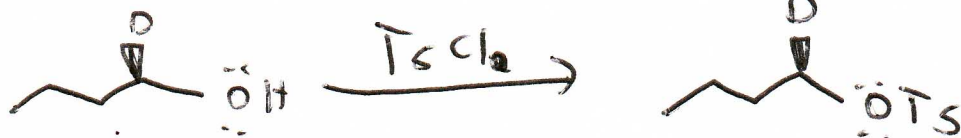
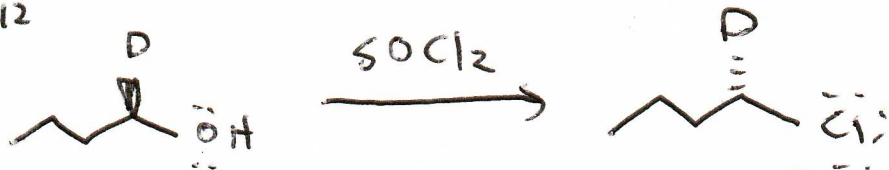


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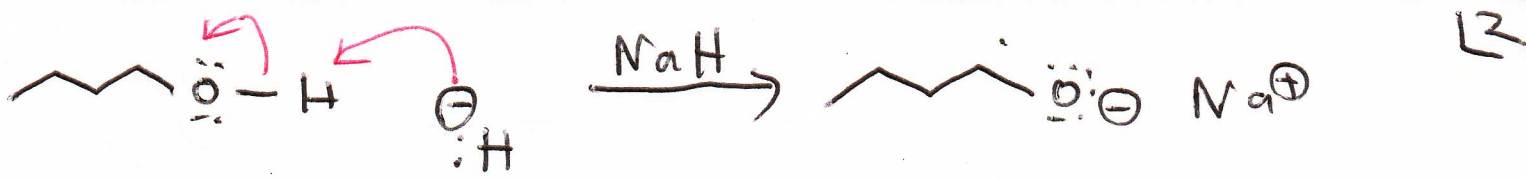
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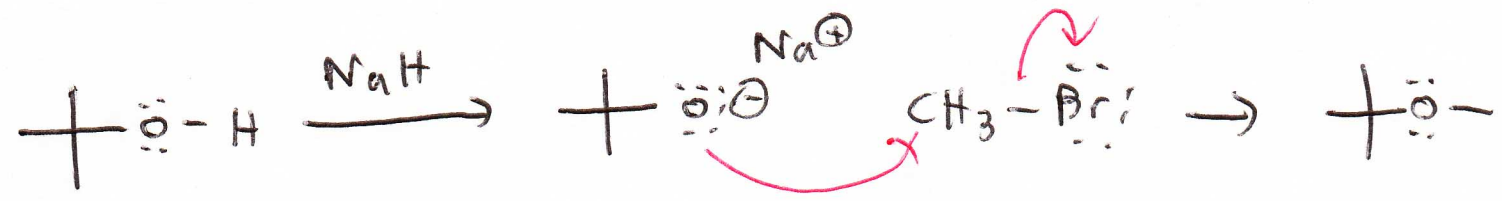
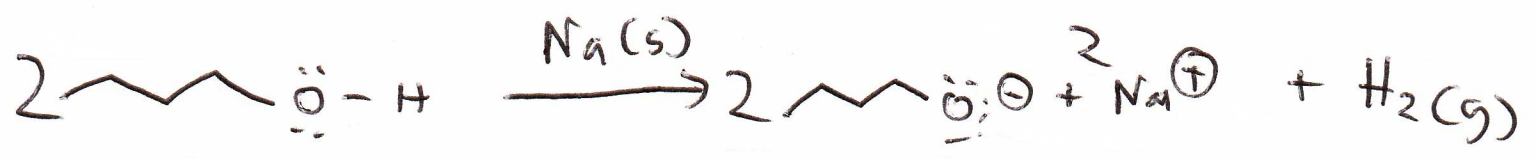
NaI is soluble in acetone, while NaBr and NaCl are not. As this rxn proceeds, NaBr (or NaCl) precipitate out of sol'n, which prevents Br⁻ (or Cl⁻) from re-attacking the alkyl iodide as it's formed.

Alkylation

Williamson ether synthesis S_N2



RO^- alkoxide CH_3O^- methoxide
CCO ethoxide

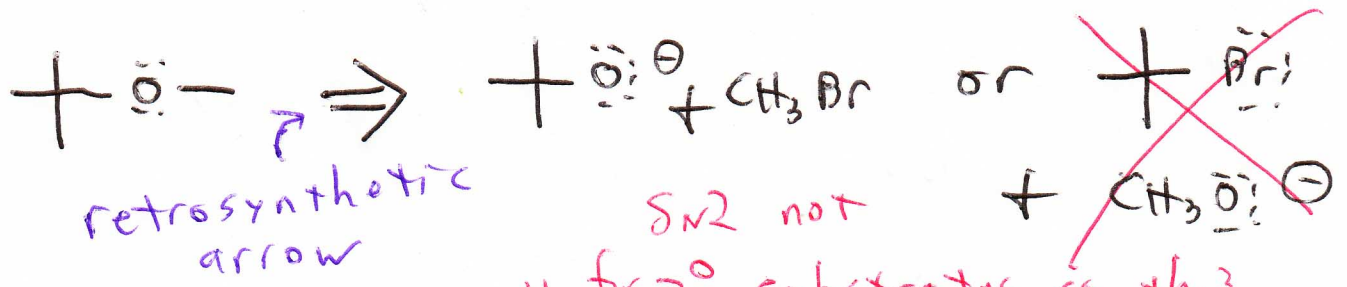


Ether nomenclature

common: CCOC diethyl ether
CCOC methyl ethyl ether

systematic: CCCCOC 1-ethoxybutane
 (longer chain as parent, shorter as substituent)

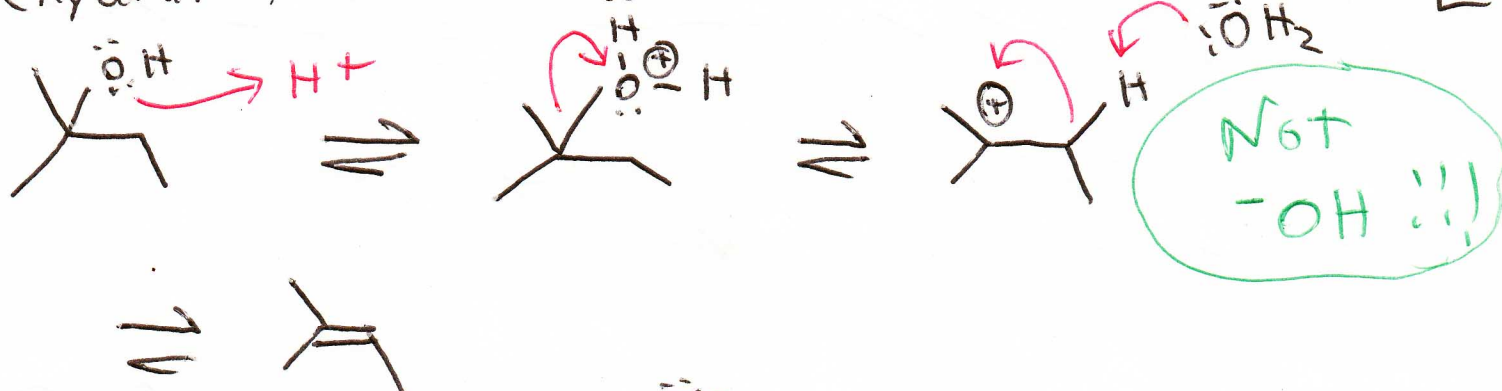
Synthesis strategy \rightarrow retrosynthesis



retrosynthetic arrow

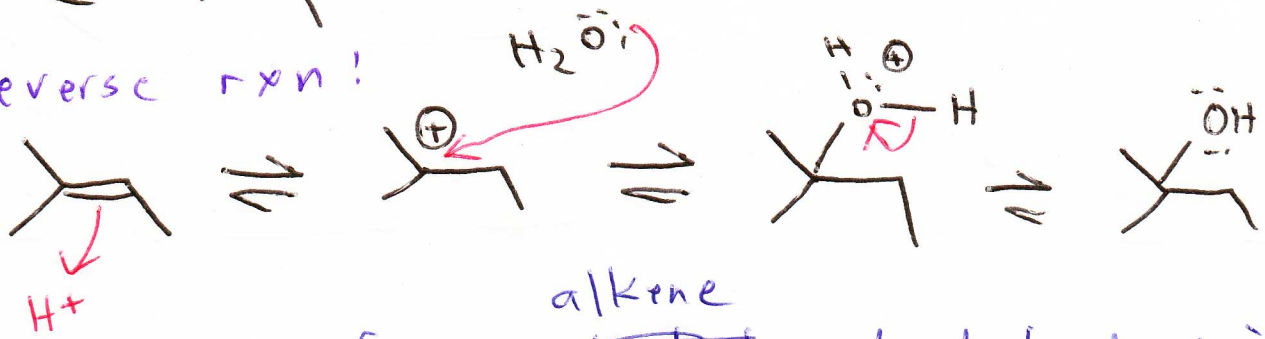
S_N2 not possible for 3° substrates, so this method is not possible

Dehydration of alcohols



Not -OH!!!

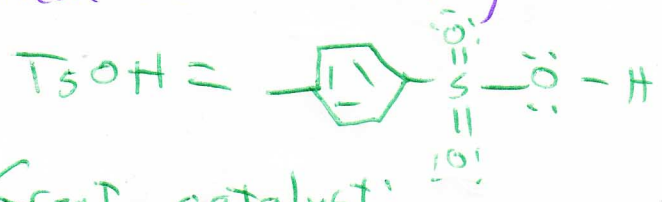
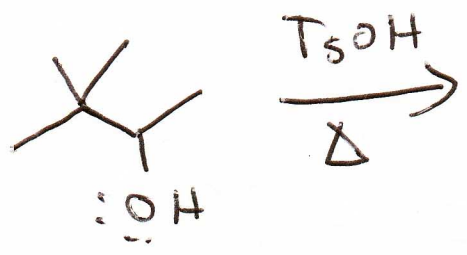
Reverse rxn!



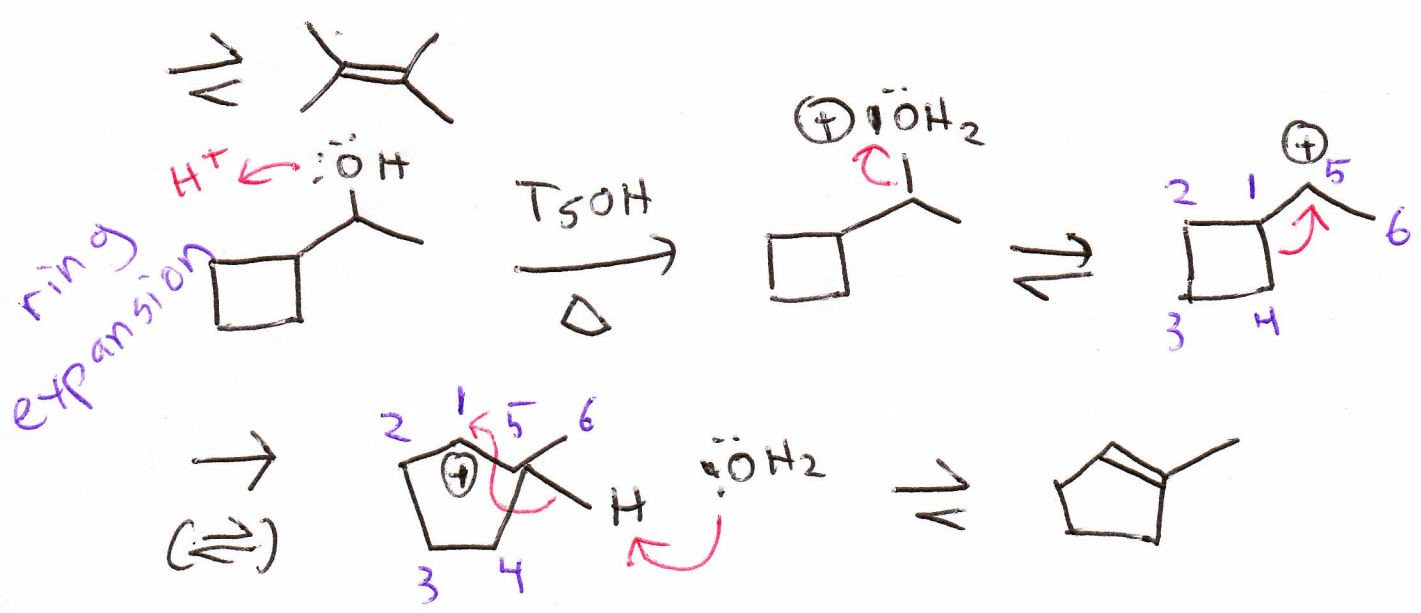
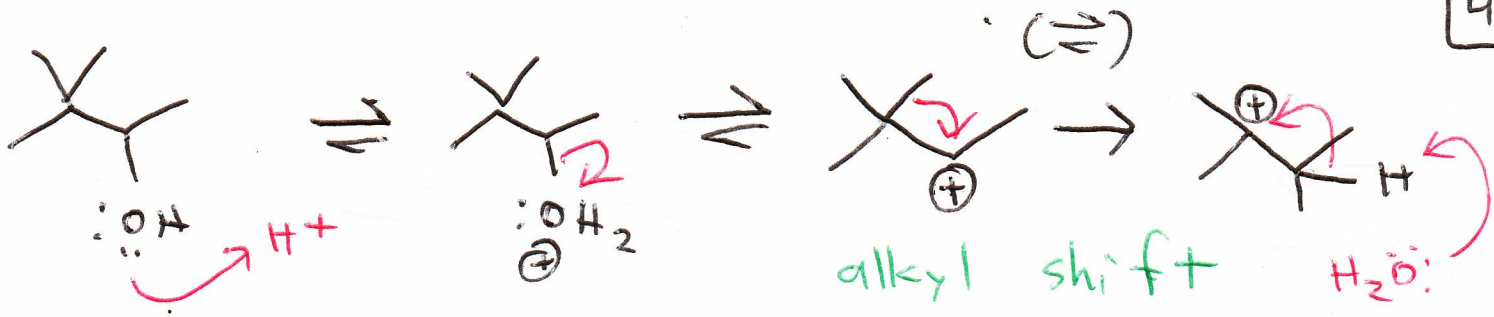
Hydration of an ~~alcohol~~ alkene and dehydration of an alcohol are just reverse rxns of each other. Dehydration is favored by high temperatures + removal of water (Le Chatelier's). Hydration is favored by adding water. Both rxns involve an acid catalyst.

(Dean-Stark Trap)

Acid-catalyzed dehydration normally favors the more thermodynamically stable alkene (most substituents; internal vs terminal; trans vs cis)



Great catalyst:
 cheap solid
 strong acid
 conjugate is non-nucleophilic



Pinacol rearrangement

