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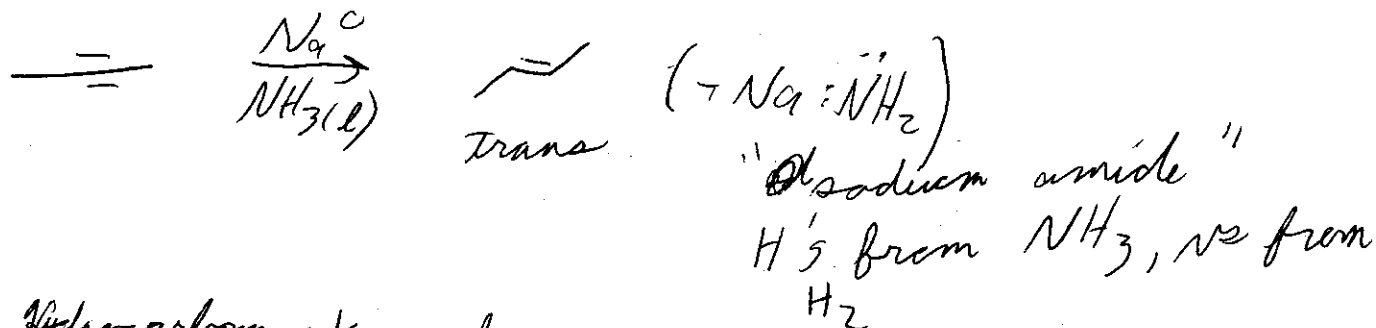
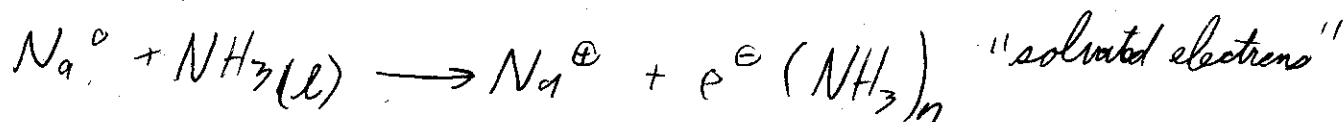
Exam 3 on Monday, same room as previously

Recall rxns of alkynes

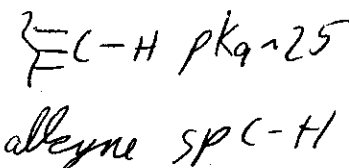
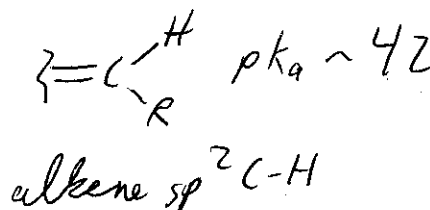
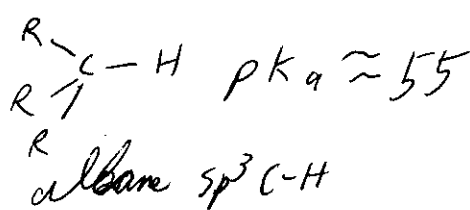


alternative approach produces trans alkenes from alkynes

"Dissolving metal rxn"



Hydrocarbon  $\text{pK}_a$  values

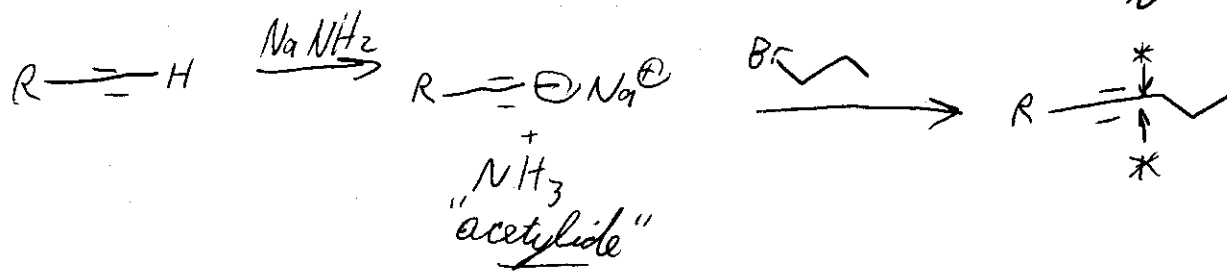


proportion of s-character is critical, More s  $\Rightarrow$  better accommodation of  $\text{e}^-$ s

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Synthetic ramification via alkyne derived anions as nucleophiles to form new C-C bonds via  $S_N2$  rxns



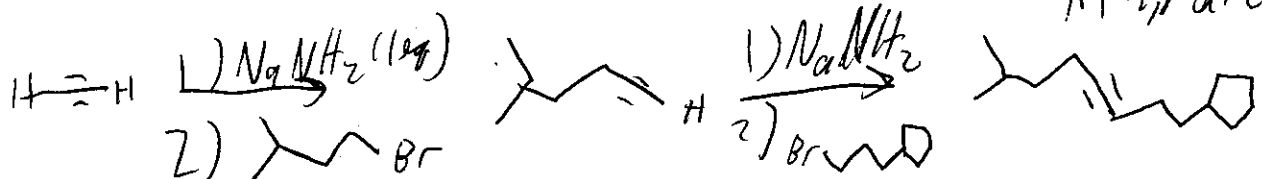
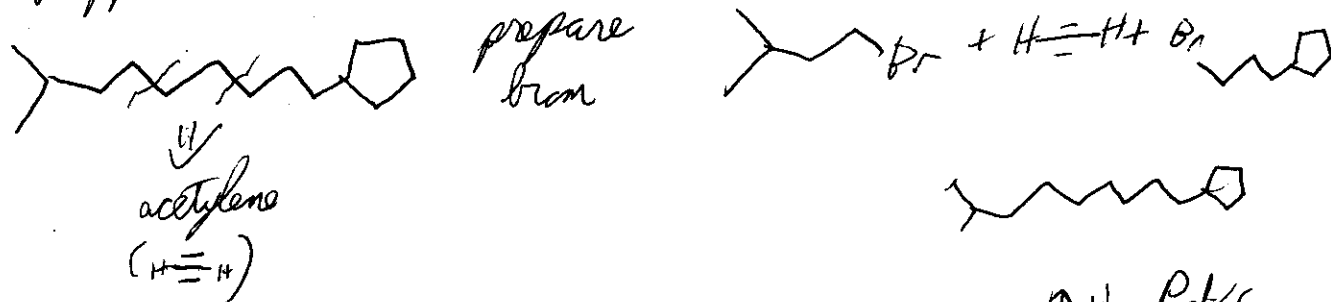
Note: For Li, Mg, Cu C-M from C-halogen

vs

$R-C\equiv C^{\ominus}$  via deprotonation

Limitation in C-C bond formation via  $R-C\equiv C^{\ominus}Na^{\oplus}$ :  
only 1° alkyl halides ( $\neq CH_3$ ) are effective (E2 elimination for 2° R-X)

Ex of application:



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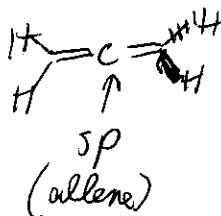
Ch. 15 - Dienes, Resonance, &amp; Aromaticity

Rec Problems: 3-5, 11-24, 27-30, 33-45, 50-73

Diene = Di-alkene, i.e., has 2 C=C units

Classify dienes into three groups, based largely on spacing b/w C=C units

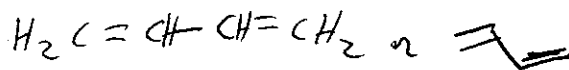
1) allenes



3D structure has H twist



2) Conjugated dienes - C=C units are immediately adjacent



(1,3-butadiene)

caution This spacing is necessary for conjugation but not sufficient

3) Ordinary dienes  $\rightarrow$  at least one  $sp^3$  atom (C) b/w C=C units