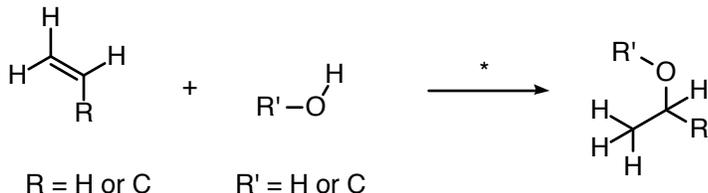


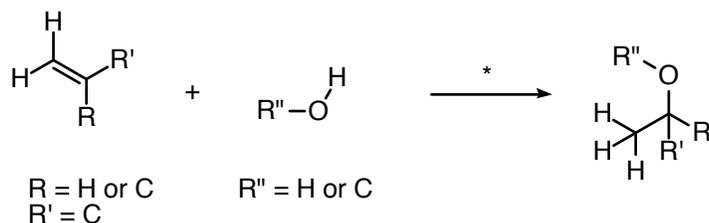
#1.



\* Notes:

- Filename: AdditionOfROHAcrossMonosubstitutedAlkene
- Acid-catalyzed addition of water or alcohol across ethene or a monosubstituted alkene
- Requires an acid catalyst, such as  $\text{H}_2\text{SO}_4$  or  $\text{H}_3\text{PO}_4$
- For monosubstituted alkenes, this rule should demonstrate the preference for substitution at the 2° carbon

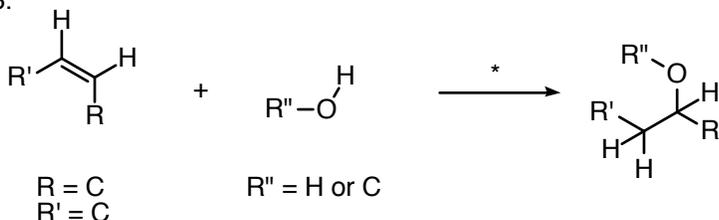
#2.



\* Notes:

- Filename: AdditionOfROHAcrossGemDisubstitutedAlkene
- Acid-catalyzed addition of water or alcohol across a monosubstituted or 1,1-disubstituted alkene
- Requires an acid catalyst, such as  $\text{H}_2\text{SO}_4$  or  $\text{H}_3\text{PO}_4$
- This rule should demonstrate the preference for substitution at the more substituted carbon

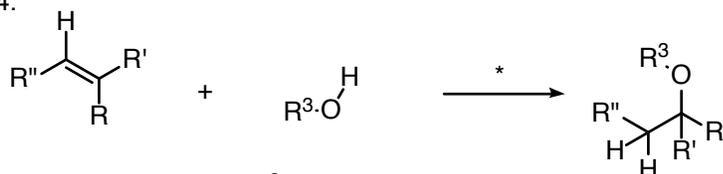
#3.



\* Notes:

- Filename: AdditionOfROHAcrossVicDisubstitutedAlkene
- Acid-catalyzed addition of water or alcohol across a 1,2-disubstituted alkene
- Requires an acid catalyst, such as  $\text{H}_2\text{SO}_4$  or  $\text{H}_3\text{PO}_4$
- Given the R and R' groups have the same constraints, this rule should demonstrate there is no preference for substitution, and I am curious how the algorithm will proceed with this in the case of an unsymmetrical alkene starting material. Will it recognize that either/both product is possible and potentially carry both through potential synthesis pathways, or will it arbitrarily decide on one product, ignoring the other?

#4.



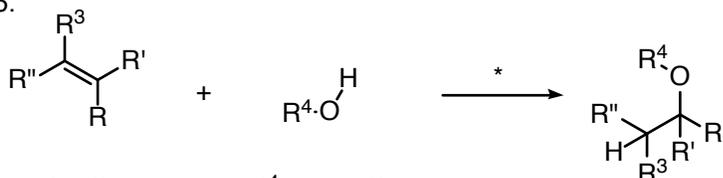
R = C  
R' = C  
R'' = C

R<sup>3</sup> = H or C

\* Notes:

- Filename: AdditionOfROHAcrossTrisubstitutedAlkene
  - Acid-catalyzed addition of water or alcohol across a trisubstituted alkene
  - Requires an acid catalyst, such as H<sub>2</sub>SO<sub>4</sub> or H<sub>3</sub>PO<sub>4</sub>
  - This rule should demonstrate the preference for substitution at the more substituted carbon
- 

#5.



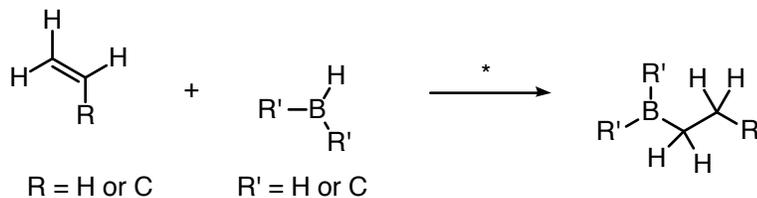
R = C  
R' = C  
R'' = C  
R<sup>3</sup> = C

R<sup>4</sup> = H or C

\* Notes:

- Filename: AdditionOfROHAcrossTetrasubstitutedAlkene
  - Acid-catalyzed addition of water or alcohol across a tetrasubstituted alkene
  - Requires an acid catalyst, such as H<sub>2</sub>SO<sub>4</sub> or H<sub>3</sub>PO<sub>4</sub>
  - As with #3 above, there is no preference for position of reaction
- 

#6.



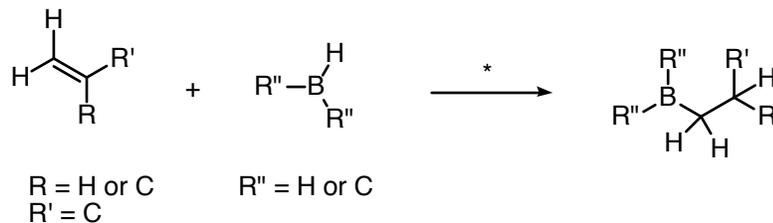
R = H or C

R' = H or C

\* Notes:

- Filename: HydroborationOfMonosubstitutedAlkene
  - Hydroboration of ethene or a monosubstituted alkene
  - For monosubstituted alkenes, this rule should demonstrate the preference for substitution at the less substituted carbon
- 

#7.



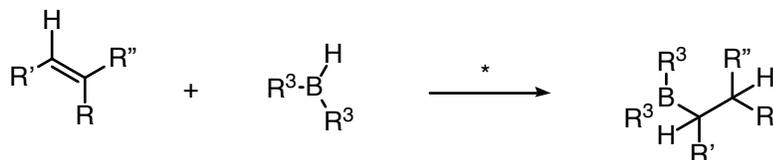
R = H or C  
R' = C

R'' = H or C

\* Notes:

- Filename: HydroborationOfGemDisubstitutedAlkene
- Hydroboration of a monosubstituted or 1,1-disubstituted alkene
- For monosubstituted alkenes, this rule should demonstrate the preference for substitution at the less substituted carbon

#8.



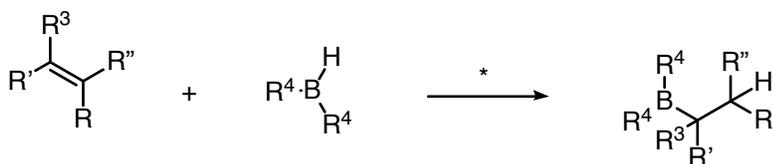
R = C  
R' = C  
R'' = H or C

R<sup>3</sup> = H or C

\* Notes:

- Filename: HydroborationOfDiOrTrisubstitutedAlkene
- Hydroboration of a 1,2-disubstituted or trisubstituted alkene
- Preference for substitution on the less substituted carbon for the trisubstituted, but no preference for 1,2-disubstituted (as with #3 above)

#9.



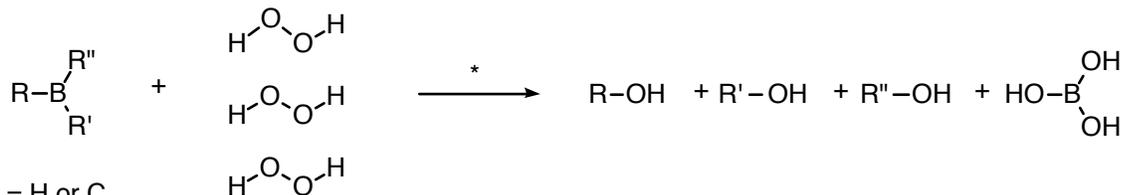
R = C  
R' = C  
R'' = C  
R<sup>3</sup> = C

R<sup>4</sup> = H or C

\* Notes:

- Filename: HydroborationOfTetrasubstitutedAlkene
- Hydroboration of a tetrasubstituted alkene
- No preference (as with #3 above)

#10.



R = H or C  
R' = H or C  
R'' = H or C

\* Notes:

- Filename: OxidationOfBorane
- Oxidative work up of a borane
- Requires a base, like NaOH, the identity of which is inconsequential for the balanced equation

#11.

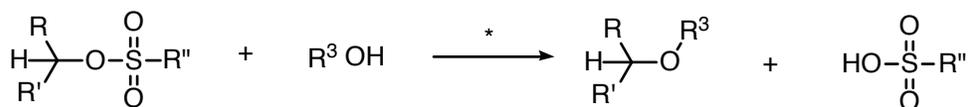


R = C      R' = C

\* Notes:

- Filename: SulfonylationOfAlcohol
- Sulfonylation of an alcohol (which converts the oxygen into a suitable leaving group for substitution and elimination reactions)
- Requires a base to neutralize the HCl generated, but the identity of this base is inconsequential.

#12.



R = H or C  
R' = H or C  
R'' = C

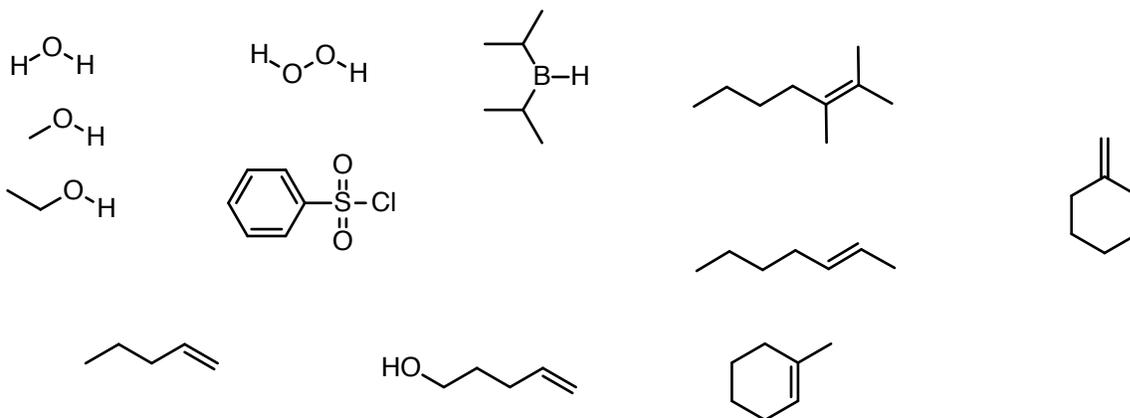
\* Notes:

- Filename: EtherFormationBySulfonateDisplacement
- Synthesis of an ether by  $S_N2$  reaction
- Requires a base (e.g. NaH) to deprotonate the nucleophile prior to nucleophilic substitution (and therefore generate the sulfonate salt rather than the sulfonic acid), but this is inconsequential to the balanced equation.
- The definitions of R and R' are meant to signify that the sulfonate must be methyl, 1°, or 2° (but cannot be 3°)

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**Available Chemicals:**



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**Targets:**

