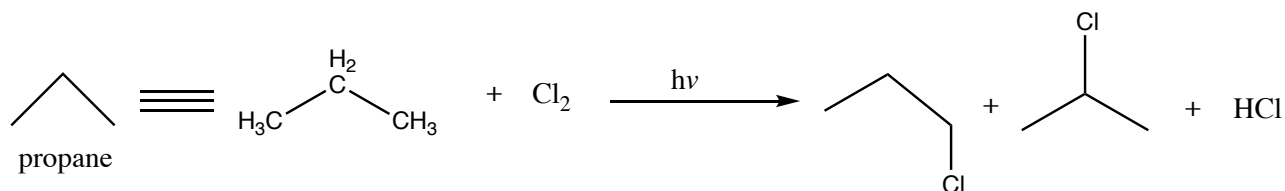
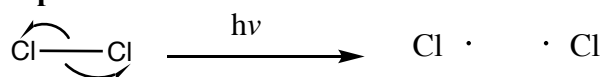


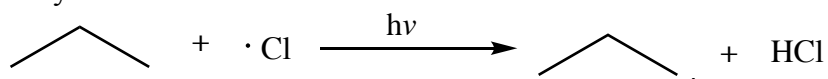
REVIEW: Halogenation of Alkanes

Mechanism (Radical Substitution)

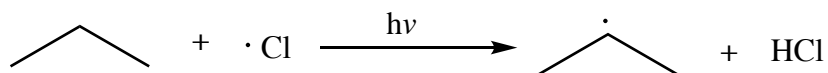
A. Chlorination of Propane

**1. Initiation Step****2. Propagation Step**

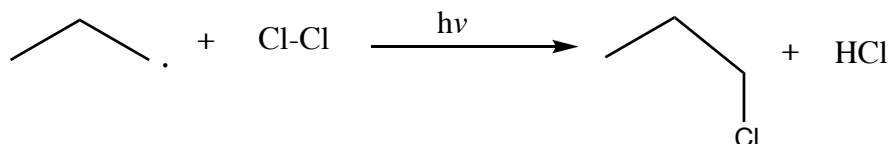
a. Primary radical formation



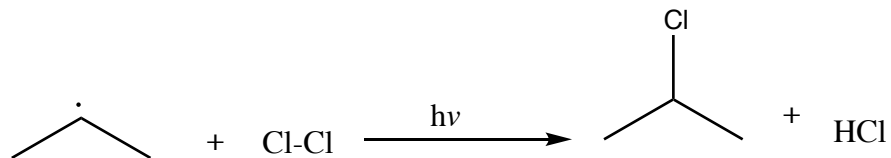
b. Secondary radical formation

**3. Termination Step**

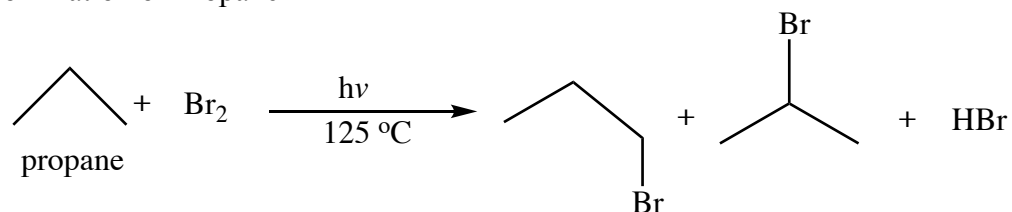
a. Lesser stable reaction (due to primary radical propane)



b. More stable reaction (due to secondary radical propane)

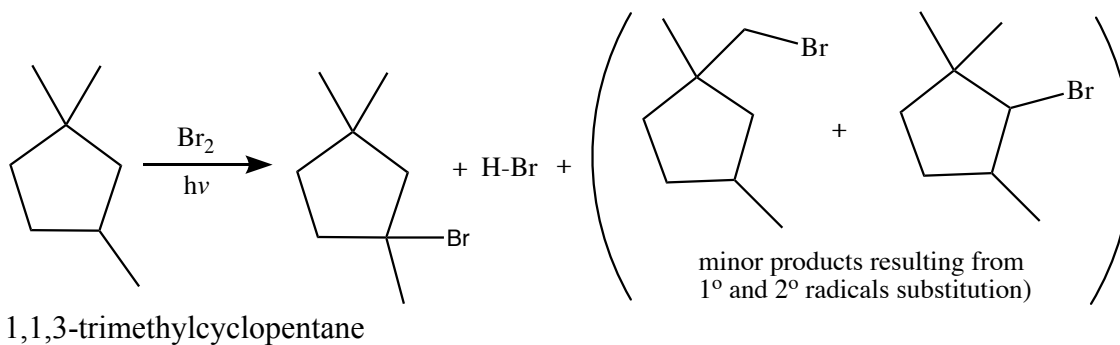


B. Bromination of Propane

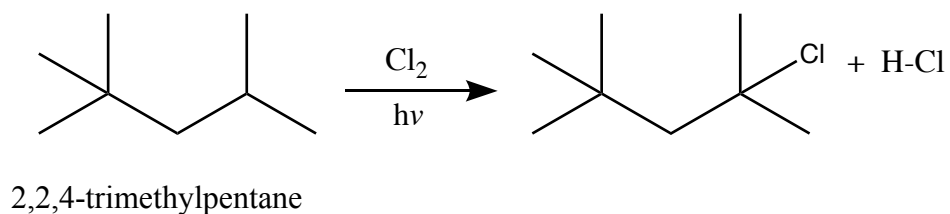


More Examples

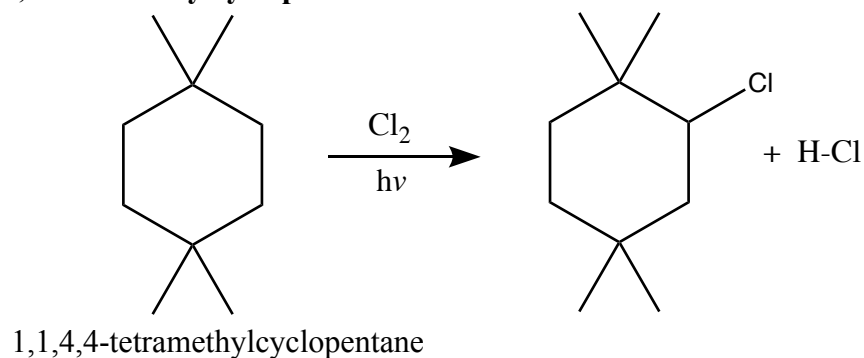
A. 1,1,3-trimethylcyclopentane bromination



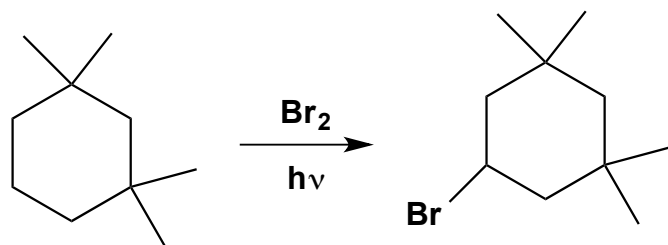
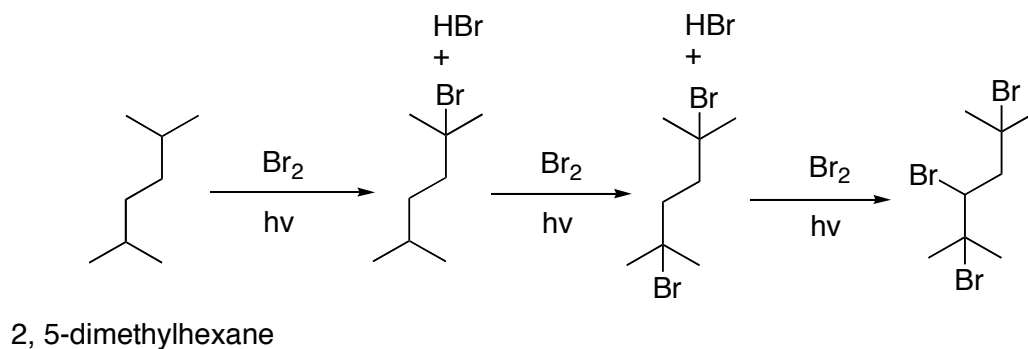
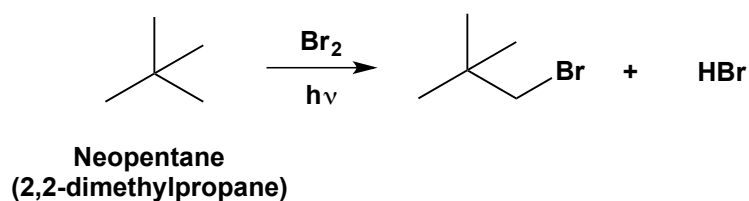
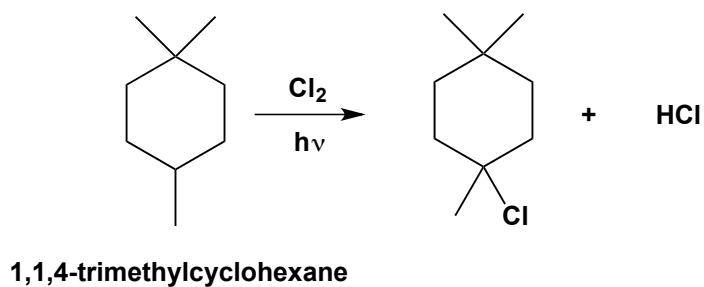
B. 2,2,4-trimethylpentane chlorination



C. 1,1,4,4-tetramethylcyclohexane chlorination



Additional examples for your reference:

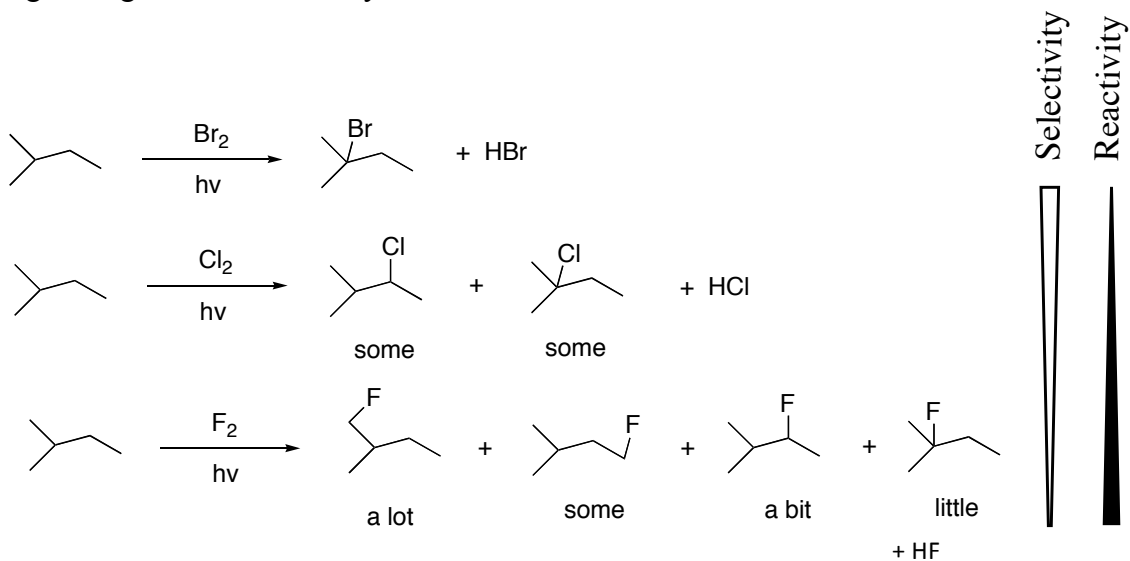


1,1,3,3-tetramethylcyclohexane

Note that the bromine is furthest from the methyl groups due to destabilizing steric interactions. Out of the methylene groups available, this one is the easiest for the bromine to access.

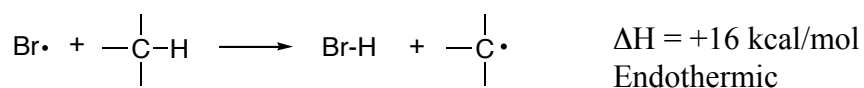
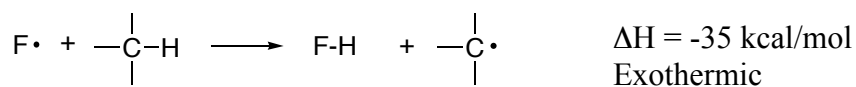
Reactivity and Selectivity (Hammond Postulate)

e.g. Halogenation of 2-methylbutane



A. Reactivity TREND:

F₂ > Cl₂ > Br₂ > I₂ Iodine does not react



B. Selectivity TREND:

Br \cdot > Cl \cdot > F \cdot

most selective	least selective
endothermic	exothermic

Bromine atom "searches" the molecule to create the most stable radical

Fluorine atom is small and feels the loss of an electron much more than bromine

- Fluorine is less precise and reacts immediately