

# **ALCOOLS**

**DÉFINITION**

**NOMENCLATURE**

**RÉACTIVITÉ**

**ACIDITÉ - RÉACTION AVEC LES BASES**

**FORMATION ET RÉACTION DES ALCOOLATES**

**ACTION DES ACIDES MINÉRAUX**

**ACIDES HALOGÉNÉS -  $\text{H}_2\text{SO}_4$  ,  $\text{H}_3\text{PO}_4$**

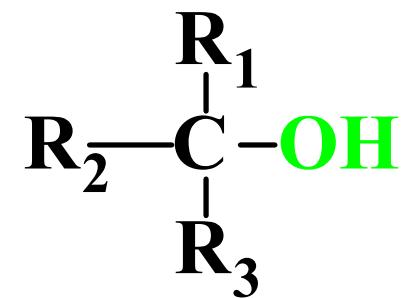
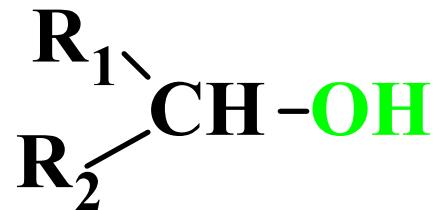
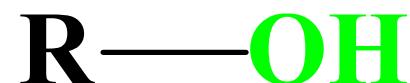
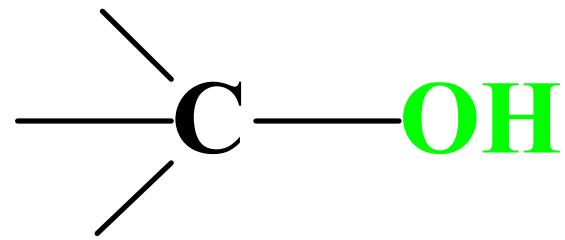
**RÉACTIVITÉ DÛE À LA NUCLÉOPHILIE DE L'OXYGÈNE**

**RÉACTION AVEC  $\text{PX}_3$  ou  $\text{SOCl}_2$**

**FORMATION DES SULFONATES**

**OXYDATION**

# ALCOOLS



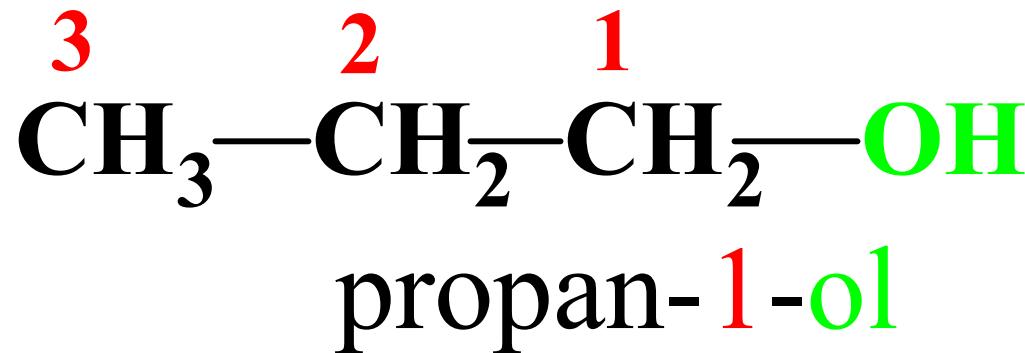
alcool primaire

alcool secondaire

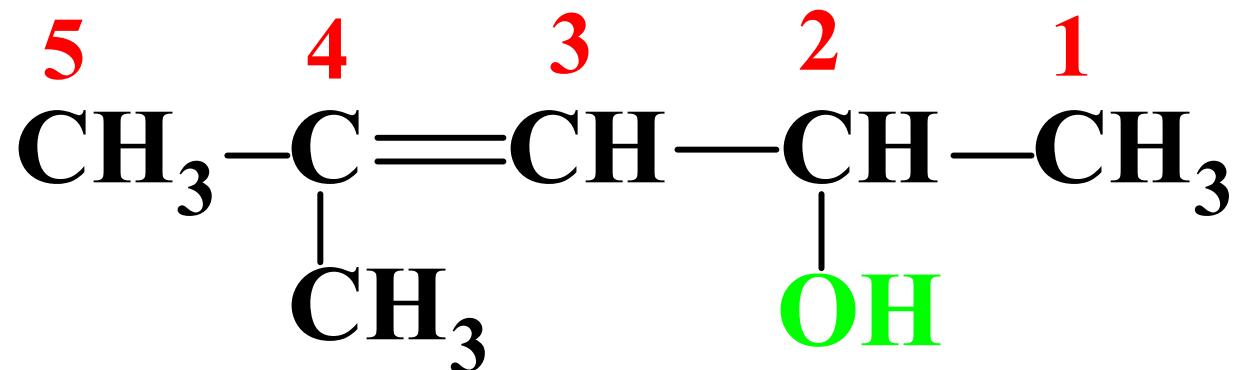
alcool tertiaire

$\text{R}, \text{R}_1, \text{R}_2, \text{R}_3$  = radical hydrocarboné

# NOMENCLATURE

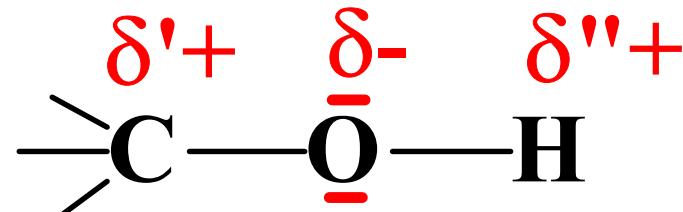


hydrocarbure    **indice de position**    ol



4-méthylpent-3-èn-2-ol

# RÉACTIVITÉ DES ALCOOLS



- ✿  $\begin{array}{c} \delta - \quad \delta'' + \\ \diagup \quad \diagdown \\ -\text{O}-\text{H} \end{array}$  ← attaque par les bases fortes →  $\text{RO}^\ominus$   
 $\text{RO}^\ominus$  = nucléophile fort, base forte
- ✿  $\begin{array}{c} \delta - \\ \diagup \\ -\text{O}- \end{array}$   $\text{ROH}$  = nucléophile faible, base faible
- ✿  $\begin{array}{c} \delta' + \quad \delta - \\ \diagup \quad \diagdown \\ \text{C} - \text{O} \text{H} \end{array}$  substitution, élimination

MAIS OH "MAUVAIS GROUPE PARTANT

# **ROH RÉACTIF NUCLÉOPHILE**



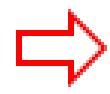
**ADDITION SUR C INSATURÉ**



**DÉRIVÉS CARBONYLÉS**

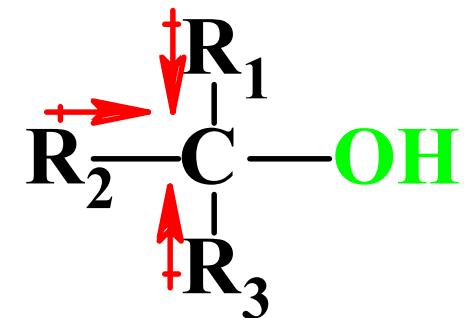
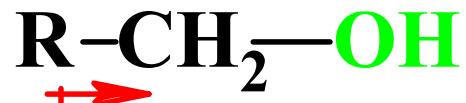


**SUBSTITUTION SUR C INSATURÉ**



**ACIDES ET DÉRIVÉS**

# ACIDITÉ DES ALCOOLS



pK - 16

pK - 19

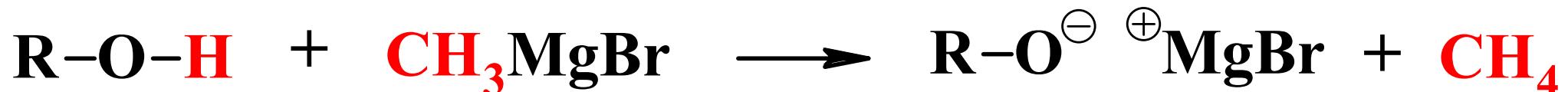


# FORMATION DES ALCOOLATES

## ◆ Action des bases

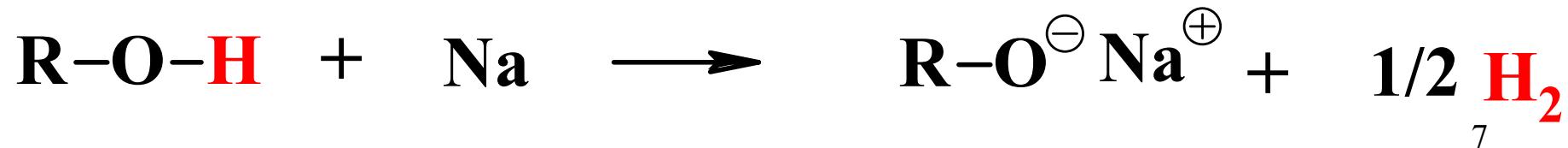


$$\text{pK ROH / RO}^- - 17 \quad \text{pK NH}_3 / \text{NH}_2^- - 35$$



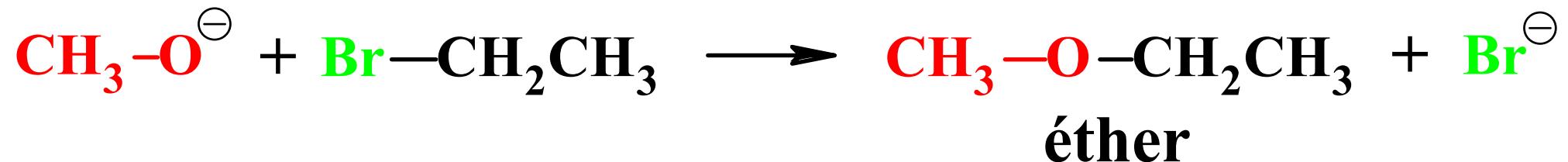
$$\text{pK CH}_4 / \text{CH}_3^- - 50$$

## ◆ Action des métaux alcalins (Na, K)



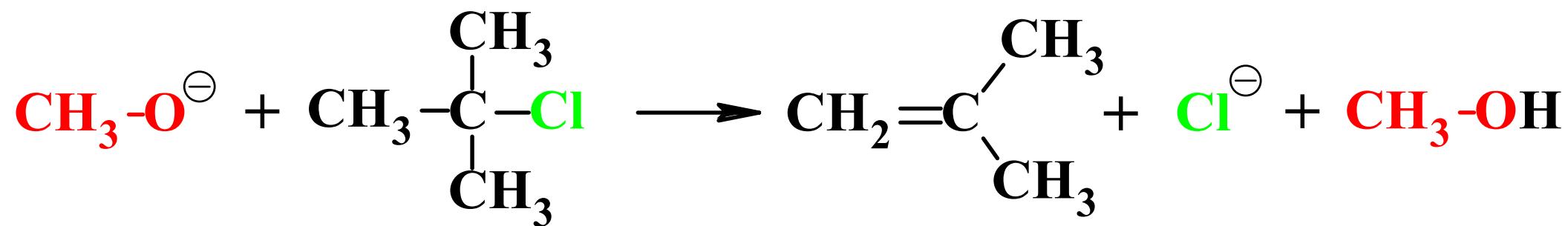
# RÉACTION DES ALCOOLATES

◆ Nucléophile  $\Rightarrow$  Substitution

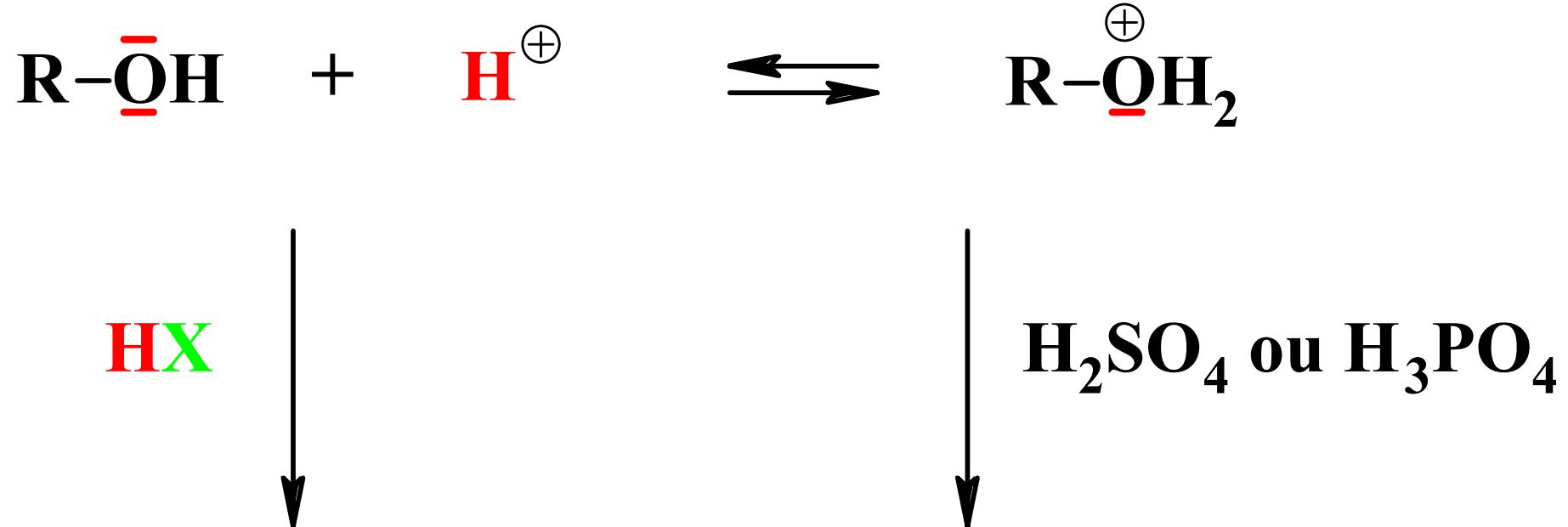


Réaction de WILLIAMSON

◆ Base  $\Rightarrow$  Élimination



# ACTION DES ACIDES MINÉRAUX



◆ SUBSTITUTION

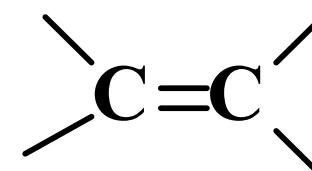


dérivé halogéné

◆ SUBSTITUTION



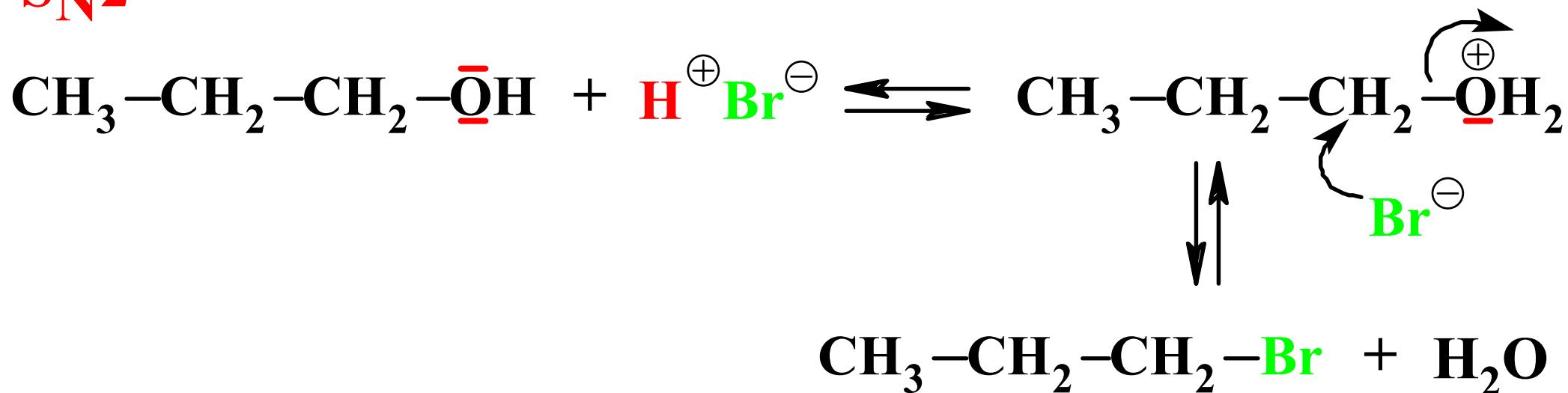
◆ ÉLIMINATION



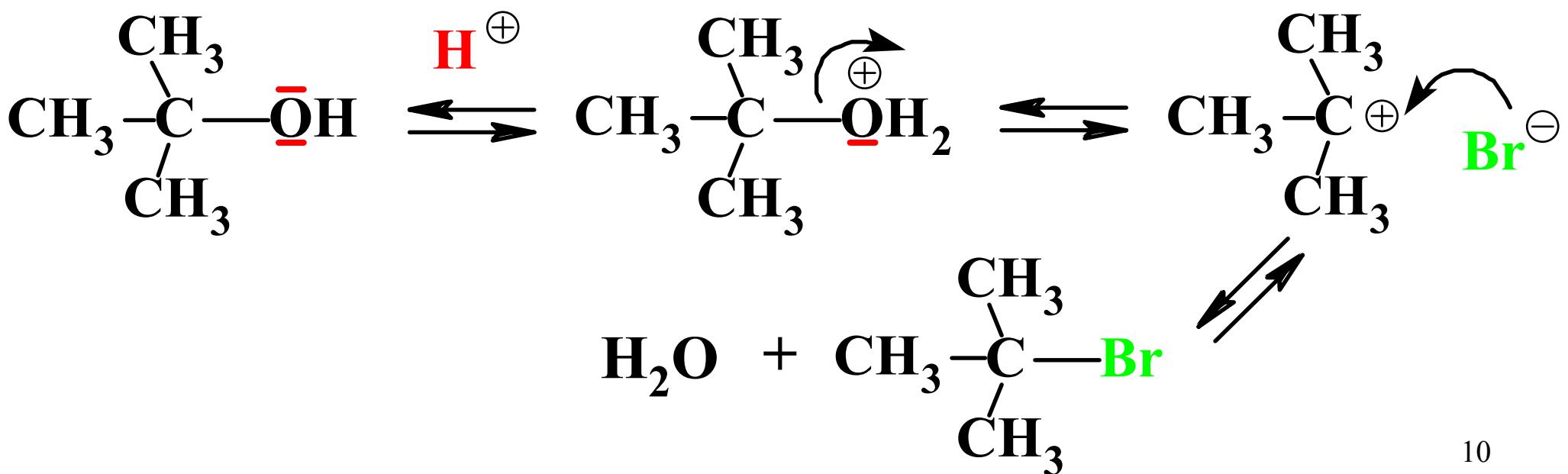
alcène

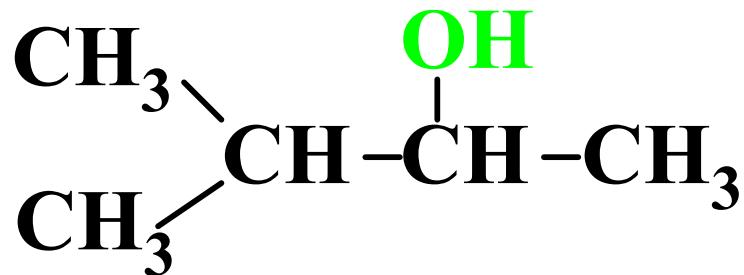
# ACTION DE HCl - HBr - HI

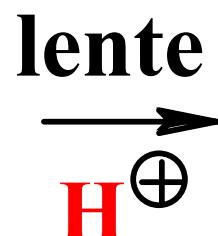
**S<sub>N</sub>2**

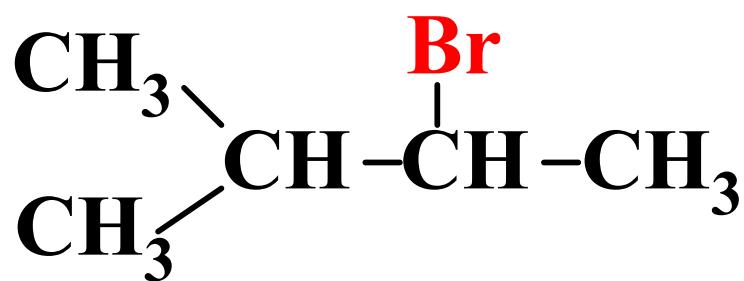
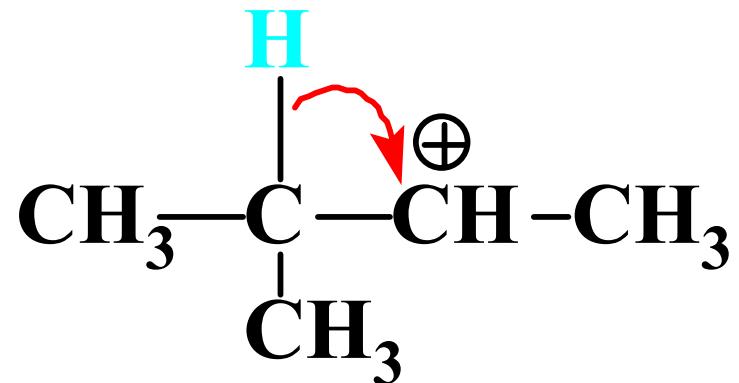


**S<sub>N</sub>1**

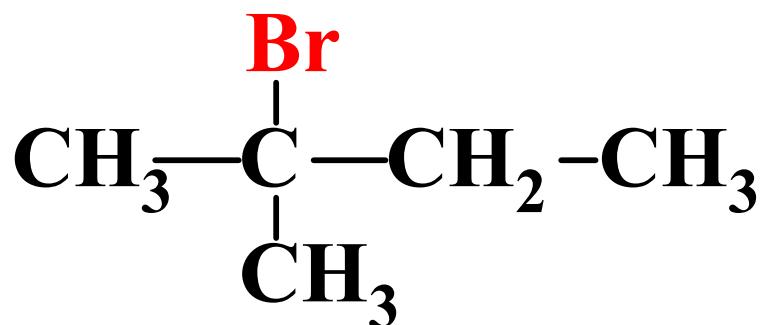
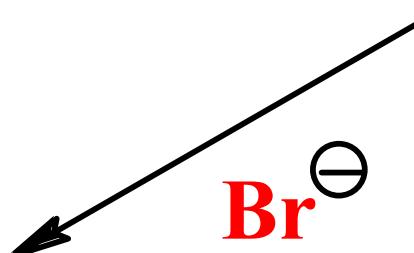




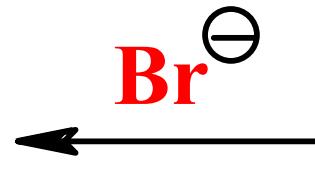
lente  


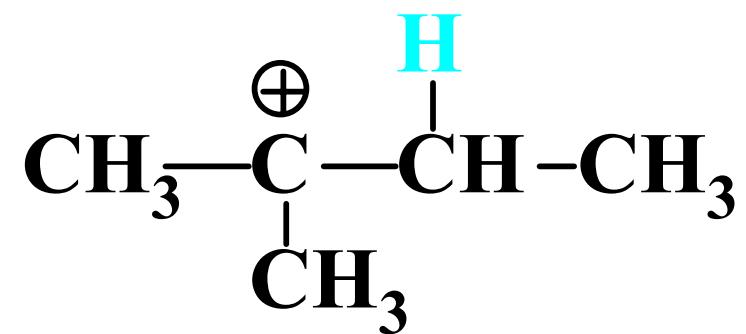


minoritaire

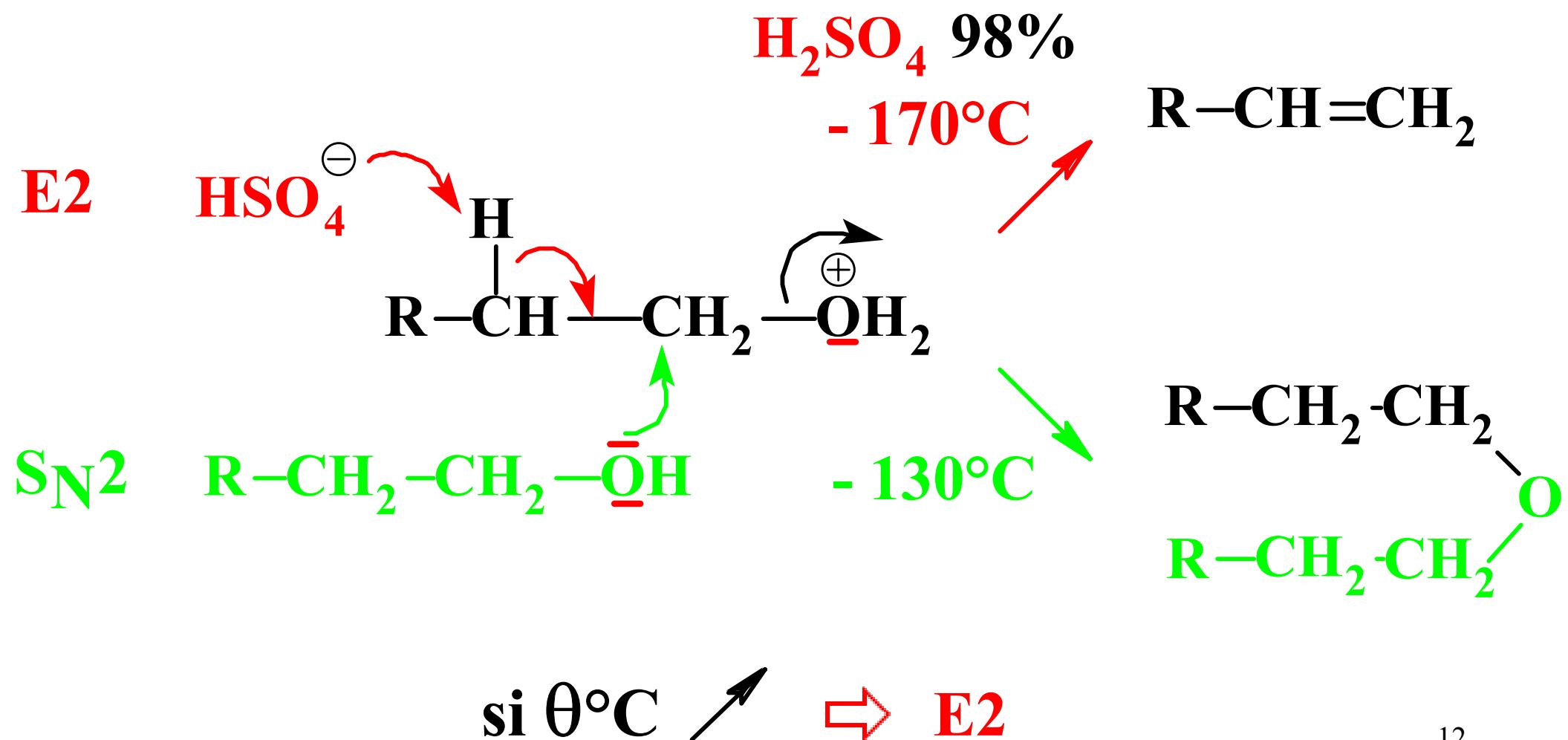
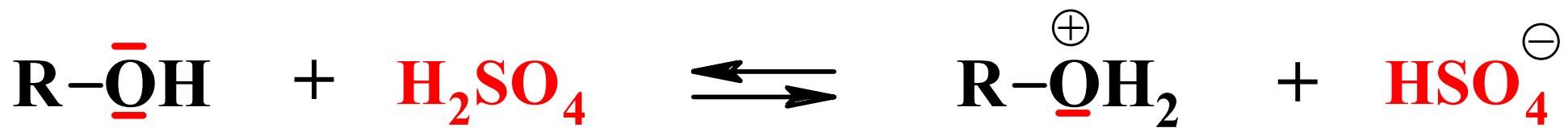


majoritaire

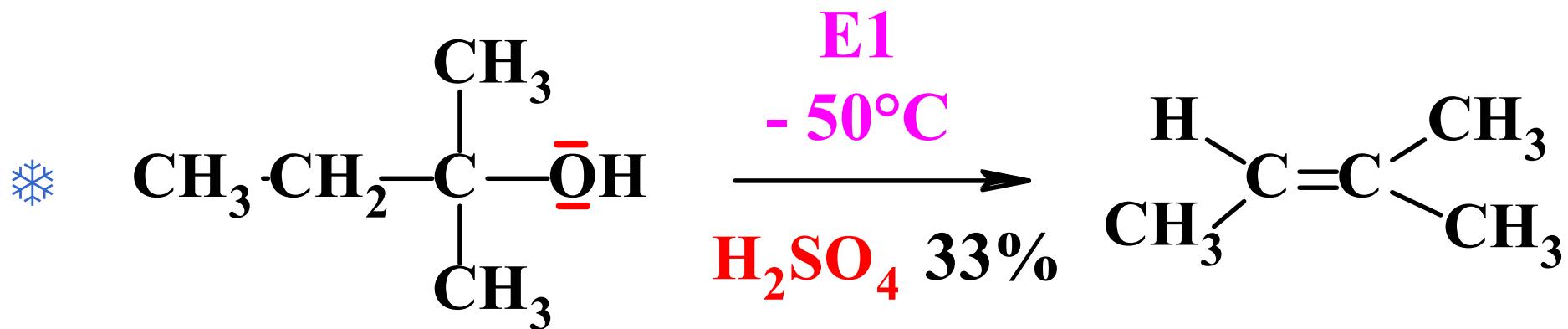
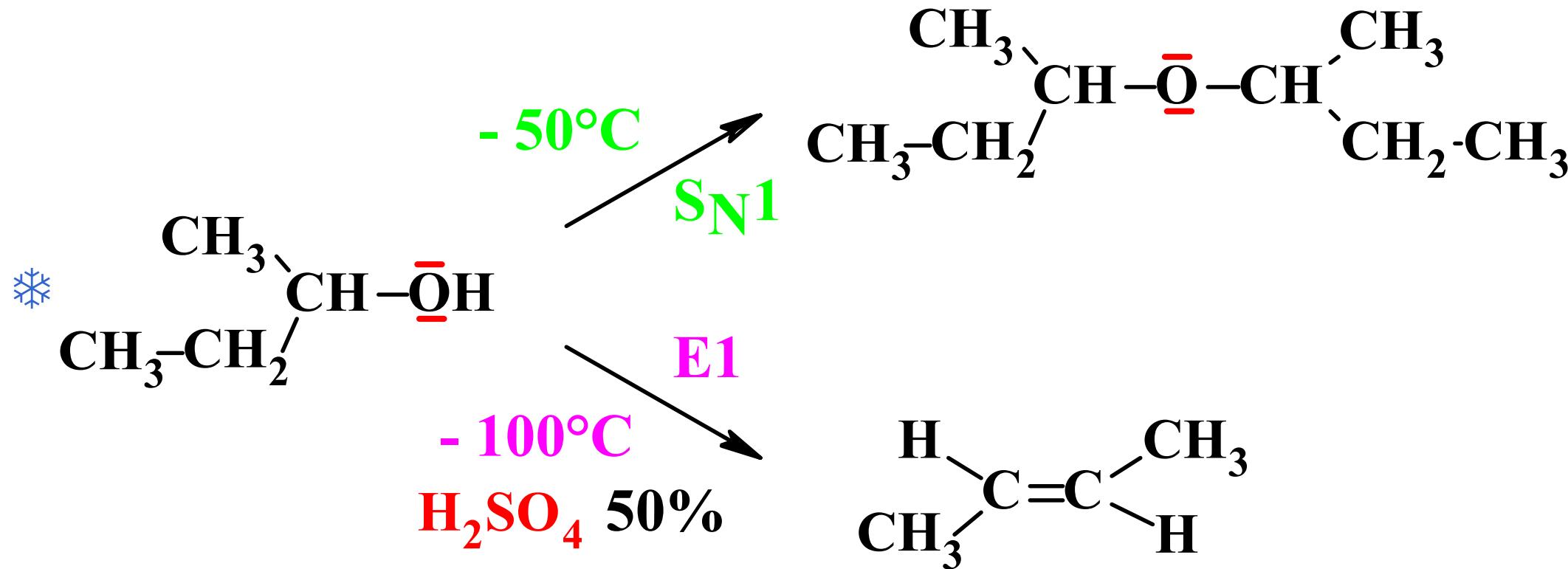




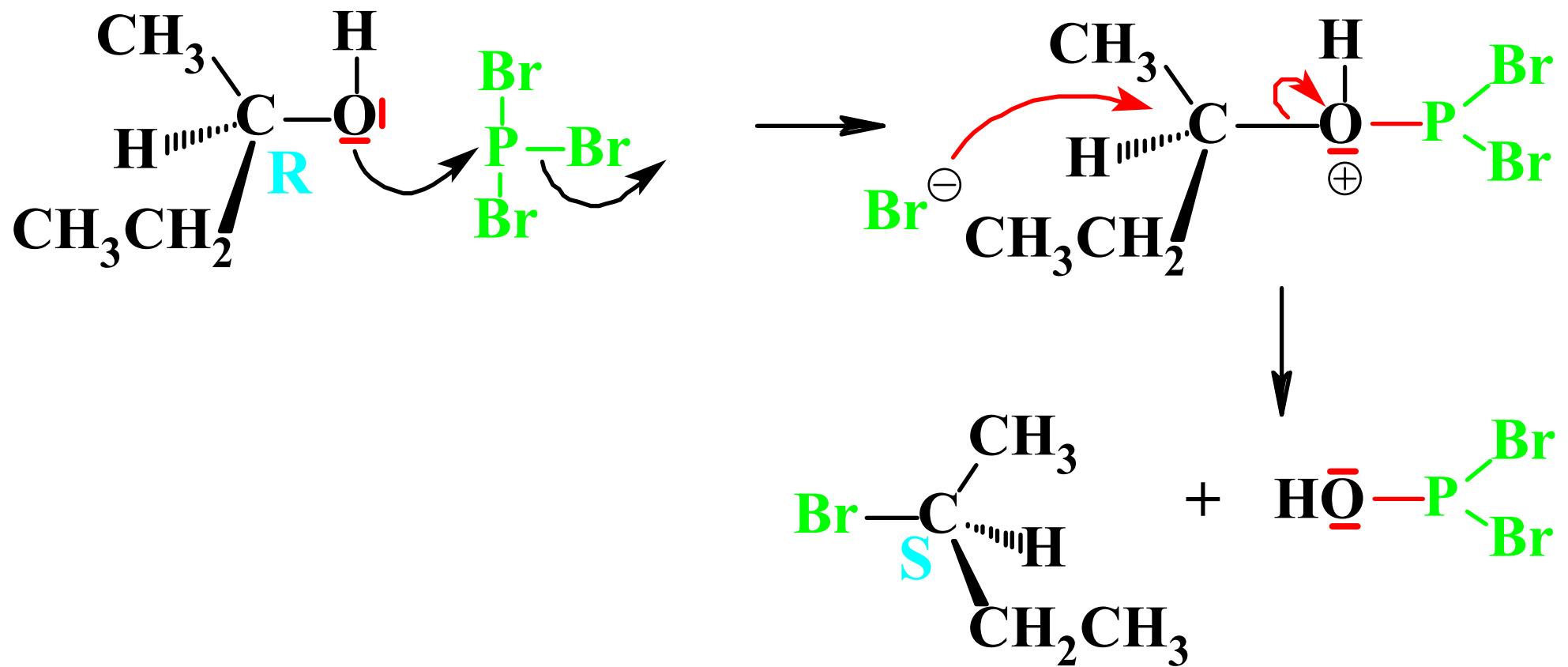
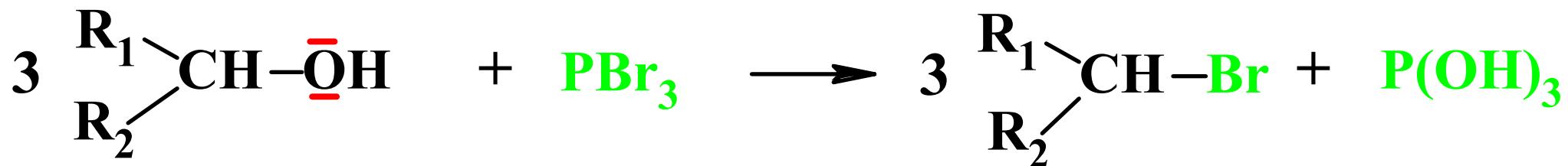
# ACTION DE $\text{H}_2\text{SO}_4$ ou $\text{H}_3\text{PO}_4$



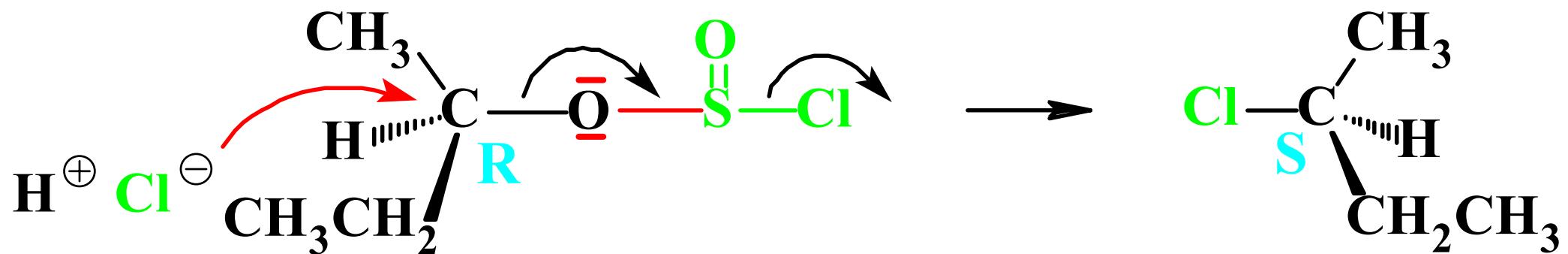
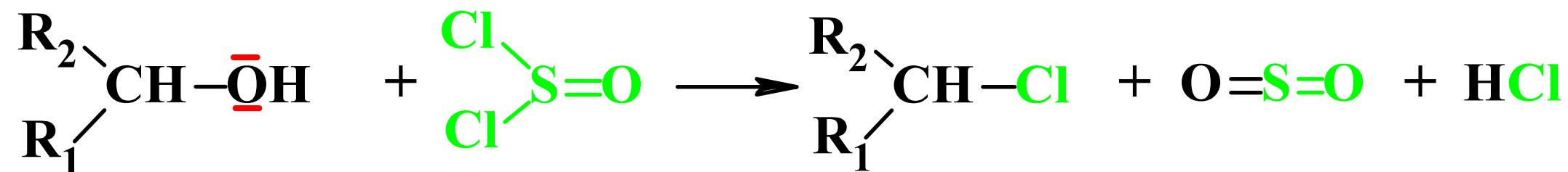
# ACTION DE H<sub>2</sub>SO<sub>4</sub> ou H<sub>3</sub>PO<sub>4</sub>



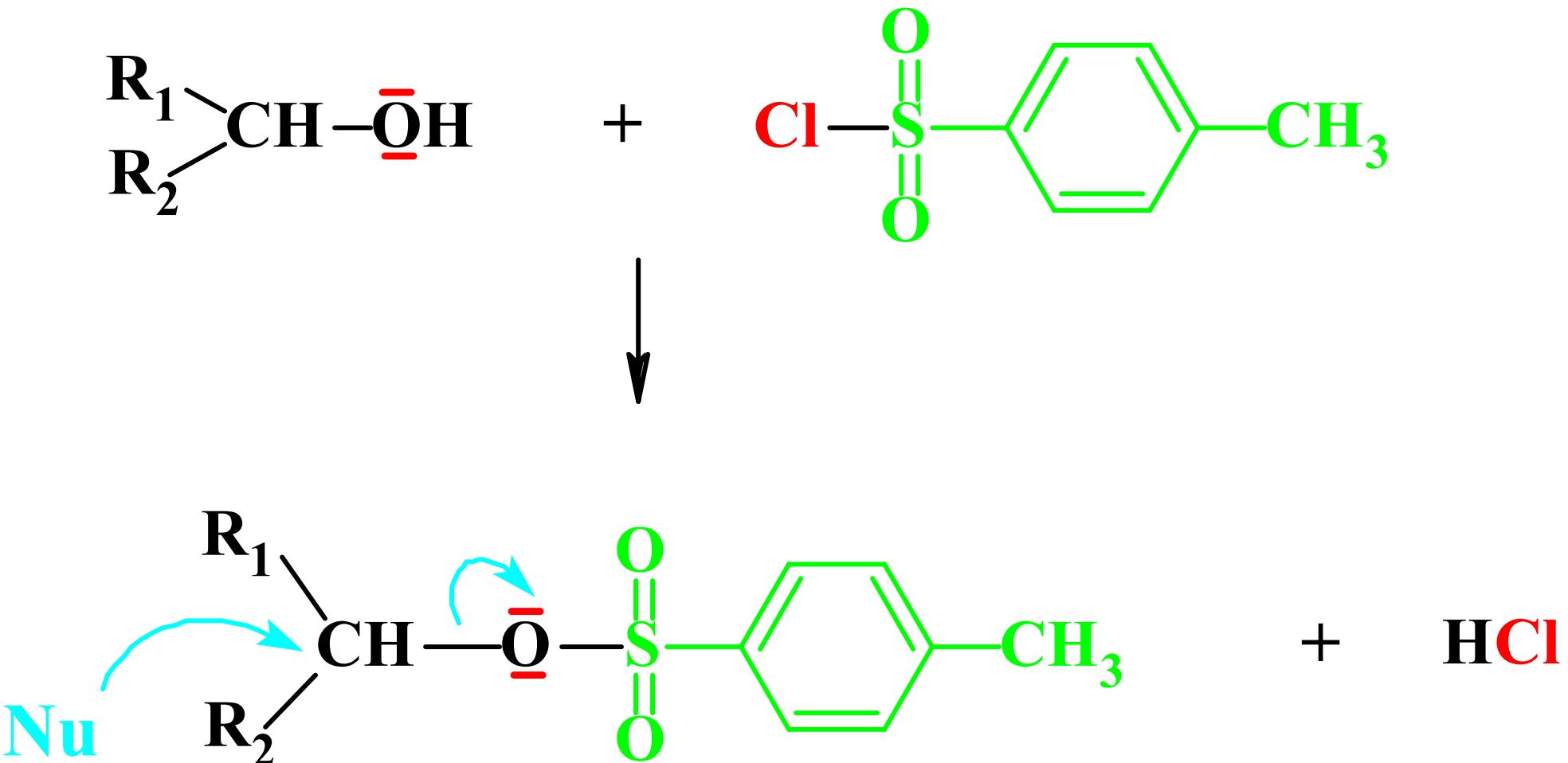
## ACTION DE $\text{PCl}_3$ ou $\text{PBr}_3$



## ACTION DE $\text{SOCl}_2$ (CHLORURE DE THIONYLE)

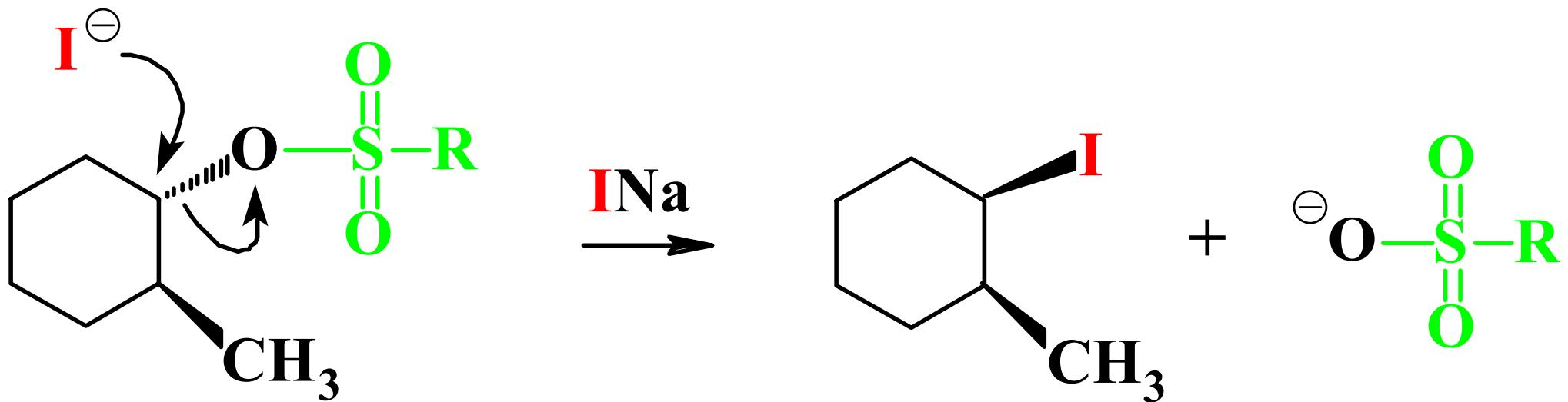
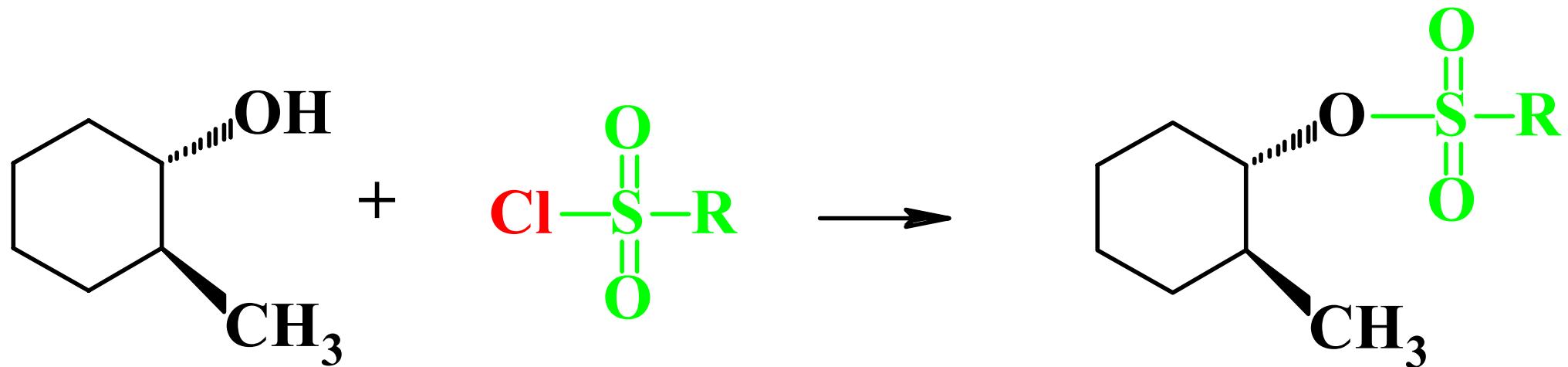


# ESTER SULFONIQUE (SULFONATE)

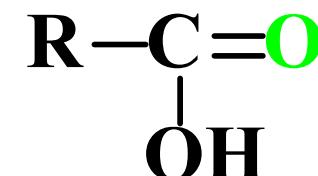
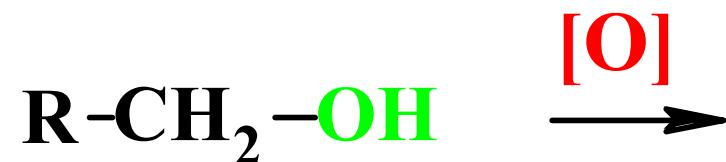


SUBSTITUTION    NUCLÉOPHILE  
ÉLIMINATION

# RÉACTIVITÉ DES SULFONATES

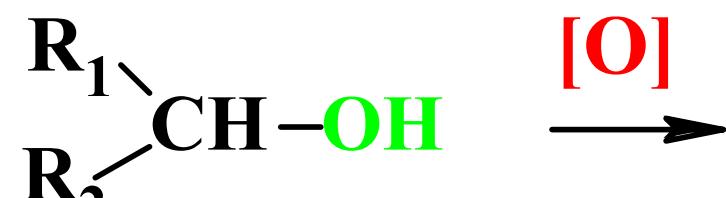


# OXYDATION DES ALCOOLS

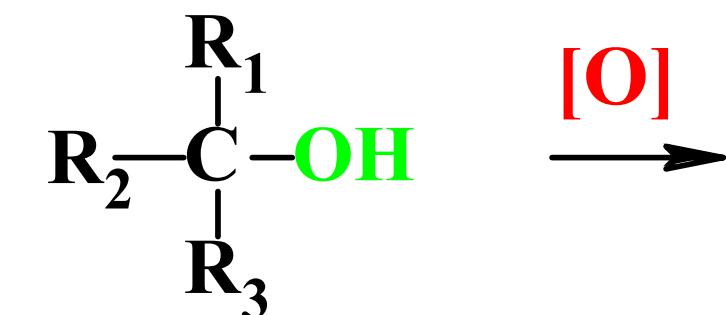


aldéhyde

acide



cétone



rien ou composés de dégradation

$\text{KMnO}_4$

$\text{K}_2\text{CrO}_7$

$\text{H}_2\text{CrO}_4$

# OXYDATION DES ALCOOLS

