3A ORGANIC REACTIONS

ALCOHOL OXIDATION

* 1° are oxidized to either aldehydes or carboxylic acids, depending on the strength of oxidiser, while 2° produce ketones. 3° alcohols are **unreactive** with laboratory oxidisers such as dichromate and permanganate but can of course be combusted in oxygen given suitable *activation energy*.



* With 1° alcohol oxidation mild oxidizers such as Cr₂O₇²- may produce an intermediate aldehyde.

● Under extreme oxidation, such as in *COMBUSTION*, all organic compounds will be oxidised through to CO₂ and H₂O!

1st Principles:

Semi-Structure:

$$\Rightarrow 5(CH_3CH_2CH_2OH + H_2O \rightarrow CH_3CH_2COOH + 4H^+ + 4e^-)$$

Add Inorganic Reduction half equation for the particular Oxidising Agent

$$\Rightarrow \qquad 4(MnO_4^- + 8H^+ + 5e^- \rightarrow Mn^{+2} + 4H_2O)$$

$$5CH_3CH_2OH + 4MnO_4^- + 12H^+ \rightarrow 5CH_3CH_2COOH + 4Mn^{+2} + 11H_2O$$

 $\Rightarrow \underline{5CH_3CH_2CH_2OH_{(1)} + 4MnO_4}\underline{\cdot}_{(aq)} + \underline{12H^+_{(aq)}} \rightarrow \underline{5CH_3CH_2COOH_{(aq)} + 4Mn^{+2}_{(aq)} + 11H_2O_{(1)}}$

1st Principles:

Semi-Structure:

$$\Rightarrow \qquad 3(CH_3CHOHCH_2CH_3 \quad \rightarrow \quad CH_3COCH_2CH_3 \ + \ 2H^+ + \ 2e^- \)$$

Add Inorganic Reduction half equation for the particular Oxidising Agent

$$\Rightarrow \frac{\text{Cr}_2\text{O}_7^{2-} + 14\text{H}^+ + 6\text{e}^- \rightarrow 2\text{Cr}^{3+} + 7\text{H}_2\text{O}}{8}$$

$$3CH_{3}CHOHCH_{2}CH_{3} \ + \ Cr_{2}O_{7}{^{2-}} \ + \ 44H^{+} \ \rightarrow \ 3CH_{3}COCH_{2}CH_{3} \ + \ 2Cr^{3+} \ + \ 6\cdot H^{+} \ + \ 7H_{2}O^{-}$$

$$3CH_{3}CHOHCH_{2}CH_{3} \ + \ Cr_{2}O_{7}{}^{2\text{-}} \ + \ 8H^{+} \ \rightarrow \ 3CH_{3}COCH_{2}CH_{3} \ + \ 2Cr^{3+} \ + \ 7H_{2}O$$

$$\Rightarrow \underline{3CH_3CHOHCH_2CH_{3 (l)} + Cr_2O_7{}^{2-}}_{(aq)} + \underline{8H^+}_{(aq)} \rightarrow \underline{3CH_3COCH_2CH_{3 (l)} + 2Cr^{3+}}_{(aq)} + \underline{7H_2O_{(l)}}_{(l)}$$

CARBOXYLIC ACID NEUTRALISATION

* As with all acids an organic acid can be neutralised by base to leave a **salt** and **water**.

Despite the "weak" nature of many organic acids, they neutralise base just as effectively as a "strong" acid. The base assists in removal of the acid proton, it does not have to simply ionise itself!

$$-\overset{\vdash}{C} - \overset{\vdash}{C} - \overset{\circlearrowleft}{C} + \overset{\lor}{Na} \overset{\circlearrowleft}{OH} \rightarrow -\overset{\vdash}{C} - \overset{\circlearrowleft}{C} - \overset{\circlearrowleft}{C} - \overset{\circlearrowleft}{Na} \overset{}{\bullet} + \overset{H_2O}{\bullet}$$

$$\Rightarrow \frac{\text{CH}_3\text{CH}_2\text{COOH}_{(aq)} + \text{OH}^{-}_{(aq)}}{\text{(Propanoate ion)}} \rightarrow \frac{\text{CH}_3\text{ CH}_2\text{COO}^{-}_{(aq)} + \text{H}_2\text{O}_{(1)}}{\text{(Propanoate ion)}}$$

* NB. Spectator ion omitted (Na⁺). Being a weak acid the molecular species is the most abundant chemical species so you should not cancel the acid to H⁺!

ESTERIFICATION

 \Rightarrow

* In the presence of an **acid catalyst ALCOHOLS** and **CARBOXYLIC ACIDS** can combine to form esters. These esters have a fruity smell.

$$-\frac{1}{C} - \frac{1}{C} - \frac{1$$

 $\frac{\text{CH}_3\text{COOH}_{(1)} + \text{CH}_3\text{CH}_2\text{OH}_{(1)}}{\text{(Ethyl Ethanoate)}} \rightarrow \frac{\text{CH}_3\text{COOCH}_2\text{CH}_3_{(1)} + \text{H}_2\text{O}_{(1)}}{\text{(Ethyl Ethanoate)}}$

REVERSE ESTERIFICATION- Base + Ester

* As esterification is an **acid catalysed** process it can be reversed by the addition of a base to an ester.

$$-\frac{1}{c} - \frac{1}{c} - \frac{1}{c} - \frac{1}{c} - \frac{1}{c} - \frac{1}{c} + \frac{1}{Na} \underbrace{OH}$$

$$+ \frac{1}{c} - \frac{1}$$

- $\Rightarrow \qquad CH_{3}CH_{2}COOCH_{2}CH_{3\;(l)}\;+\;NaOH_{\;(aq)}\;\;\rightarrow \qquad CH_{3}CH_{2}COONa_{(aq)} \quad +\;CH_{3}CH_{2}OH_{\;(l)}$
- $\Rightarrow \qquad \underline{\text{CH}_3\text{CH}_2\text{COOCH}_2\text{CH}_3\text{ (i)} + \text{OH}^2\text{ (aq)}} \rightarrow \qquad \underline{\text{CH}_3\text{CH}_2\text{COO}^2\text{ (aq)}} + \underline{\text{CH}_3\text{CH}_2\text{OH}}\text{ (i)}$

ADDITION

* Hydrocarbons with UNSATURATED parts (double or triple bonds) can be added onto producing saturated molecules:

 $\underline{\text{CH}_2\text{CHCH}_3}_{(g)} + \underline{\text{Br}_2}_{(l)} \rightarrow \underline{\text{CH}_2\text{BrCHBr CH}_3}_{(l)}$

SUBSTITUTION

* **SATURATED** hydrocarbons (alkanes) or aromatic compounds (benzene) cannot undergo addition, only the interchange or *substitution* of parts of their structure:

 $\Rightarrow \qquad \begin{array}{c} \text{hv} \\ \Rightarrow \qquad \underline{\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_3}_{(g)} + \underline{\text{Cl}_2}_{(g)} \rightarrow \underline{\text{CH}_3\text{CH}_2\text{CH}_2\text{CH}_2\text{Cl}_{(l)}} + \underline{\text{HCl}_{(g)}} \\ \end{array}$

OR

 $\Rightarrow \qquad \frac{\text{hv}}{\text{C}_6\text{H}_6.\text{(1)}} + \frac{\text{Cl}_2}{\text{O}} \rightarrow \frac{\text{C}_6\text{H}_5\text{Cl}.\text{(1)}}{\text{Cl}} + \frac{\text{HCl}}{\text{HCl}}$

COMBUSTION

- * All Hydrocarbons and carbohydrates will burn to produce carbon dioxide and water.
 - \Rightarrow <u>2CH₃CH₂CH₂CH₃(g) + 13 O₂(g) \rightarrow 8 CO₂(g) + 10 H₂O (g)</u>

