|  | Statement | Guidance |
| :---: | :---: | :---: |
| 8.1.U1 | A Brønsted-Lowry acid is a proton/ $\mathrm{H}+$ donor and a Brønsted-Lowry base is a proton/ $\mathrm{H}+$ acceptor. | The location of the proton transferred should be clearly indicated. For example, $\mathrm{CH}_{3} \mathrm{COOH} / \mathrm{CH}_{3} \mathrm{COO}^{-}$rather than $\mathrm{C}_{2} \mathrm{H}_{4} \mathrm{O}_{2} / \mathrm{C}_{2} \mathrm{H}_{3} \mathrm{O}_{2}^{-}$ |
| 8.1.U2 | Amphiprotic species can act as both Brønsted-Lowry acids and bases | The difference between the terms amphoteric and amphiprotic should be covered |
| 8.1.U3 | A pair of species differing by a single proton is called a conjugate acid-base pair | Students should know the representation of a proton in aqueous solution as both $\mathrm{H}^{+}(\mathrm{aq})$ and $\mathrm{H}_{3} \mathrm{O}^{+}(\mathrm{aq})$. |
| 8.2.U4 | Most acids have observable characteristic chemical reactions with reactive metals, metal oxides, metal hydroxides, hydrogen carbonates and carbonates. | Bases which are not hydroxides, such as ammonia, soluble carbonates and hydrogen carbonates should be covered |
| 8.2.U5 | Salt and water are produced in exothermic neutralization reactions |  |
| 8.3.U6 | $\mathrm{pH}=\log \left[\mathrm{H}_{\text {(ar) }}\right]$ and $[\mathrm{H}]=10{ }^{-\mathrm{pH}}$. |  |
| 8.3.U7 | A change of one pH unit represents a 10 -fold change in the hydrogen ion concentration $[\mathrm{H}]$. |  |
| 8.3.U8 | pH values distinguish between acidic, neutral and alkaline solutions. | Students should be concerned only with strong acids and bases in this sub-topic |
| 8.3.U9 | The ionic product constant, $\mathrm{K}_{\mathrm{w}}=\left[\mathrm{H}^{+}\right][\mathrm{OH}]=10^{-14}$ at 298 K . | Equations involving $\mathrm{H}_{3} \mathrm{O}^{+}$instead of $\mathrm{H}+$ may be applied |
| 8.4.U10 | Strong and weak acids and bases differ in the extent of ionization. | The terms ionization and dissociation can be used interchangeably. |
| 8.4.U11 | Strong acids and bases of equal concentrations have higher conductivities than weak acids and bases |  |
| 8.4.U12 | A strong acid is a good proton donor and has a weak conjugate base |  |
| 8.4.U13 | A strong base is a good proton acceptor and has a weak conjugate acid. |  |
| 8.5.U14 | Rain is naturally acidic because of dissolved $\mathrm{CO}_{2}$ and has a pH of 5.6. Acid deposition has a pH below 5.6. |  |
| 8.5.U15 | Acid deposition is formed when nitrogen or sulfur oxides dissolve in water to form $\mathrm{HNO}_{3}, \mathrm{HNO}_{2}, \mathrm{H}_{2} \mathrm{SO}_{4}$ and $\mathrm{H}_{2} \mathrm{SO}_{3}$. |  |
| 8.5.U16 | Sources of the oxides of sulfur and nitrogen and the effects of acid deposition should be covered |  |
| 18.1.U17 | A Lewis acid is a lone pair acceptor and a Lewis base is a lone pair donor | Both organic and inorganic examples should be studied. |
| 18.1.U18 | When a Lewis base reacts with a Lewis acid a coordinate bond is formed. |  |
| 18.1.U19 | A nucleophile is a Lewis base and an electrophile is a Lewis acid. | Relations between Brønsted-Lowry and Lewis acids and bases should be discussed |
| 18.2.U20 | The expression for the dissociation constant of a weak acid $\left(K_{a}\right)$ and a weak base ( $K_{b}$ ). | Only examples involving the transfer of one proton will be assessed. |
| 18.2.U21 | For a conjugate acid base pair, $K_{\mathrm{a}} \times K_{\mathrm{b}}=K_{\mathrm{w}}$ | The value $K_{w}$ depends on the temperature. Students should state when approximations are used in equilibrium calculations |
| 18.2.U22 | The relationship between $K_{\mathrm{a}}$ and $\mathrm{p} K_{\mathrm{a}}$ is $\left(\mathrm{p} K_{\mathrm{a}}=-\log K_{\mathrm{a}}\right)$, and between $K_{\mathrm{b}}$ and $\mathrm{p} K_{\mathrm{b}}$ is $\left(\mathrm{p} K_{\mathrm{b}}=-\log K_{\mathrm{b}}\right)$ | Calculations of pH at temperatures other than 298 K can be assessed |
| 18.3.U23 | The characteristics of the pH curves produced by the different combinations of strong and weak acids and bases. | Only examples involving the transfer of one proton will be assessed. Important features are: <br> - intercept with pH axis <br> - equivalence point <br> - buffer region <br> - points where $\mathrm{p} K_{\mathrm{a}}=\mathrm{pH}$ or $\mathrm{pK}_{\mathrm{b}}=\mathrm{pOH}$. |
| 18.3.U24 | An acid-base indicator is a weak acid or a weak base where | For an indicator which is a weak acid: |


|  | the components of the conjugate acid-base pair have different colours | $\begin{aligned} & -\mathrm{HIn}(\mathrm{aq}) \neq \mathrm{H}+(\mathrm{aq})+\operatorname{In}-(\mathrm{aq}) \\ & \text { Colour A Colour B } \\ & \text { - The colour change can be considered to take } \\ & \text { place over a range of pKa } \pm 1 \text {. } \\ & \text { For an indicator which is a weak base: } \\ & -\mathrm{BOH}(\mathrm{aq}) \rightleftarrows \mathrm{B}+(\mathrm{aq})+\mathrm{OH}-(\mathrm{aq}) \\ & \text { Colour A } \quad \text { Colour B } \end{aligned}$ |
| :---: | :---: | :---: |
| 18.3.U25 | The relationship between the pH range of an acid-base indicator, which is a weak acid, and its $p K_{a}$ value. |  |
| 18.3.U26 | The buffer region on the pH curve represents the region where small additions of acid or base result in little or no change in pH. |  |
| 18.3.U27 | The composition and action of a buffer solution |  |

