

- M1.** (a) Hydrogen bonding (1)
between H₂O and NH₃ (1) 2
- (b) (i) $\text{NH}_3 + \text{H}_2\text{O} \rightleftharpoons \text{NH}_4^+ + \text{OH}^-$ (1)
(ii) Ammonia is weak base (1)
NOT partially ionised
Equilibrium to left or incomplete reaction (1) 3
- (c) A proton donor (1) 1
- (d) *Buffer solution*: A solution which resists change in pH (1)
when small amounts of acid or base added or on dilution (1)
Reagent: NH₄Cl (1)
Allow a correct strong acid 3
- (e) (i) $K_a = \frac{[\text{H}^+][\text{A}^-]}{[\text{HA}]}$ (1)
 $= \frac{[\text{H}^+][0.125 \times 4]}{1.00}$ (1)
 $[\text{H}^+] = 1.70 \times 10^{-5} / 0.125 \times 4 = 3.40 \times 10^{-5}$ (1)
 $\text{pH} = -\log_{10} [\text{H}^+] = 4.47$ (1)
Allow pH conseq to [H⁺] if 2 place decimals given
(ii) $\text{H}^+ + \text{CH}_3\text{COO}^- \rightarrow \text{CH}_3\text{COOH}$ (1) 5
- [14]**

M2.C

[1]

M3. Penalise pH given to 1 dp first time it would have scored only

(a) (i) $K_w = [\text{H}^+][\text{OH}^-]$ (1)

(ii) $\text{pH} = -\log [\text{H}^+]$ (1)

or in words or below unless contradiction

(iii) *Calculation:* $[\text{H}^+] = \sqrt{5.48 \times 10^{-14}}$ (1)

$$= 2.34 \times 10^{-7}$$

$$\therefore \text{pH} = 6.63 \text{ or } 6.64$$
 (1)

Explanation: pure water $\therefore [\text{H}^+] = [\text{OH}^-]$ (1)

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(b) (i) $[\text{OH}^-] = 0.150$ $\therefore [\text{H}^+] = 10^{-14}/0.15 = 6.66 \times 10^{-14}$

or $\text{pOH} = 0.82$

$$\therefore \text{pH} = 13.18$$
 (1)

or $\text{pH} = 13.17$

(ii) moles $\text{OH}^- = (35 \times 10^{-3}) \times 0.150 = 5.25 \times 10^{-3}$ (1)^a

moles $\text{H}^+ = (40 \times 10^{-3}) \times 0.120 = 4.8(0) \times 10^{-3}$ (1)^b

$$\therefore \text{excess moles of } \text{OH}^- = 4.5 \times 10^{-4}$$
 (1)^c

$$\therefore [\text{OH}^-] = (4.5(0) \times 10^{-4}) \times 1000/\underline{75}^d$$
 (1)^e
$$= 6.0(0) \times 10^{-3}$$

$$[\text{H}^+] = \frac{10^{-14}}{6.00 \times 10^{-3}} = 1.66 \times 10^{-12} \text{ or } \text{pOH} = 2.22$$

$$\therefore \text{pH} = 11.78$$
 (1)^f
or 11.77

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(c) (i) $K_a = \frac{[H^+][X^-]}{[HX]} \quad (1)$

(ii) $[H^+] = 1.80 \times 10^{-2} \times 0.150 = 2.70 \times 10^{-3} \quad (1)$

$$K_a = \frac{[H^+]^2}{[HX]} (1) = \frac{(2.70 \times 10^{-3})^2}{0.150} = 4.86 \times 10^{-5} \quad (1) \text{ mol dm}^{-3} \quad (1)$$

$$\text{or } \frac{(2.70 \times 10^{-3})^2}{0.1473} = 4.95 \times 10^{-5}$$

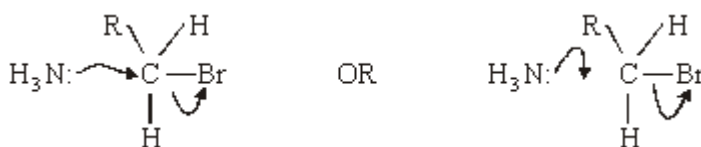
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Notes

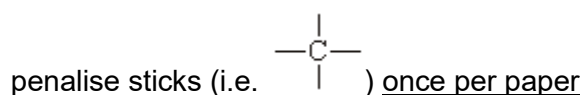
- (a) If K_w includes H_2O allow 6.63 if seen otherwise no marks likely
- (b) (ii) If no vol, max 4 for a, b, c, f answer = 10.65
 If wrong volume max 5 for a, b, c, e, f
 If no subtraction max 3 for a, b, d
 If missing 1000 max 5 for a, b, c, d, f answer = 8.78
 If uses excess as acid, max 4 for a, b, d, f answer = 2.22
 If uses excess as acid and no volume, max 2 for a, b answer = 3.35
- (c) If wrong K_a in (i) max 2 in part (ii) for $[H^+]$ (1) and conseq units (1)
but mark on fully from minor errors
eg no [] or charges missing

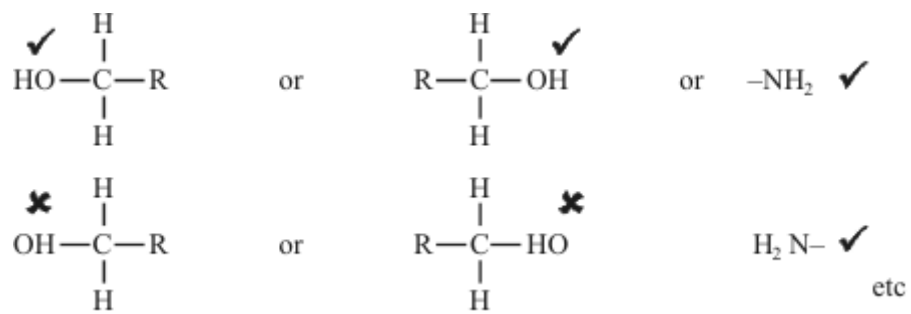
Organic points

- (1) Curly arrows: must show movement of a pair of electrons, i.e. from bond to atom or from lp to atom / space e.g.



- (2) Structures





Penalise once per paper

allow CH_3- or $-\text{CH}_3$ or $\begin{array}{c} \text{CH}_3 \\ | \end{array}$ or CH_3
 or $\text{H}_3\text{C}-$

[18]