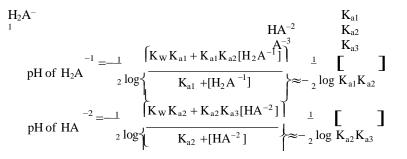
particles to come close thus inhibits the formation of larger aggregates

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titration is one with pKa close to the equivalence point pH

Preview from Notesale.co.uk Page 17 of 46 endpoint with the volume recorded as $V_{0\mbox{-}Ph}.$ On the same solution, methylred is then added and an additional volume is required to reach the end point recorded as $V_{Ph\mbox{-}MR}$

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- Step 1: Assign oxidation numbers to each of the species in the reaction NO_2 MnO_4^{-1} ⇇ NO₃ MnO₂ + +3 - 2+7 - 2+5 -2+4
- Step 2: Identify oxidation and reduction reactions and indicate the number of electrons lost or gained, respectively

| Oxidation: | NO ₂ ⁻¹ | ⇆ | NO_3^{-1} | + | 2e- |
|------------|--|---|-------------|---|---------|
| Reduction: | $^{+3}$ MnO ₄ ⁻¹ | + | +5 3e- | ₽ | MnO_2 |
| | +7 | | | | +4 |

Step 3: Balance the reaction by multiplying a factor on both sides of the reaction so that the numbers of electrons on both reactions are the same 3 NO_2^{-1} ← $3 NO_3^{-1}$ Ovidation x3. 60 +

| | Oxfuation 2 | () . | +3 | - | +5 | т | 06- | |
|---|-----------------------|-------------|------------------------|---|-----------------------|---|-------------------|--|
| | Reduction | ×2: | 2 MnO_4^{-1} | + | 6e- | ⇆ | 2 MnO_2 | |
| _ | | | +7 | | | | +4 | |
| | 3 NO_2^{-1} | + | 2 MnO_4^{-1} | 与 | 3 NO_3^{-1} | + | 2 MnO_2 | |

- Step 4: Balance the charges (by adding H⁺ or HO⁻) and number of hydrogen and/or oxygen atoms (by adding H₂O) on both sides of the equation
- In acidic medium, add H_2O to the oxygen-deficient side and supply H^+ to balance the hydrogen

 $3 \text{ NO}_2^{-1} + 2 \text{ MnO}_4^{-1} + 2 \text{ H}^+ \leftrightarrows 3 \text{ NO}_3^{-1} + 2 \text{ MnO}_2 + \text{H}_2\text{O}$

In basic medium, balance assuming reaction was in acidic medium. Neutralize H⁺ by adding HO⁻¹ on both sides of the reaction and simplify $3 \text{ NO}_2^{-1} + 2 \text{ MnO}_4^{-1} + 2 \text{ H}^+ + 2 \text{ HO}^{-1} \leftrightarrows 3 \text{ NO}_3^{-1} + 2 \text{ MnO}_2 + \text{ H}_2\text{ O} + 2 \text{ H}_2\text{ H} + 2 \text{ H} + 2 \text{ H}_2\text{ H} + 2 \text{ H} + 2 \text{ H} + 2 \text{ H} + 2 \text{$ \Rightarrow 3 NO₃⁻¹ + 2 MnO₂ + 10 $3 \text{ NO}_2^{-1} + 2 \text{ MnO}_4^{-1} +$ $2 H_2O$ 10^{-1} $+ 2 \text{ HO}^{-1}$ $3 \text{ NO}_2^{-1} + 2 \text{ MnO}_4^{-1} + \text{H}_2\text{O} \rightleftharpoons 3 \text{ NO}_3^{-1}$

3. Standard electrode potential

- The potential of a half-cell maction what the standard herrogen electrode (SHE) used as anode when the crivities of all reactant and products are taken as unity, that is, 1 M converting and 1 at 1 at a lat in sure
- Userly listed as standard reduction potential ($\varepsilon_{red}^{\circ}$) where a positive value implies that the located was used as a cathode and the SHE as anode High value of reduction potential indicates that the electrode is a good oxidizing
 - agent

Nernst equation

- Formulated by Walther Hermann Nernst (1864-1941)
- Accounts for the effect of concentration on electrode potentials

For a half-cell reduction reaction...Ox
$$\leftrightarrows$$
 Red + ne⁻⁻
 $\epsilon = \epsilon^{\circ} - \frac{\text{RT}}{\ln \frac{a_{\text{Red}}}{\ln \frac{a_{\text{RR}}}{\ln \frac{a_{\text{RR}}}{\ln$

where ε_{Red} = actual cell potential [V], $\varepsilon_{Red}^{\circ}$ = standard reduction potential [V], $R = 8.314 \text{ J-mol}^{-1}\text{-K}^{-1}$, T = temperature [K], n = number of electrons that appear in the half-cell reaction [mol], a = activity [] and F, Faraday's $constant = 96485.3399 coul-(mol e^{-})^{-1}$

At 25°C and for a given cell...

$$\varepsilon_{\text{cell}} = \varepsilon_{\text{cell}}^{\circ} - \frac{0.05916}{n} \log Q$$

h

$$COD = \underbrace{\left(V_{Fe^{+2}}^{mL,blank} - V_{Fe^{+2}}^{mL,sample} \right) \left(M_{Fe^{+2}} \right) \times \underbrace{\frac{1 \, \text{mmol} \, K_2 C r_2 O_7}{6 \, \text{mmol} \, Fe^{+2}} \times \underbrace{\frac{6 \, \text{mmol} \, O_2}{4 \, \text{mmol} \, K_2 C r_2 O_7}}_{1 \, \text{mmol} \, O_2} \times \underbrace{\frac{32 \, \text{mg} \, O_2}{1 \, \text{mmol} \, O_2}}_{1 \, \text{mmol} \, O_2}$$

volume sample (L)

Molecular Absorption Spectrometry

A. Absorption Process and Beer-Lambert Law

- If a beam of light passes through a glass container filled with liquid, the emergent radiation is always less powerful than that entering
- Consider a block of absorbing matter where a beam of monochromatic radiation of radiant power, P_0 strikes the surface perpendicularly and passes through the length of the material, **b**
- The emergent or transmitted radiation will always have less radiant power, **P** than the entering or incident radiation
- The fraction of incident radiation transmitted by the solution, **P/P**₀ is called transmittance and related to absorbance according to the equation:

A =
$$-\log T = -\log \frac{P}{P_0}$$

where A = absorbance, T = transmittance, $P_0 = incident$ radiant power [W], and P = transmitted radiant power [W]

• Beer-Lambert's law states that the absorbance is directly proportional to the concentration the best borbing species and to the path length

where $\varepsilon = \text{molar a sort vir (2-mol^{-1}-cm^{-1}]}$, b = path length [cm] and c = concentration [1 to 1-2-1]

B. Quantitative Analysis

1. Standard addition method

- Involves addition of several increments of a standard solution to aliquots of samples of the same size and the resulting solution upon adding the color development reagent is then diluted to a fixed volume (V_T) and measured for its absorbance
- Assume several identical aliquots of the unknown solution of volume V_x were treated with several increments of standard solution of volume V_s of known concentration C_s and diluted to a fixed final volume V_T .
- If each of these solutions were assumed to obey Beer's law, the absorbance (As)of each solution is described by:

$$A_{s} = \frac{\varepsilon b V_{S} C_{S}}{V_{T}} + \frac{\varepsilon b V_{X} C_{X}}{V_{T}} = k V C_{S} + k V C_{S} = m C_{S} + b$$

where $k = \varepsilon b/V_T$, $m = kV_S$ and $b = kC_XV_X$

d. 67.98%

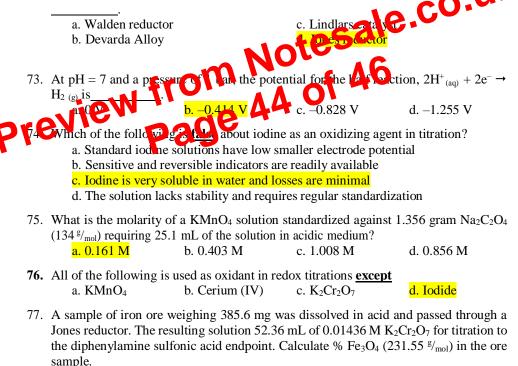
70. The 300 mg sample of impure Na_2SO_4 (142.04) was dissolved in sufficient water and the sulfate was precipitated by the addition of 35.00 mL of 0.1022 M BaCl₂. The precipitate was removed by filtration and the remaining BaCl₂ consumed 6.79 mL of 0.2467 M EDTA for titration to the Calmagite endpoint. Calculate the purity of the sample. a. 80% c. 90% d. 95%

b. 85%

71. A 0.8521 gram sample of an alloy was found to contain Cu (63.55) and Zn (65.41) with small amounts of Pb (207.2) and Hg (200.59). The sample was dissolved in nitric acid and diluted to 500 mL. A 10 mL aliquot was treated with KI to mask the Hg and the resulting solution required 7.06 mL of 0.0348 M EDTA solution. A second 25 mL aliquot was treated with ascorbic acid and the pH was adjusted to 2.00to reduce Hg⁺² and the metallic Hg was removed from the solution. To this solution, thiourea was then added to mask the Cu and the resulting solution required 8.58 mL for titration. The lead ion was titrated in a 250 mL in the presence of NaCN to mask Cu, Zn and Hg and required 3.11 mL for titration. Calculate the percentage of Cuand Hg in the sample of alloy.

a. 47% Cu and 3% Hg b. 44% Cu and 5% Hg c. 53% Cu and 7% Hg d. 56% Cu and 5% Hg

72. Commonly, the analyte in a sample is present in two different oxidation states. Prereduction is then necessary before titration. One of the metallic reductors is zitor soaked in a dilute solution of mercuric chloride. This reductor is known as



a. 15.05% b. 45.15% c. 90.30%

78. A sample of pyrolusite weighing 0.2400 gram was treated with excess KI. The iodine liberated required 46.24 mL of 0.1105 M Na₂S₂O₃ solution. Calculate % MnO₂ (86.94)