# THE UNITED REPUBLIC OF TANZANIA

### NATIONAL EXAMINATIONS COUNCIL

## ADVANCED CERTIFICATE OF SECONDARY EDUCATION EXAMINATION

132/3A CHEMISTRY 3A

(For Both School and Private Candidates)

Time: 3 Hours ANSWERS Year: 2009

### **Instructions**

- 1. This paper consists of THREE questions.
- 2. Answer all questions.



1. You are provided with the following:

MM: A solution made by dissolving 3.25 g of impure potassium dichromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) in 500 cm<sup>3</sup> of distilled water

NN: A solution made by dissolving 12.40 g of sodium thiosulphate pentahydrate (Na<sub>2</sub>S<sub>2</sub>O<sub>3</sub>·5H<sub>2</sub>O) in 500 cm<sup>3</sup> of distilled water

PP: 10% potassium iodide (KI) solution

Also provided: starch indicator and 1M sulphuric acid

Procedure

- (i) Pipette 25.00 cm<sup>3</sup> of MM into a clean titration flask
- (ii) Add to it 25.00 cm<sup>3</sup> of sulphuric acid
- (iii) Add to it 25.00 cm<sup>3</sup> of PP

The liberated iodine reacts with thiosulphate according to the reaction:

$$Cr_2O_7^{2-} + 6I^- + 14H^+ - - > 2Cr^{3+} + 3I_2 + 7H_2O$$

$$I_2 + 2S_2O_3^{2-} ----> S_4O_6^{2-} + 2I^-$$

- (iv) Titrate the iodine against NN using starch as indicator
- (v) Repeat for consistent values

Summary

25.00 cm<sup>3</sup> of MM liberated sufficient iodine required to oxidize 23.60 cm<sup>3</sup> of NN

Questions

- (a) Write balanced redox equations:
- (i)  $Cr_2O_7^{2-} + 6I^- + 14H^+ - > 2Cr^{3+} + 3I_2 + 7H_2O$
- (ii)  $I_2 + 2S_2O_3^{2-} > S_4O_6^{2-} + 2I^-$

# (b) Calculate the molarity of potassium dichromate (K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>) in MM

Volume of NN used =  $23.60 \text{ cm}^3 = 0.02360 \text{ dm}^3$ 

Mass of  $Na_2S_2O_3 \cdot 5H_2O = 12.40 \text{ g}$ 

Molar mass = 248 g/mol

Moles in  $500 \text{ cm}^3 = 12.40 \div 248 = 0.05 \text{ mol}$ 

Molarity =  $0.05 \div 0.5 = 0.1 \text{ mol/dm}^3$ 

Moles of thiosulphate in 23.60 cm<sup>3</sup> =  $0.1 \times 0.02360 = 0.00236$  mol

From the second equation, 1 mol I<sub>2</sub> reacts with 2 mol S<sub>2</sub>O<sub>3</sub><sup>2-</sup>

Moles of  $I_2 = 0.00236 \div 2 = 0.00118$  mol

From the first equation, 1 mol Cr<sub>2</sub>O<sub>7</sub><sup>2-</sup> liberates 3 mol I<sub>2</sub>

Moles of  $Cr_2O_7^{2-} = 0.00118 \div 3 = 0.000393$  mol

Molarity =  $0.000393 \div 0.025 = 0.01572 \text{ mol/dm}^3$ 

(c) Calculate the concentration of K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> in g/dm<sup>3</sup>

Molar mass of  $K_2Cr_2O_7 = 294$  g/mol

Concentration in  $g/dm^3 = 0.01572 \times 294 = 4.621 g/dm^3$ 

(d) Calculate the percentage impurity of the chromate salt

Actual mass used =  $3.25 \text{ g in } 500 \text{ cm}^3 \rightarrow \text{scaled to } 6.50 \text{ g in } 1 \text{ dm}^3$ 

Mass of pure  $K_2Cr_2O_7 = 4.621$  g

Percentage purity =  $(4.621 \div 6.50) \times 100 = 71.09\%$ 

Percentage impurity = 100 - 71.09 = 28.91%

2. You are provided with the following:

MN: A solution of 0.5 M sodium thiosulphate

PQ: A solution of 0.1 M nitric acid

A stopwatch

A thermometer (0–100 °C)

Theory

A white precipitate of amorphous sulphur can be obtained by the reaction:

$$3S_2O_3^{2-} + 2H^+ - - > 3SO_4^{2-} + 3S \downarrow + H_2O$$

This sulphur causes the solution to turn opaque. The rate of the reaction is measured by the time it takes for the solution to obscure a marked letter beneath the beaker.

#### Procedure

- (i) Draw letter X on paper and place a small beaker on it
- (ii) Place 200 cm<sup>3</sup> of water into a 250 cm<sup>3</sup> beaker as water bath
- (iii) Pipette 10.0 cm<sup>3</sup> of MN and 10.0 cm<sup>3</sup> of PQ into separate boiling tubes
- (iv) Warm to required temperature in the bath (e.g., 50 °C)
- (v) Pour both into small beaker on letter X and start stopwatch
- (vi) Record time (in seconds) taken for letter X to disappear
- (vii) Repeat at 60 °C, 70 °C and 80 °C

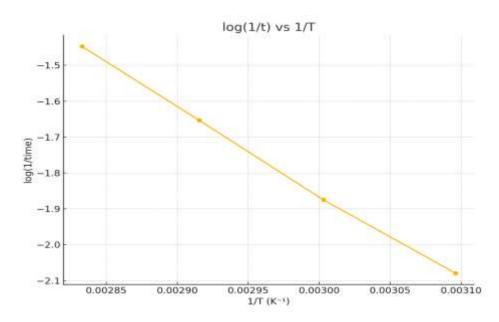
#### Results table (example)

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## (a) Plot a graph of log(1/t) against 1/T ( $K^{-1}$ )



# (b) The slope of the graph log(1/t) vs 1/T is approximately -2410.88.

This slope was obtained from linear regression using the temperature and time data provided.

(c)Using the equation  $K = Ae^{-(-Ea/RT)}$ , and the linear form log(1/t) = log A - Ea/(2.303RT):

Activation energy (Ea) =  $-\text{slope} \times 2.303 \times R$ 

$$= -(-2410.88) \times 2.303 \times 8.314$$

= 46209.90 J/mol

= 46.21 kJ/mol

Pre-exponential factor (A) =  $10^{\text{intercept}}$ 

$$= 2.42 \times 10^5 \; s^{\text{--}1}$$

Therefore,

Activation energy Ea = 46.21 kJ/mol

Frequency factor  $A = 2.42 \times 10^5 \text{ s}^{-1}$ 

3. You are provided with:

V = 0.5 g of succinic acid

W = 0.2 M sodium hydroxide

Distilled water

Diethyl ether

Phenolphthalein indicator (P.O.P)

#### Procedure

- (i) Fill burette with W
- (ii) Measure 50 cm<sup>3</sup> distilled water into separating funnel
- (iii) Add 30 cm<sup>3</sup> ether
- (iv) Add 0.5 g of succinic acid
- (v) Shake vigorously and allow layers to separate
- (vi) Tap off aqueous layer, pipette 25 cm³ into flask

Add P.O.P and titrate with W until pink appears

- (a) Volume of NaOH used =  $20.0 \text{ cm}^3$
- (b) Balanced equation:

$$HOOC-CH_2CH_2-COOH + 2NaOH -----> NaOOC-CH_2CH_2-COONa + 2H_2O$$

(c)(i) Molarity of acid in aqueous layer:

Moles NaOH = 
$$0.2 \times 20 \div 1000 = 0.004$$
 mol

Moles of acid = 
$$0.004 \div 2 = 0.002 \text{ mol}$$

Volume = 
$$25 \text{ cm}^3 = 0.025 \text{ dm}^3$$

Molarity = 
$$0.002 \div 0.025 = 0.08 \text{ mol/dm}^3$$

(ii) Mass of acid in 25 cm<sup>3</sup> =  $0.002 \times 118 = 0.236$  g

So in 50 cm<sup>3</sup>:

Concentration =  $0.236 \times 2 = 0.472$  g in 50 cm<sup>3</sup>

$$g/dm^3 = 0.472 \times 1000 \div 50 = 9.44 \ g/dm^3$$

(iii) Ether layer has 0.5 - 0.236 = 0.264 g

 $Volume = 30 \text{ cm}^3$ 

Concentration =  $0.264 \times 1000 \div 30 = 8.80 \text{ g/dm}^3$ 

(iv) Partition coefficient

 $K = concentration in water \div concentration in ether = 9.44 \div 8.80 = 1.07$