

FACULTY OF ENGINEERING AND THE BUILT ENVIRONMENT

InSTEM

QUALIFICATION: INTRODUCTION TO SCIENCE, TECHNOLOGY, ENGINEERING AND MATHEMATICS								
QUALIFICATIO	N CODE: 04STEM	LEVEL: 4						
COURSE CODE	: ICH402S	COURSE NAME: INTRODUCTION TO CHEMISTRY B						
SESSION:	NOVEMBER 2022	PAPER:	N/A					
DURATION:	3 HOURS	MARKS:	100					

	FIRST OPPORTUNITY EXAMINATION PAPER
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INSTRUCTIONS

- 1. Answer all questions.
- 2. Write all the answers in ink.
- 3. No books, notes, correction fluid (Tippex) or cell phones allowed.
- 4. Pocket calculators are allowed.
- 5. You are not allowed to borrow or lend any equipment or stationary.
- 6. Periodic Table on page 12

Question 1 [15]

Dinitrogen pentoxide, N_2O_5 , is dissolved in an inert solvent (solv) and the rate of Decomposition of N_2O_5 is investigated. This reaction produces nitrogen dioxide, which remains in solution, and oxygen gas.

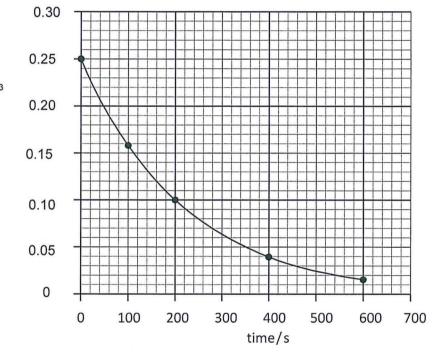
$$N_2O_5(solv) \rightarrow 2NO_2(solv) + \frac{1}{2}O_2(g)$$

- 1.1 Suggest what measurements could be used to follow the rate of this reaction from the given information. (1)
- 1.2 In a separate experiment, the rate of the decomposition of $N_2O_5(g)$ is investigated.

$$N_2O_5(g) \rightarrow 2NO2(g) + \frac{1}{2}O_2(g)$$

The graph below shows the results obtained:

 $[N_2O_5]/moldm^{-3}$



The reaction is first order with respect to N_2O_5 . This can be confirmed from the graph using half-lives.

- 1.2.1 Explain the term half-life of a reaction. (1)
- 1.2.2 Determine the half-life of this reaction. (1)
- 1.2.3 Suggest the effect on the half-life of this reaction if the initial concentration of N_2O_5 Is halved. (1)
- 1.2.4 Use the graph in 1.2 to determine the rate of reaction at 200 s. Show your working and the unit.(2)



The rate equation for this reaction is shown:

rate = $k [N_2O_5]$

1.2.5 Use your answer to 1.2.4 to calculate the value of the rate constant, k, for this reaction and state its units. (1)

1.3 Nitrogen dioxide reacts with ozone, O₃, as shown:

$$2NO_2 + O_3 \rightarrow N_2O_5 + O_2$$

The rate equation for this reaction is rate = $k [NO_2][O_3]$.

Suggest a possible two-step mechanism for this reaction.

(2)

1.4 Gases **A** and **B** react as shown in the following equation:

$$2A(g) + B(g) \rightarrow C(g) + D(g)$$

The initial rate of the reaction was measured in a series of experiments at a constant temperature. The following rate equation was determined:

rate =
$$k[A]^2$$

An incomplete table of data for the reaction between A and B is shown in the Table below.

Experiment	Initial [A] / mol dm ⁻³	Initial [B] / mol dm ⁻³	Initial rate / mol dm ⁻³ s ⁻
1	4.2 × 10 ⁻³	2.8 × 10 ⁻³	3.3 × 10 ⁻⁵
2	7.9 × 10 ⁻³	2.8 × 10 ⁻³	
3		5.6 × 10 ⁻³	1.8×10^{-4}

- 1.4.1 Use the data from Experiment 1 to calculate a value for the rate constant, k, at this temperature. What is the units of k? (3)
- 1.4.2 Use your value of k from Question 1.4.1 to complete the Table for the reaction between A and B.
 (If you have been unable to calculate an answer for Question 1 (a), you may assume a value of 2.3 This is not the correct answer.)
- 1.4.3 The reaction is zero order with respect to B. State the significance of this zero order for the mechanism of the reaction. (1)



Question 2 [15]

When dilute sulfuric acid is electrolysed, water is split into hydrogen and oxygen.

$$2H_2O(I) \rightarrow 2H_2(g) + O_2(g)$$

A current of x A is passed through the solution for 14.0 minutes. 462 cm³ of hydrogen are produced at the cathode, measured under room conditions.

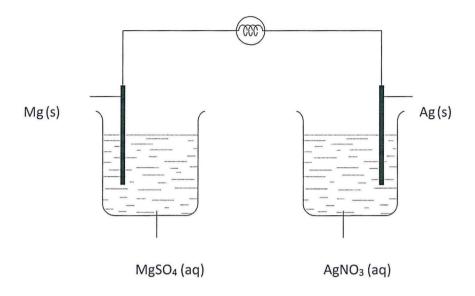
- 2.1 Calculate the number of hydrogen molecules produced during the electrolysis. (2)
- 2.2 Calculate the total number of electrons transferred to produce this number of hydrogen molecules. (1)
- 2.3 Calculate the quantity of charge, in coulombs, of the total number of electrons calculated in 2.2. (1)
- 2.4 Calculate the current, x, passed during this experiment. (1)
- 2.5 The standard entropies, ΔS^{θ} , of three species are given in the table.

species	S ^e /JK ⁻¹ mol ⁻¹
H ₂ O(I)	+70
H ₂ (g)	+131
O ₂ (g)	+205

- 2.5.1 Calculate ΔS° for the reaction $2H_2O(I) \rightarrow 2H_2(g) + O_2(g)$. (1)
- 2.5.2 ΔH° for the reaction $2H_2O(I) \rightarrow 2H_2(g) + O_2(g)$ is +572 kJ mol⁻¹. Calculate ΔG° for this reaction at 298 K. (2)
- 2.5.3 Predict the effect of increasing temperature on the spontaneity of this reaction.Explain your answer. (1)



2.6 The diagram shows an incomplete voltaic cell with a light bulb in the circuit.



- 2.6.1 Identify the missing component of the cell and its function. (2)
- 2.6.2 Deduce the half-equations for the reaction at each electrode when current flows. (2)
- 2.6.3 Redraw the diagram with the location and direction of electron movement when current flows. (1)
- 2.6.4 Calculate the cell potential, in V, using the given data below: (1)

120	Oxidized species	_	Reduced species	E*(V)
	Li*(aq) + e*	*	LJ(s)	-3.04
	K*(aq) + e~	-	K(s)	-2.93
	Ca2*(aq) + Ze*	Allemage	Ca(s)	-2.87
	Na*(aq)+e*	**	Na(s)	-2.71
	Mg2*(aq) + Ze*		Mg(s)	-2.37
	Ai2*(aq)+3e*	*	Al(s)	-1.66
	Mn2*(aq) + 2e"	-	Mn(s)	-1.18
	$H_2O(1) + e^{-}$	-	$\frac{1}{2}H_2(g) + OH^{-}(sq)$	-0.83
	Zn2*(aq) + Ze-	-	Zn(z)	-0.76
	Fe ^{2*} (aq) + 2e ⁻	-	Fe(s)	-0.45
	Nt2*(aq) + 2e*	-	Ni(s)	-0.26
	Sm ²⁺ (aq) + Ze*	-	Sn(s)	-0.14
	Pb2+(aq) + Ze-	-	Pb(s)	-0.13
	H*(aq)+e*	Manage	±H₂(g)	0.00

Oxidized species	_	Reduced species	E*(V)
Os ² *(aq) + e ⁻	-	Cu*(aq)	+0.15
SO ₄ *(aq) + 4H*(aq) + 2e*	-	$H_2SO_3(aq) + H_2O(1)$	+0.17
Cu ^{1*} (aq) + 2e ⁻	-	Cu(x)	+0.34
$\frac{1}{2}O_2(g) + H_2O(1) + 2e^{-}$	-	20H*(aq)	+0,40
Cu*(aq) + e-	-	Cu(s)	+0.52
}/ ₄ (s) + e⁻	-	I*(aq)	+0.54
Fe ⁴⁺ (aq) + e ⁻	-	Fe2+(aq)	+0.77
Ag*(aq) + e*	-	Ag(s)	+0.90
$\frac{1}{4}Br_2(1) + e^-$	-	Br ~(aq)	+1.09
$\frac{1}{3}O_2(g) + 2H_2(aq) + 2e^{-}$	-	H ₂ O(I)	+1.23
Cr ₂ O ₂ ² (aq) + 14H*(aq) + 6e ⁻	-	$2Cr^{3+}(aq) + 7H_2O(l)$	+1.36
2Cl₄(g) + e⁻	*	Cl*(aq)	+1.36
MnU7(aq) +8H*(aq) +5e*		$Mn^{2*} + 4H_2O(0)$	+1.51
$\frac{1}{2}F_{2}(g) + e^{-}$		F=(aq)	+287



- 3.1 Calcium hydroxide is slightly soluble in water.
- 3.1.1. Write an equation to show the dissociation of calcium hydroxide, Ca(OH)₂(s), in aqueous solution. Include state symbols. (1)
- 3.1.2 Calculate the solubility, in mol dm⁻³, of Ca(OH)₂.

$$[K_{sp}: Ca(OH)_2, 5.02 \times 10^{-6} \text{ mol}^3 \text{ dm}^{-9}]$$
 (2)

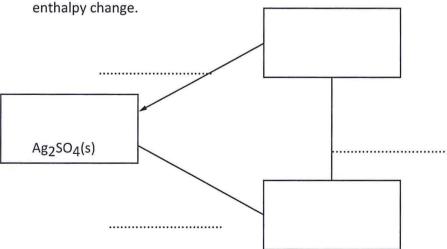
- 3.1.3 Suggest how the solubility of Ca(OH)₂ in aqueous NaOH compares to its solubility in water. Explain your reasoning. (1)
- 3.2 Silver sulfate, Ag_2SO_4 , is sparingly soluble in water. The concentration of its saturated solution is 2.5×10^{-2} mol dm⁻³ at 298 K.
- 3.2.1 Write an expression for the solubility product, K_{sp}, of Ag₂SO₄, and state its units.(1)
- 3.2.2 Calculate the value for K_{sp}(Ag₂SO₄) at 298 K.
- 3.2.3 Using Ag_2SO_4 as an example, redraw and complete the following Hess' Law energy cycle relating the: (5)

(1)

- lattice energy, ΔH_{latt}° ,
- enthalpy change of solution, ΔH_{sol}° , and
- enthalpy change of hydration, ΔH_{hyd}° .

On your diagram:

- include the relevant species in the two empty boxes,
- label each enthalpy change with its appropriate symbol,
 - complete the remaining two arrows showing the correct direction of





Pt | Fe³⁺, Fe²⁺ | Ag⁺ | Ag Ag₂SO₄(s) is formed at the Ag electrode Use the Data in 2.6.4 to calculate the value of under standard conditions, stating which electrode is the positive one. (1)How would the actual Ecell of the above cell compare to the under standard 3.3.2 (1)conditions? Explain your answer. How would the E_{cell} of the above cell change, if at all, if a few cm³ of 3.3.3 concentrated Na₂SO₄(aq) were added to: the beaker containing $Fe^{3+}(aq) + Fe^{2+}(aq)$, the beaker containing Ag₂SO₄(aq)? (2)Explain any changes in Ecell you have stated in 3.3.3. (1)3.3.4 3.3.5 Solutions of iron(III) sulfate are acidic due to the following equilibrium. $[Fe(H_2O)_6]^{3+}(aq) [Fe(H_2O)_5(OH)]^{2+}(aq) + H^+(aq) K_a = 8.9 \times 10^{-4} \text{ mol dm}^{-3}$ Calculate the pH of a 0.1 mol dm⁻³ solution of iron(III) sulfate, Fe₂(SO₄)₃. (2)Question 4 [10] 4.1 Solution Y is hydrochloric acid, HCl (aq). Solution Z is aqueous 4-chlorobutanoic acid, $Cl(CH_2)_3CO_2H(aq)$. The pK_a of $Cl(CH_2)_3CO_2H(aq)$ is 4.52. The pH of both solutions is 4.00. 4.1.1 Write an expression for the K_a of Cl (CH₂)₃CO₂H(aq).0 (1)4.1.2 Write a mathematical expression to describe the relationship between Ka and pKa. (1)Calculate [H⁺] in solutions Y and Z. (1)4.1.3 4.1.4 Calculate the ratio: [HCl] dissolved in solution Y [CI(CH₂)₃CO₂H] dissolved in solution Z (2)

Another electrochemical cell is set up as follows:

3.3



4.2	A buffer solution of pH 5.00 is produced by adding sodium propanoate to 5.00 g of propanoic acid in 100 cm^3 of distilled water. Calculate the mass of sodium propanoate that must be used to produce this buffer solution. The Ka of propanoic acid is 1.35×10^{-5} mol dm ⁻³ .										
	[Mr: propanoic acid, 74.0; sod	ium propa	noate, 96.0]								
4.3	Some dilute sulfuric acid is mixed described in 4.2. The final pH of Explain this observation.			ouffer solution	(2)						
Questi	cion 5				[12]						
	ne, C_3H_6 , reacts with H_2O in the poular formula C_3H_8O .	resence o	f an acid catalyst to	form an alcohol wit	h						
5.1	Name this type of reaction.				(1)						
5.2	Name the catalyst used and st occur.	ate the co	nditions needed for	this reaction to	(2)						
5.3	Redraw and Complete the tab $pi(\pi)$ bonds present in proper			ma (σ) bonds and	(2)						
		σ	π								
	С ₃ Н ₆										
	C ₃ H ₈ O										
5.45.4.15.4.2	The reaction of propene, C ₃ H ₆ . In step 1 C ₃ H ₆ reacts with the or Draw structures to identify the can form in step 1. Explain you Name the major organic produce.	catalyst, H e more sta ur answer.	t, to form a carboca ble and less stable c	tion. arbocations which							
	H₂O.				(1)						
5.5	2-bromopropane reacts to for certain conditions.	m propene	e, hydrogen bromide	e and water under							
5.5.1 5.5.2	Name this type of reaction. Describe the reagents and con	ıditions ne	eded to favour this	reaction.	(1) (2)						



Question 6 [9]

The structure of phenylethanoic acid is shown:

- Phenylethanoic acid can be synthesised using benzene as the starting material. In the first stage of this synthesis, benzene reacts with chloromethane in the presence of an A/Cl_3 catalyst to form methylbenzene. Chloromethane reacts with A/Cl_3 to form two ions. One of these is the carbocation ${}^+CH_3$.
- 6.1.1 Write an equation for the reaction between chloromethane and A/C/₃. (1)
- 6.1.2 Redraw the diagrams below and draw the mechanism of the reaction between benzene and ⁺CH₃. Include all relevant curly arrows, charges and the structure of the intermediate. (3)

6.2 A three-step synthesis of phenylethanoic acid from methylbenzene is shown:

- 6.2.1 State reagents and conditions for step 1. (1)
- 6.2.2 Suggest the structure of compound Q. (1)
- 6.2.3 State reagents and conditions for steps 2 and 3. (2)
- 6.2.4 Draw the structure of an organic by-product that forms in step 1. (1)



Question 7

[10]

- 7.1 The reason for the wide variety of organic compounds is isomerism, either structural isomerism or stereoisomerism.
- 7.1.1 Explain the meaning of the term structural isomerism. (2)
- 7.1.2 Explain the meaning of the term stereoisomerism. (2)
- 7.2 Pent-1-ene, $CH_2=CH(CH_2)_2CH_3$, does not show stereoisomerism.
- 7.2.1 Give **two** reasons why pent-1-ene does not show stereoisomerism. (2)
- 7.2.2 A structural isomer of pent-1-ene is used as the monomer to form a polymer. The repeat unit of this polymer is shown:

Draw the displayed formula of the monomer used to make this polymer. Give the name of the monomer (2)

7.2.3 A different structural isomer of pent-1-ene shows geometrical isomerism. Draw the structure of one of the two geometrical isomers with the formula C_5H_{10} . Give the full name of this isomer. (2)



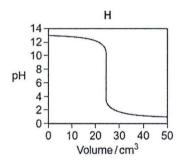
Question 8

[6]

Titration curves, labelled E, F, G and H, for combinations of different aqueous solutions of acids and bases are shown in the diagrams below. All solutions have concentrations of 0.1 mol dm^{-3} .

PH 6-4-2-0 10 20 30 40 50 Volume/cm³ PH 6-4-2-0 10 20 30 40 50 Volume/cm³

PH 6-4-2-0 10 20 30 40 50 Volume/cm³



8.1 From the curves E, F, G and H, choose the curve produced by the addition of:

8.1.1 sodium hydroxide to 25 cm³ of ethanoic acid

(1)

8.1.2 ammonia to 25 cm³ hydrobromic acid (HBr)

(1)

8.1.3 hydrochloric acid to 25 cm³ of potassium hydroxide

(1)



8.2 The table below shows information about some acid-base indicators.

Indicator	pH range	Lower pH colour	Higher pH colour
pentamethoxy red	1.2-3.2	violet	colourless
naphthyl red	3.7–5.0	red	yellow
4-nitrophenol	5.6-7.0	colourless	yellow
cresol purple	7.6-9.2	yellow	purple

- 8.2.1 Which indicator in the Table could be used for the titration that produces curve E but not for the titration that produces curve F? (1)
 8.2.2 Give the colour change at the end point of the titration that produces curve H when naphthyl red is used as the indicator. (1)
- 8.2.3 A beaker contains 25 cm³ of a buffer solution at pH = 6.0

 Two drops of each of the four indicators in Table 2 are added to this solution.

 State the colour of the mixture of indicators in this buffer solution.

You should assume that the indicators do not react with each other.

Question 9 [4]

9.1 Write the equations for the following decay processes:

Positron emission from silicon-26 (2)

(1)

9.2 The radioisotope fluorine-21 had an initial mass of 80 milligrams. 20 milligrams of sample remained unchanged after 8.32 s. What is the half life of F-21? (2)

THE END



The Periodic Table of the Elements

1	2											3	4	5	6	7	0
							1.0 H										(18) 4.0 He
(1)	(2)			Key			hydrogen 1					(13)	(14)	(15)	(16)	(17)	helium 2
6.9 Li lithium 3	9.0 Be beryllium 4			ive atomic symbol name c (proton)								10.8 B boron 5	12.0 C carbon 6	14.0 N nitrogen 7	16.0 O oxygen 8	19.0 F fluorine 9	20.2 Ne neon 10
23.0 Na sodium 11	24.3 Mg magnesium 12	(3)	(4)	<i>(</i> 5)	(6)	(7)	(8)	<i>(</i> 9)	(10)	(11)	(12)	27.0 Al aluminium 13	28.1 SI silicon 14	31.0 P phosphorus 15	32.1 S sulfur 16	35.5 Cl chlorine 17	39.9 Ar argon 18
39.1 K potassium 19	40.1 Ca calcium 20	45.0 Sc scandium 21	47.9 Ti titanium 22	50.9 V vanadium 23	52.0 Cr chromium 24	54.9 Mn manganese 25	55.8 Fe iron 26	58.9 Co cobalt 27	58.7 Ni nickel 28	63.5 Cu copper 29	85.4 Zn zinc 30	69.7 Ga gallium 31	72.6 G9 germanium 32	74.9 As arsenic 33	79.0 Se selenium 34	79.9 Br bromine 35	83.8 Kr krypton 36
85.5 Rb rubidium 37	87.6 Sr strontium 38	88.9 Y yttrium 39	91.2 Zr zirconium 40	92.9 Nb niobium 41	96.0 Mo molybdenum 42	[97] Tc technetium 43	101.1 Ru ruthenium 44	102.9 Rh rhodium 45	106.4 Pd palladium 46	107.9 Ag silver 47	112.4 Cd cadmium 48	114.8 In Indium 49	118.7 Sn tin 50	121.8 Sb antimony 51	127.6 Te tellurium 52	126.9 lodine 53	131.3 Xe xenon 54
132.9 Cs caeslum 55	137.3 Ba barlum 56	138.9 La * lanthanum 57	178.5 Hf hamlum 72	180.9 Ta tantalum 73	183.8 W tungsten 74	186.2 Re rhenlum 75	190.2 Os osmlum 76	192.2 Ir Irldlum 77	195.1 Pt platinum 78	197.0 Au gold 79	200.6 Hg mercury 80	204.4 Tl thalllum 81	207.2 Pb lead 82	209.0 Bi blamuth 83	[209] Po polonium 84	[210] At astatine 85	[222] Rn radon 86
[223] Fr francium 87	[226] Ra radium 88	[227] Ac † actinium 89	[267] Rf rutherfordium 104	[270] Db dubnium 105	[269] Sg seaborgium 106	[270] Bh bohrium 107	[270] Hs hassium 108	[278] Mt meitnerium 109	[281] Ds darmstadtium 110	[281] Rg roentgenium 111	[285] Cn copernicium 112	[286] Nh nihonium 113	[289] Fl flerovium 114	[289] Mc mascovium 115	[293] Lv livermorium 116	[294] Ts tennessine 117	[294] Og ogenesson 118
* 58 – 7	* 58 - 71 Lanthanides		140.1 Ce cerium 58	59	144.2 Nd neodymium 60	61	150.4 Sm samarium 62	152.0 Eu europium 63	157.3 Gd gadolinium 64	158.9 Tb terbium 65	162.5 Dy dysprosium 66	164.9 Ho holmium 67	167.3 Er erbium 68	168.9 Tm thulium 69	173.0 Yb ytterbium 70	175.0 Lu lutetium 71	
† 90 – 1	03 Actin	des		232.0 Th thorium 90	231.0 Pa protactinium 91	238.0 U uranium 92	[237] Np neptunium 93	[244] Pu plutonium 94	[243] Arn americium 95	[247] Cm curium 96	[247] Bk berkelium 97	[251] Cf californium 98	[252] E8 einsteinium 99	[257] Fm fermium 100	[258] Md mendelevium 101	[259] No nobelium 102	[262] Lr lawrencium 103

