UNIVERSITY OF SWAZILAND

FINAL EXAMINATION 2007

TITLE OF PAPER:

INTRODUCTORY

INORGANIC

CHEMISTRY

COURSE NUMBER:

C201

TIME ALLOWED:

THREE (3) HOURS

INSTRUCTIONS:

THERE ARE SIX (6) QUESTIONS. ANSWER ANY FOUR (4) QUESTIONS. EACH QUESTION IS WORTH 25

MARKS.

A TABLE OF CONSTANTS AND A PERIODIC TABLE ARE **ATTACHED**

NON-PROGRAMMABLE ELECTRONIC CALCULATORS MAY BE **USED**

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QUESTION ONE

(a)	The entrance doors at airports are often controlled by photoelectric cells. When the maximum wavelength of light that could be used for such systems cesium cathodes if electrons are ejected from cesium with a kinetic energy of 9.6×10^{-20} J upon irradiation with 500 nm light? [6]	with
(b)	 The He⁺ ion is a one electron-system similar to hydrogen, except that Z Consider the third and fifth energy levels in the species He⁺. (i) Calculate the frequency of the radiation which this species would all in going from the lower of these levels to the upper level. (ii) Assuming circular orbits of the simple Bohr theory, calculate the rate the radii of the orbits associated with these levels. (iii) Calculate the first ionisation potential of He⁺. [9] 	bsorb tio of
(c)	Write out the ground-state electronic configurations and predict the numb unpaired electrons of each of the following species: (i) Cu ²⁺ (ii) Mn ³⁻ (iii) V [3]	
(d)	Calculate the de Broglie wavelength of an electron moving at one-tenth the of light.	
(e)	What is an orbital? Draw the shapes of the orbitals with $n = 3$ [5]]
QUE	STION TWO	
(a)	Relate the tendency of atoms to gain or lose electrons to the types of bonds form.	_
(b)	If Ge is added to GaAs, the Ge is about equally distributed between the Ga an sites. (i) Which sites would the Ge prefer if Se is added also? (ii) Would GaAs doped with Se be an n-type or p-type semiconductor? [4]	
(c)	The promotion of an electron from the valence band into the conduction band pure TiO ₂ by light absorption requires a wavelength of less than 350 nm. Calculate the energy gap in eV between the valence and conduction bands. [2]	
(d)	Distinguish intrinsic from extrinsic semiconduction and give an example of each in actual compounds.	
(e)	 (i) Complete the following reaction: ²³²₉₀Th +? → ¹₀n + ²³⁵₉₂U (ii) Write equations showing how (1) ²³⁸₉₂U undergoes α decay. (2) ¹³₇N undergoes β⁺ emission. 	

QUESTION THREE

(a)	Determine the hybridization for the central atom in each of the following. (i) BeF ₂ (ii) TeCl ₄ (iii) OPCl ₃							
	in eac	h case predict	the geometry usin	g the VSEPR theory.	[9]			
(b)	energy (i)	(which is a la Give the m showing the	rge assumption), olecular-orbital endistribution of ele	nergy level diagram for the ctrons in the molecular orbi	he molecule BN,			
	(ii)	(1) BN	bond orders in the bond orders in the (2) BN		[7]			
(c)		n π bonding a	`,	olecular orbitals that result				
(d)	In wh	ich of the fo	llossing govelent	compounds is the central	atom obeying the			
(d)	In which of the following covalent compounds is the central atom obeying octet rule?							
	(i) CH	I 4	(ii) H ₂ S	(iii) ClF ₃	[3]			
_		N FOUR			4 2200 1-I aT-1			
(a)			rgies for Cl, Cl in these difference	and Cl ⁺ are 343, 1250 a.s.	nd 2300 kJmol ⁻ , [6]			
(b)	(i) (ii)	sh of the follow S ²⁻ , Br ⁻ Fe ²⁺ , Co ³⁺ O ²⁻ , Se ²⁻	wing pairs, which i	is the larger ion? Explain.	[6]			
(c)			s, determine the e	ffective nuclear charge (Z*)) for the following			
* 2	electr (i) (ii) (iii)	The outerm A 3d electro	ost electron in an on in a nickel atom on in a nickel atom	ı. ·	[7]			
(d)	Provi (i) (ii)	Water readi	ns for the following decomposes Boounds have high have low melting	Cl ₃ whereas it has no effect of the melting points while n	on CCl ₄ . nolecular covalen [6]			

QUESTION FIVE

(ii)

(iii)

Give the uses of CO.

How can CO be detected?

(a) Give equations to show the reaction of hydrogen with: N₂ using a catalyst of activated Fe at 380-450 °C and 200 atmospheres pressure. (ii) CO over a Cu/Zn catalyst at 300 °C. [4] (b) Account for the following: The variation in boiling points of the Group VIA hydrides whose values are 100, -60, -42 and -2.3 °C for H₂O, H₂S, H₂Se and H₂Te respectively. (ii) The trend in the bond angles for the hydrides of the Group VA elements which are as follows: NH₃, 106.6°; PH₃, 93.6°; AsH₃, 91.8°; SbH₃, 91.3°. [10] (c) (i) What products are formed when Li, Na, and K are each burnt in oxygen? (ii) How do these products react with water? (d) On the basis of inductive effect, the Lewis acidity of the boron halides is expected to be BF₃>BCl₃>BBr₃. Experimentally, the opposite is observed. Explain this anomaly. [5] **QUESTION SIX** (a) A total of 0.4008 g of an alkaline earth metal element (A) reacts with oxygen to give 0.5608 g of an alkaline oxide (B). The metallic element (A) reacts with nitrogen gas to form the compound (C) and with hydrogen gas to form the compound (D). On reacting the compound (D) with water, a gas (E) was given off and a sparingly soluble compound (F) was formed. Compound (F) is also obtained when oxide (B) is dissolved in water. Compound (F) gave a strongly basic aqueous solution, which can be neutralised with hydrochloric acid to form the salt (G). When a sample of the salt (G) was moistened with nitric acid and introduced into a Bunsen burner flame, a red colour was imparted into the Bunsen flame. Bubbling carbon dioxide gas through a solution of compound (F) gave a milky solution of compound (H), which finally became a clear solution of compound (I) as more carbon dioxide passed through the solution. Calculate the molar mass of the element (A) and hence identify the (i) element (A). [3] (ii) Identify, with reasons, the compounds (B) to (I). [4] Write balanced equations for all the reactions involved. [4] (iii) Calculate the volume of 0.750 M HCl required to neutralise the strongly (iv) basic solution obtained from 0.5608 g of the metal oxide (B). [4] Compare and contrast the structures of trimethylamine and trisilylamine. [4] (b) (c) (i) How would you prepare CO in the laboratory?

[6]

General data and fundamental constants

Quantity	Symbol	Value
Speed of light	c	2.997 924 58 X 10 ⁸ m s ⁻¹
Elementary charge	e .	1.602 177 X 10 ⁻¹⁹ C
Faraday constant	$F = N_A e$	9.6485 X 10 ⁴ C mol ⁻¹
Boltzmann constant	k	1.380 66 X 10 ²³ J K ⁻¹
Gas constant	$R = N_A k$	8.314 51 J K ⁻¹ mol ⁻¹
		8.205 78 X 10 ⁻² dm ³ atm K ⁻¹ mol ⁻¹
		6.2364 X 10 L Torr K ⁻¹ mol ⁻¹
Planck constant	h	6.626 08 X 10 ⁻³⁴ J s
	$\hbar = h/2\pi$	1.054 57 X 10 ⁻³⁴ J s
Avogadro constant	N_A	6.022 14 X 10 ²³ mol ⁻¹
Atomic mass unit	u	1.660 54 X 10 ⁻²⁷ Kg
Mass		•
electron	m_e	9.109 39 X 10 ⁻³¹ Kg
proton	m_p	1.672 62 X 10 ⁻²⁷ Kg
neutron	m_n	1.674 93 X 10 ⁻²⁷ Kg
Vacuum permittivity	$\varepsilon_o = 1/c^2 \mu_o$	$8.854\ 19\ X\ 10^{-12}\ J^{-1}\ C^2\ m^{-1}$
	$4\pi\varepsilon_{o}$	$1.112 65 \times 10^{-10} \text{ J}^{-1} \text{ C}^2 \text{ m}^{-1}$
Vacuum permeability	μ_o	$4\pi \times 10^{-7} \text{ J s}^2 \text{ C}^{-2} \text{ m}^{-1}$
		$4\pi \times 10^{-7} \text{ T}^2 \text{ J}^{-1} \text{ C}^{-2} \text{ m}^3$
Magneton		
Bohr	$\mu_B = e\hbar/2m_e$	9.274 02 X 10 ⁻²⁴ J T ⁻¹
nuclear	$\mu_N = e \hbar/2m_p$	5.050 79 X 10 ⁻²⁷ J T ⁻¹
g value	g _e	2.002 32
Bohr radius	$a_o = 4\pi \varepsilon_o \hbar/m_e e^2$	5.291 77 X 10 ⁻¹¹ m
Fine-structure constant	$\alpha = \mu_0 e^2 c/2h$	7.297 35 X 10 ⁻³
Rydberg constant	$R_{\infty} = m_e e^4 / 8h^3 c \varepsilon_0^2$	1.097 37 X 10 ⁷ m ⁻¹
Standard acceleration		
of free fall	g	9.806 65 m s ⁻²
Gravitational constant	G	6.672 59 X 10 ⁻¹¹ N m ² Kg ⁻²
Conversion factors		

Conversion factors

1 cal 1 eV			4.184 1.602	joules (2 X 10	19 J			le	96 48	1 X 10 ⁻⁷ J 96 485 kJ mol ⁻¹ 23.061 kcal mol ⁻¹	
f femto 10 ⁻¹⁵	p pico 10 ⁻¹²	n nano 10 ⁻⁹	μ micro 10 ⁻⁶	m milli 10 ⁻³		deci	k kilo 10 ³	M mega 10 ⁶	G giga 109	Prefixes	

 $\begin{array}{l} \textbf{Spectrochemical Series} \\ \Gamma < Br^{-} < S^{2-} < Cl^{-} < NO_{3}^{-} < F^{-} < OH^{-} < EtOH < C_{2}O_{4}^{2-} < H_{2}O < EDTA < (NH_{3}, \ py) < \\ en < dipy < NO_{2}^{-} < CN^{-} < CO \end{array}$

PERIODIC TABLE OF ELEMENTS

*	Ť.	7	6	Us.	4	ω	2	-	PERIODS
**Actinide Series	ınthanic	Fr. 87	132.91 Cs 55	85,468 Rb 37	39.098 K 19	22.990 Na 11	6:941 Li 3	1.00 8 1	IA I
e Series	*Lanthanide Series	R.a 88	137.33 Ba 56	87.62 Sr 38	40.078 Ca 20	24.305 Mg 12	9.012 Be 4		2 IIA
	, 1	*** Ac	138.91 *La 57	88.906 Y 39	44.956 Sc 21			•	3
232.04 Th 90	140.12 Ce 58	(261) Rf 104	178.49 Hf 72	91.224 Zr 40	47.88 Ti 22				IVB
231.04 Pa 91	140.91 Pr 59	(262) Ha 105	180.95 Ta 73	92.906 Nb 41	50.942 V 23				VB
238.03 U 92	144.24 Nd 60	(263) Unh 106	183.85 W 74	95.94 Mo 42	51.996 Cr 24	TRAN			VIB V
237.05 Np 93	(145) Pm 61	(262) Uns 107	186.21 Re 75	98.907 Tc 43	54.938 Mn 25	TRANSITION ELEMENTS			7 VIIB
(244) Pu 94	150.36 Sm 62	(265) Uno 108	190.2 Os 76	101.07 Ru 44	55.847 Fe 26	ELEM			8 G
(243) Am 95	151.96 Eu 63	(266) Une 109	192.22 Ir 77	102.91 Rth 45	58.933 Co 27	ENTS			GROUPS 9 VIIIB
(247) Cm 96	157.25 Gd 64	(267) Uun 110	195.08 Pt 78	106.42 Pd 46	58.69 Ni 28				10
(247) Bk 97	158.93 Tb 65		196.97 Au 79	107.87 Ag 47	63.546 Cu 29		Atomic mas Symbol Atomic No		B 11
(251) CY 98	162.50 Dy 66		200.59 Hg 80	112.41 Cd 48	65.39 Zn 30		Atomic mass Symbol Atomic No.		12 IIB
(252) Es 99	164.93 Ho 67		204.38 T1 81	114.82 In 49	69.723 Ga 31	26.982 AI 13	¥0.811 →B 5		13 IIIA
(257) Fm 100	167.26 Er 68		207.2 Pb 82	118.71 Sn 50	72.61 Ge 32	28.086 Si 14	12.011 C 6		14 IVA
(258) Md 101	168.93 Tm 69		208.98 Bi 83	121.75 Sb 51	74.922 As 33	30.974 P 15	14.007 N 7		15 VA
(259) No 102	173.04 Yb 70		(209) Po 84	127.60 Te 52	78.96 Se 34	32.06 S 16	15.999 O 8		16 VIA
(260) Lr 103	174.97 Lu 71		(210) At 85	126.90 I 53	79.904 Br 35	35.453 CI 17	18.998 F 9		17 VIIA
			(222) Rn 86	131.29 Xe 54	83.80 Kr 36	39.948 Ar 18	20.180 Ne 10	4.003 He 2	18

() indicates the mass number of the isotope with the longest half-life.