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PSF 1303 HOME ASSIGNMENT I 2009/2010

1. (a) Write the full electron configurations ($1S^2$, $2S^2$.....) of the following ions.
K^+, Ca^{2+}, Cl^-, S^{2-}
With which noble gas are these ions isoelectronic.

(b) Write balanced chemical equations for the following reactions;
(i) reaction of aqueous $H_2C_2O_4$ with NaOH
(ii) oxidation of Fe^{2+} by $K_2Cr_2O_7$ in acidic medium
(iii) dissociation of N_2O_4

(c) The density of germanium, Ge is 5.23 g cm^{-3}. Express in SI units,
(i) the volume
(ii) the mass
of a cube of germanium with an edge length of 80.5 mm

(d) The percentage of H_3PO_4 in syrupy phosphoric acid is 98% by w/v. Density of syrupy phosphoric acid is 1.89 g cm^{-3}. Calculate the volume of syrupy phosphoric acid needed to prepare 1.5 dm^3 of 3.8 M (mol dm^{-3}) aqueous H_3PO_4 solution.

(e) A compound of nitrogen (N) and oxygen(O) has the weight composition of 1.52 g N and 3.47 g O. The molar mass of this compound is known to be between 90 g and 95 g. Determine the molecular formula of the compound.(N = 14, O = 16)

2. (a) (i) Define the term 'first ionisation energy' of an element.
(ii) Write an equation to illustrate the second ionization energy of lithium.

(b) Explain why the first electron affinity of oxygen atoms,
$O(g) + e^- \rightarrow O^-(g)$ is exothermic,
whereas the second stage,
$O^-(g) + e^- \rightarrow O^{2-}(g)$ is endothermic.

© Draw dot and cross diagrams of the following molecules.
(i) CCl_4 (ii) CO_3^{2-} (iii) IF_5 (iv) H_3O^+

(d) (i) What is meant by the term "dipole moment"?
(ii) Explain why CHF_3 possess a dipole moment but CF_4 molecule does not?

(f) Explain the following:
(i) The radius of a cation is less than that of the relevant atom whereas radius of an anion is higher than that of the relevant atom.
(ii) Lithium and Magnesium show some similar chemical properties, although the two elements belong to different groups.
(iii) Lithium iodide and caesium iodide are both ionic compounds. Which has the greater covalent character.
(iv) Ammonia has the highest boiling point among the hydrides of group V elements.
(v) Copper and sodium are not classified in the same block of the periodic table.

ශ්‍රී ලංකා විවෘත විශ්වවිද්‍යාලය

PSF 1303 2009/2010

පැවරුම I

(1) a) පහත දැක්වෙන අයනයන්හි ඉලෙක්ට්‍රෝනික වින්‍යාසයන් s හා p ඉලෙක්ට්‍රෝන ($1s^2 \ldots$) අනුසාරයෙන් ලියා දක්වන්න.

K^+ , Ca^{2+} , Cl^- , S^{2-}

මෙම අයන සමඉලෙක්ට්‍රෝනික (isoelectronic) වන්නේ කුමන නිශ්ක්‍රිය වායුන් සමඟ ද?

b) පහත ප්‍රතික්‍රියා සඳහා තුලිත රසායනික සමීකරණ ලියන්න.

i) ඇලිය $H_2C_2O_4$ සහ $NaOH$ අතර ප්‍රතික්‍රියාව.

ii) $K_2Cr_2O_7$ මඟින් ආම්ලික මාධ්‍යයේදී Fe^{2+} ඔක්සිකරණය

iii) N_2O_4 හි වියෝජනය.

c) ජර්මේනියම් (Ge) වල ඝනත්වය 5.23 g cm^{-3}, දිග 80.5 mm වූ ජර්මේනියම් සෑන්ඩයක,

i) පරිමාව

ii) ස්කන්ධය

SI ඒකක වලින් දක්වන්න.

d) ඉතා ඵොස්ෆරික් අම්ලයේ H_3PO_4 ප්‍රතිශතය 98% w/w. මෙම ඵොස්ෆොරික් අම්ලයේ ඝනත්වය 1.89 g cm^{-3} වේ. 3.8 M (mol dm^{-3}) සාන්ද්‍රණයෙන් යුත් ඇලිය ඵොස්ෆරික් අම්ල ද්‍රාවණ 1.5 dm^3 සෑදීම සඳහා අවශ්‍යවන මෙම ඵොස්ෆරික් අම්ල පරිමාව ගණනය කරන්න.

e). N හා O වලින් සමන්විත සංයෝගයක ඛර අනුව N 1.52 g ද, O 3.47 g අඩංගු වේ. මෙම සංයෝගයේ මවුලික ස්කන්ධය 90 g හා 95 g අතර වේ. මෙම සංයෝගයේ අනුක සූත්‍රය නිර්ණය කරන්න. (N = 14 , O = 16)

(2) a) i) යුලිදුවයක "පළමු අයනීකරණ ශක්තිය" අර්ථ දක්වන්න.

ii) ලිතියම් (Li) හි දෙවන අයනීකරණ ශක්තිය විදහා දැක්වීමට සමීකරණයක් ලියන්න.

b) ඔක්සිජන් පරමාණුවේ පළමුවන ඉලෙක්ට්‍රෝන බන්ධුතාවය

$O_{(g)} + e \longrightarrow O^-_{(g)}$ ආකාරයෙනි.

එදෙවන අවස්ථාවේ දී,

$O^-_{(g)} + e \longrightarrow O^{2-}_{(g)}$ ආර අවශෝෂකද වන්නේ මන්දැයි පැහැදිලි කරන්න.

c) පහත අණු සඳහා කිනම් කනිෂ්ඨ සවහන් අදින්න.

i) CCl_4 ii) CO_3^{2-} iii) IF_5 iv) H_3O^+

d). i) "ද්විධ්‍රැව ඝූර්ණය" (dipole moment) යන්නෙන් අදහස් වන්නේ කුමක්ද?

ii) CF_4 සඳහා ද්විධ්‍රැව ඝූර්ණයක් නැතත් CHF_3 සඳහා ද්විධ්‍රැව ඝූර්ණයක් පවතින්නේ මන්දැයි පැහැදිලි කරන්න.

f) පහත දී පැහැදිලි කරන්න.

i) නැටරායනික අරය, අදාළ පරමාණුක අරයට වඩා වැඩි වෙතත්, ඇනායනික අරය අදාළ පරමාණුක අරයට වඩා වැඩිවේ.

ii) ලිතියම් සහ මැග්නීසියම් කාණ්ඩ 2 ට අයත් වුවද සමාන රසායනික ගුණ පෙන්වයි.

iii) ලිතියම් ඔක්සයිඩ් සහ සීසියම් ඔක්සයිඩ් යන දෙකම අයනික සංයෝග වන නමුත් වැඩිම සාපේක්ෂ අණුක ගුණ

දික්වාන්නේ කුමන සංයෝජනය ද ?

iv) V වන කාණ්ඩයේ මූලද්‍රව්‍යයන්ගේ හයිඩ්‍රයිඩ අඩුවන්
වැනිම කාර්යාංකය ඇමෝනියා ..

v) ෆොස්පරස් සහ සෝඩියම් ආවර්තිතා වගුවේ එකම
ආවර්තයේ වර්ගීකරණය කර දක්වන්න.

PSF 1303 மதிப்பீடு 01 2009/2010

1. (a) பின்வரும் அயன்களுக்குரிய இலத்திரன் நிலையமைப்பை $1s^2, 2s^2 \ldots$ எனும் வடிவில் எழுதுக? இவ் அயன்கள் எச்சடத்துவ வாயுவுடன் சம இலத்திரன் நிலையமைப்பைக் கொண்டுள்ளன?

$$K^+ , Ca^{2+} , Cl^- , S^{2-}$$

(b) பின்வரும் தாக்கங்களுக்குரிய சமப்படுத்தப்பட்ட இரசாயன சமன்பாடுகளை எழுதுக?

(i) $H_2C_2O_4$ நீர்க்கரைசலுடன் NaOH இன் தாக்கம்

(ii) அமில ஊடகத்தில் $K_2Cr_2O_7$ இனால் Fe^{2+} இன் ஒட்சியேற்றம்

(iii) N_2O_4 இன் பிரிகைத் தாக்கம்

(c) ஜேர்மானியத்தின், Ge அடர்த்தி 5.23 gcm^{-3} ஆகும். SI அலகில் 80.5 mm நீளமுடைய கனவடிவ ஜேர்மானியத்தின் ,

(i) கனவளவு

(ii) திணிவு

என்பனவற்றைக் கணிக்குக?

(d) பாகு நிலை பொஸ்போரிக்கமிலத்தில் H_3PO_4 இன் திணிவு/கனவளவு வீதம் 98% ஆகும். பாகு நிலை பொஸ்போரிக்கமிலத்தின் அடர்த்தி 1.89 gcm^{-3}. 3.8M ($Mol\ dm^{-3}$) செறிவுடைய H_3PO_4 நீர்க் கரைசலின் 1.5 dm^3 கரைசலைத் தயாரிக்கத் தேவையான பொஸ்போரிக்கமிலத்தின் கனவளவைக் கணிக்குக?

(e) ஒரு சேர்வை 1.52 g நைதரசனையும் 3.47 g ஒட்சிசனையும் திணிவு அமைப்பில் கொண்டுள்ளது. இந்த சேர்வையின் மூலரத் திணிவு 90-95 g இற்கு இடைப்பட்டதாகும். இச்சேர்வையின் மூலக்கூற்றுச் சூத்திரத்தைப் பெறுக? (N=14 , O=16)

2.(a) (i) மூலகம் ஒன்றின் முதலாம் அயனாக்கற் சக்தி எனும் பதத்தை வரையறுக்குக?
(ii) வீதியுத்தின் இரண்டாம் அயனாக்கற் சக்தி எனும் பதத்தை வீளரிப்பதற்குப் பொருத்தமான சமன்பாடு ஒன்றை எழுதுக?

(b) ஏன் ஒட்சிசன் அணுவின் முதலாம் இலத்திரன் நாட்டற் சக்தி புற வெப்பமாகவும் இரண்டாம் இலத்திரன் நாட்டற் சக்தி அக வெப்பமாகவும் இருக்கின்றது என்பதை வீளக்குக?

$O(g) + e \longrightarrow O^-(g)$ புறவெப்பம்

$O^-(g) + e \longrightarrow O^{2-}(g)$ அகவெப்பம்

PSF 1303
Home Assignment 1 – 2009/2010
Answer guide

1) a)

K^+ - $s^2\ 2s^2\ 2p^6 3s^2\ 3p^6$
Ca^{2+} - $s^2\ 2s^2\ 2p^6 3s^2\ 3p^6$
Cl^- - $1s^2\ 2s^2\ 2p^6 3s^2\ 3p^6$
S^{2-} - $s^2\ 2s^2\ 2p^6 3s^2\ 3p^6$

Ar is isoelectronic with all these ions

b) i) $H_2C_2O_4 + 2NaOH \longrightarrow Na_2C_2O_4 + 2H_2O$

ii) $Fe^{2+} \longrightarrow Fe^{3+} + e$
$Cr_2O_7^{2-} + 14\ H^+ + 6e \longrightarrow 2Cr^{3+} + 7\ H_2O$

$6Fe^{2+} + Cr_2O_7^{2-} + 14\ H^+ \longrightarrow 6\ Fe^{3+} + 2Cr^{3+} + 7H_2O$

iii) $N_2O_4(g) \rightleftharpoons 2NO_2$

c) Density $= 5.23 \times 10^3 kgm^{-3}$

i) Volume of a cube of Ge $= (80.5 \times 10^{-3}\ m)^3$
$= 5.216 \times 10^{-4}\ m^3$

ii) The mass of a cube of Ge $= 5.23 \times 10^3\ kgm^{-3} \times 5.216 \times 10^{-4}\ m^3$

$= 27.30 \times 10^{-1}\ kg$

d) Density $= 1.89\ g\ cm^{-3}$

No: of mol in $1cm^3$ $= \frac{1.89\ g}{98\ gmol^{-1}} \times \frac{98}{100}$

$= 1.89 \times 10^{-2}\ mol$

No: of mol in $1000\ cm^3 = 1.89 \times 10\ mol$

Concentration of syrupy H_3PO_4 acid $= 18.9\ moldm^{-3}$

No: of mols in $1.5\ dm^{-3}$ of $3.8\ moldm^{-3}$ H_3PO_4 acid $= 3.8\ moldm^{-3} \times 1.5dm^{-3}$
$= 5.7\ mol$

Volume required by the syrupy H_3PO_4 acid to prepare the above H_3PO_4 (aq) solution

$$= \frac{5.7 \text{ mol}}{18.9 \text{ moldm}^{-3}}$$

$$= 0.302 \text{ dm}^3$$

$$= \underline{\underline{302 \text{ cm}^3}}$$

e)

	N	O
Weight	1.52 g	3.47 g
No: of mols	$\frac{1.52 \text{ g}}{14 \text{ gmol}^{-1}}$	$\frac{3.47 \text{ g}}{16 \text{ gmol}^{-1}}$
	0.108 mol	0.216 mol
Molar ratio	1	2

$(NO_2)x = 90$ (as)

$(14 + (16 * 2))x = 90$

$46x = 90$

$x = 2$

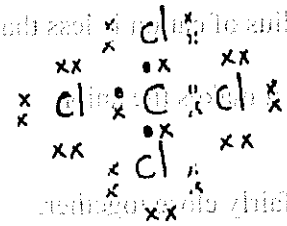
Molecular fomula $\underline{\underline{N_2O_4}}$

2) a) i) The energy required to remove an electron from an atom in the gaseous state to an infinitely large distance.

ii) $Li^{+} \longrightarrow Li^{2+} + e$

b) Electron affinity is the energy required to gain an electron to an atom in the gaseous state. The second electron affinity will take a large positive value because the second electron must be forced on against the net negative charge of the ion. Therefore that Energy should be supplied from the out side to force an electron to combine with the gaseous atom to form an ion. i.e The process is endothermic.

c) i) CCl_4



C — 4

4(Cl) — 28

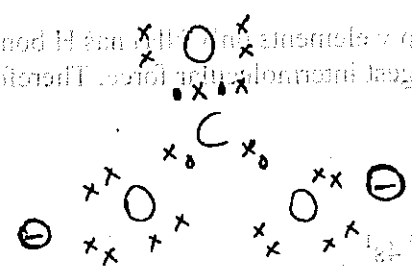
Total en: — 32

For bonding — 8

24 ÷ 2

12 pairs

ii) CO_3^{2-}



C — 4

3(O) — 18

Total — 22

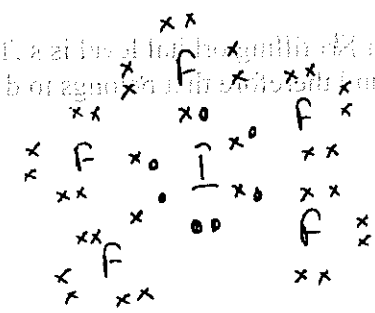
charge — +2

24 — 6

18 ÷ 2

9 pairs.

iii). IF_5



I — 7

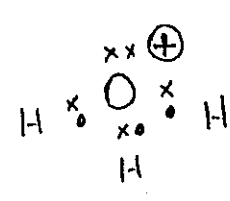
5(F) — 35

42 — 10

32 ÷ 2

16 pairs

iv). H_3O^{+}



3(H) — 3

O — 6

9 — 1

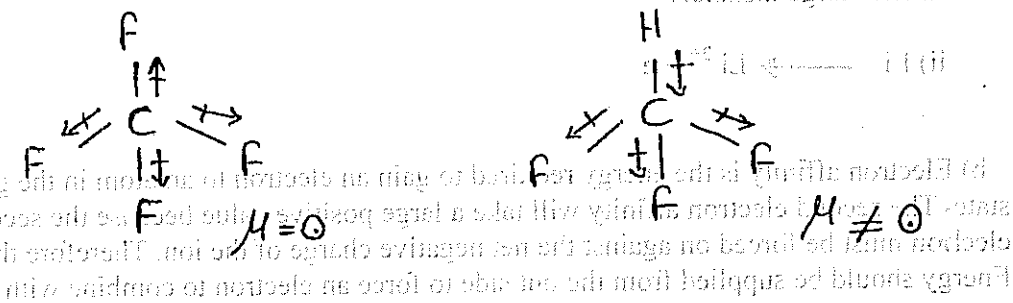
8 — 6

2 ÷ 2

d) i) The measured polarity of a covalent bond of a compound .

Dipole moment (μ) = charge of the atom x bond distance

ii)



The dipole moment of CH_4 is zero since the dipole moment of the C- F bonds are cancelled out. But in CHF_3 there is a dipole moment because there are 2 different types of dipole moments and there is no cancellation of dipole moments.

e) i) The no: of electrons in a cation is less compared to its atom therefore more(+)ve charge on nucleus attract electron cloud towards it. Therefore radius of cation is less than that of the relevant atom.

In anion, there is an en –en repulsion between electrons and it makes the anion larger than the atom.

ii) Ionic potential (Ionic charge / ionic radius) of Li and Mg are fairly close together. That means those two show diagonal relationship.

iii) LiI.Both Li and Cs belong to group I. Electro negativity difference between Li and I is lower than that of Cs and I. Therefore Li I has more covalent character.

iv) Among hydrides of group v elements only NH_3 has H bonds as its intermolecular forces. H bonds are the strongest intermolecular force. Therefore NH_3 has the highest boiling point among these.

v) Na - $1s^2\ 2s^2\ 2p^6 3s^1$

Cu - $1s^2\ 2s^2\ 2p^6 3s^2 3p^6\ 3d^{10}\ 4s^1$

For Na filling orbital level is s .Therefore that belongs to s. For Cu filling orbital level is d and therefore that belongs to d block.

1. (a) (i) Write down a **mathematical expression** for the Raoult's law, as applied to an ideal binary liquid system A and B. Identify all the terms in your expression.

(ii) A solution contain 15.00 g of urea $\{CO(NH_2)_2\}$ in 0.200 kg of water. If vapour pressure of water at 25°C is 23.7 torr, calculate the partial vapour pressure of water in solution at 25°C.

(b) A solution is formed by mixing two volatile solvents C & D, shows large positive deviations from ideal behaviour.

(i) Draw the plot of vapour pressure against mole fraction of D. (vapour pressure of pure C is higher than that of pure D)

(ii) If the solution forms a constant boiling mixture, draw the plot of temperature versus mole fraction of D. (pressure remains constant)

© A volatile liquid A has saturated vapour pressure 68 torr at 30 °C. 5 g of a non volatile substance B, is dissolved in 150 g of liquid A. Then the vapour pressure of the solution is 67 torr. Calculate the relative molecular/molar mass of B if that of liquid A is 30.

PSF 1303

2. (a) The lattice enthalpy of an ionic solid cannot be measured directly by experiment.

(i) Write an equation to define what is meant by lattice enthalpy

(ii) Name all the enthalpy changes that need to be measured to calculate the lattice enthalpy of a simple ionic compound MX. In each case write an equation to represent the change that occurs.

(b) (i) Draw the Born-Haber cycle for CaO given the following data in kJ mol^{-1}.

For Calcium	Atomisation enthalpy	+177
	1st ionisation enthalpy	+590
	2nd ionisation enthalpy	+1145
For Oxygen	Atomisation enthalpy	+249
	1st electron affinity	-141
	2nd electron affinity	+798
	Enthalpy of formation for CaO	-635

(iii) Explain why the second ionisation enthalpy of calcium is greater than the first ionisation enthalpy.

(iii) Compare the first ionisation enthalpy of calcium with that of magnesium.

ශ්‍රී ලංකා විවෘත විශ්වවිද්‍යාලය.

PSE 1303 - 2009/2010

සැවර්ණය II

1) a) i) A හා B අඩංගු තිරපූර්ණ, ද්‍රව්‍යාංශික ද්‍රව පද්ධතියක් සඳහා ද්‍රාවුල්ගේ නියමය ගණිතමය ප්‍රකාශනයක් ලෙස ලියා දක්වන්න. සියළුම පද හඳුන්වන්න.

ii) ද්‍රාවණයක ස්ථලය 0.200 kg තුල යූරියා [$CO(NH_2)_2$] 15.00 g අඩංගු වේ. ස්ථලයේ වාෂ්ප පීඩනය 25 °C දී ටෝර් (torr) 23.7 වේ. 25 °C දී මෙම ද්‍රාවණයේ ආංශික වාෂ්ප පීඩනය ගණනය කරන්න.

b). C හා D නමැති වාෂ්පශීලී ද්‍රව දෙකක් විශ්‍රව්‍ය මිශ්‍රණ සාදන ද්‍රාවණයක් තිරපූර්ණ භාෂ්පීකරණ විශාල ධන අපගමනයක් පෙන්වයි.

i) D හි ද්‍රවලික කලාපයේ සිදියෙන් වාෂ්ප පීඩනය ප්‍රස්ථාරගත කරන්න. (සංශුද්ධ C හි වාෂ්ප පීඩනය සංශුද්ධ D හි සංශුද්ධ වාෂ්ප පීඩනයට වඩා වැඩිය)

ii). මෙම ද්‍රාවණය නියත නාත මිශ්‍රණයක් සාදයි නම්, D හි ද්‍රවල කලාපයේ සිදියෙන් උච්චාන්වය ප්‍රස්ථාරගත කරන්න. (පීඩනය නියතයක්ම පවතී)

c). A නම් වාෂ්පශීලී ද්‍රවයක සංතෘප්ත වාෂ්ප පීඩනය 30 °C දී ටෝර් (torr) 68 වේ. B නම් වාෂ්පශීලී නොවන ද්‍රවයක 5 g ක්, A ද්‍රවය 150 g ක දිය කළ විට ද්‍රාවණයේ වාෂ්ප පීඩනය ටෝර් 67 වේ. A හි සාපේක්ෂ ආණුක / ද්‍රවලික ස්කන්ධය 30 නම්, B හි සාපේක්ෂ ආණුක ස්කන්ධය ගණනය කරන්න.

2) a) ඇයනික සහයක දැලිස ඒන්තැල්පිය පරිසරය මගින් හෙළිවන්ව ඇති බව පෙන්වා දෙන්න.

i) දැලිස ඒන්තැල්පිය ඇරඹ දැක්වෙන ලෙස සමීකරණයක් ලියන්න.

ii) සරල අයනික සංයෝගයක (MX) දැලිස ශක්තියේ ගණනය කිරීම සඳහා යොදා ගත හැකි සියල්ල ශක්තීන් වෙන් වී නම් කරන්න. සෑම වෙන් වීමක්ම නම් කරන්න. සෑම වෙන් වීමක්ම නිරූපණය මින් ලෙස සමීකරණ ලියා දක්වන්න.

b) i) $kJ\ mol^{-1}$ වලින් දී ඇති පහත දැන්න උපයෝගී කරගනිමින් CaO සඳහා බෝන්-හේබ් චක්‍රය ඇඳ දක්වන්න.

කැල්සියම් සඳහා,

පරමාණුකරණ ශක්තිය (atomization E)	+ 177
පළමුවන අයනීකරණ ශක්තිය	+ 590
දෙවන අයනීකරණ ශක්තිය	+ 1145

ඔක්සිජන් සඳහා,

පරමාණුකරණ ශක්තිය (atomization E)	+ 249
පළමු වන ඉලෙක්ට්‍රෝන බන්ධුතාව	- 141
දෙවන ඉලෙක්ට්‍රෝන බන්ධුතාව	+ 798
CaO හි උත්පාදන එන්තැල්පිය	- 635

iii) කැල්සියම්හි දෙවන අයනීකරණ ශක්තිය, පළමු වන අයනීකරණ ශක්තියට වඩා විශාල වන්නේ මන්දැයි පහදන්න.

iv). කැල්සියම් හි පළමුවන අයනීකරණ ශක්තිය, මැග්නීසියම් හි පළමුවන අයනීකරණ ශක්තිය හා සසඳන්න

1(a)

(i) A,B ஆகிய துவித திரவங்களைக் கொண்ட இலட்சியக் கரைசல தொகுதிக்கு பிரயோகிக்கக் கூடிய இரவோற்றின் விதியின் கணித வடிவக் கோவையினை எழுதுக.

(ii) கரைசலொன்று 0.2 kg நீரில் 15.00g யூரியாவினைக் , $(Co(NH_2)_2)$ கொண்டுள்ளது. 25°C இல் நீரின் ஆவியமுக்கம் 23.7 torr எனின் 25°C இல் கரைசலில் நீரினது பகுதி ஆவிஅமுக்கம் யாது?

(b) C,D எனும் இரண்டு ஆவிப்பறப்புடைய கரைப்பான்களைக் கலப்பதன் மூலம் உருவாக்கப்பட்ட கரைசலானது இலட்சிய நடத்தையிலிருந்து பெரிய நேர் விலகலைக் காட்டுகின்றது

(i) ஆவிஅமுக்கம் எதிர் D இன் மூல்ப்பின்னம் வரைபினை வரைக. (தூய C யினது ஆவியமுக்கம் தூய D இனது ஆவியமுக்கத்தினை விட அதிகமாகும்.)

(ii) இக்கரைசல் மாறா கொதிநிலைக் கலவையினை உருவாக்கின் வெப்பநிலை எதிர் D இன் மூல்ப்பின்ன வரைபினை வரைக. (அமுக்கம் மாறாது உள்ளது)

(c) ஆவிப்பறப்புடைய திரவம் A, 30°C இல் 68 torr நிர்ம்பல் ஆவியமுக்கத்தினைக் கொண்டுள்ளது. 5g ஆவிப்பறப்பற்ற சேர்வை B ஆனது A இன் 150g இல் கரைக்கப்பட்டது. அதன் பின்னர் கரைசலின் ஆவியமுக்கம் 67 torr ஆகும். திரவம் A இன் மூலக்கூற்று / மூலர்த்திணிவு 30 எனின் B இன் மூலர்த்திணிவினைக் கணிக்க.

2

(a) அயன் திண்மம் ஒன்றின் சாலகச் சக்தியினைப் பரிசோதனை ரீதியாக நேரடியாகத் துணியமுடியாது.

(i) சாலகச் சக்தி என்பதால் யாது விளங்குகின்றீர் என்பதனை சமன்பாடு ஒன்றின் மூலம் வரையறுக்க.

(ii) எளிய அயன் சேர்வை MX இன் சாலகச் சக்தியினைக் கணிப்பதற்கு அளவிடப்பட வேண்டிய சகல எந்தலபி மாற்றங்களையும் பெயரிடுக. ஒவ்வொரு சந்தர்ப்பத்திலும் நடைபெறும் மாற்றங்களைச் சுட்டிக் காட்டுவதற்கு சமன்பாடு ஒன்றினை எழுதுக.

(b)

(i) CaO இன் Born-Haber சக்கரத்தினை வரைக, பின்வரும் தரவுகள் $KJmol^{-1}$ இல் தரப்பட்டுள்ளன.

கல்சியத்திற்கு	அணுவாதல் வெப்பவுள்ளுறை	+ 177
	முதலாம் அயனாக்கற்சக்தி	+ 590
	இரண்டாம் அயனாக்கற்சக்தி	+ 1145

ஒட்சிசனிற்கு	அணுவாதல் வெப்பவுள்ளுறை	+ 249
	முதலாம் இலத்திரன் நாட்டற்சக்தி	- 141
	இரண்டாம் இலத்திரன் நாட்டற்சக்தி	+ 798
	CaO இன் தோன்றல் வெப்பவுள்ளுறை	- 635

a)

(ii) கல்சியத்தின் இரண்டாம் அயனாக்கல் வெப்பவுள்ளுறை முதலாம் அயனாதல் வெப்பவுள்ளுறையினை விட அதிகமாகும் என விளக்குக

(iii) கல்சியத்தின் முதலாம் அயனாக்கல் வெப்பவுள்ளுறை யினை மக்னீசியத்துடன் ஒப்பிடுக.

b)

PSF 1303

Home Assignment II - Answer guide (2009/2010)

a) i)

$$P_A = P^\circ_A \, x_A$$

or

$$P_A = P^\circ_A \, (1 - x_B)$$

P_A = Patial Vapour pressure of compound A

P°_A = Vapour pressure of pure A.

x_B = Mole fraction of A

x_B = Mole fraction of B

ii) $P_{H_2O} = P^\circ_{H_2O} \cdot x_{H_2O}$

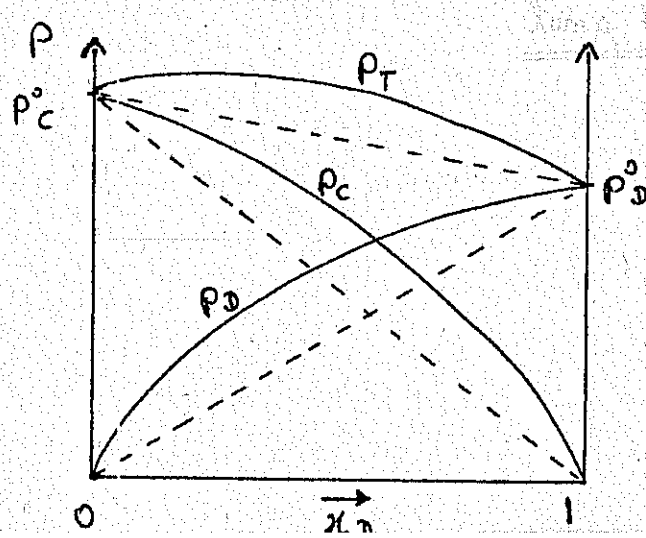
$$x_{H_2O} = \frac{0.2 \times 10^3 g / 18 g mol^{-1}}{0.2 \times 10^3 g / 18 g mol^{-1} + 15.00 g / 60 g mol^{-1}}$$

$x_{H_2O} = 0.977$

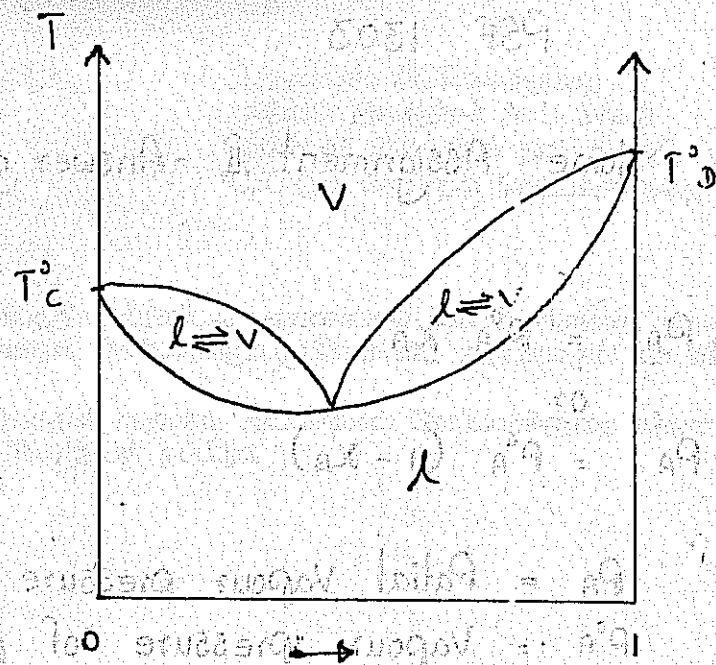
$P_{H_2O} = 23.7 \text{ torr} \times 0.977$

$P_{H_2O} = 23.17 \text{ torr}$

b) i)



ii)



c) $P_A = P^\circ_A \chi_A$

$P_A = P^\circ_A (1 - \chi_B)$

$\chi_B = \dfrac{m_B/M_B}{m_B/M_B + m_A/M_A}$

$67 \text{ torr} = 68 \text{ torr} (1 - \chi_B)$

$68 \chi_B = 1$

$\chi_B = \dfrac{1}{68}$

$\underline{\underline{M_B = 67 \text{ g mol}^{-1}}}$

a) i)

$$MX_{(s)} \longrightarrow M^+_{(g)} + X^-_{(g)}$$

ii)

i) Atomization of $M_{(s)}$

$$M_{(s)} \xrightarrow{\Delta H \text{ atom}} M_{(g)}$$

ii) Ionization enthalpy of $M_{(g)}$

$$M_{(g)} \longrightarrow M^+_{(g)} + e$$

iii) Atomization enthalpy of $X_{2\,(g)}$

$$\tfrac{1}{2}\, X_{2\,(g)} \longrightarrow X_{(g)}$$

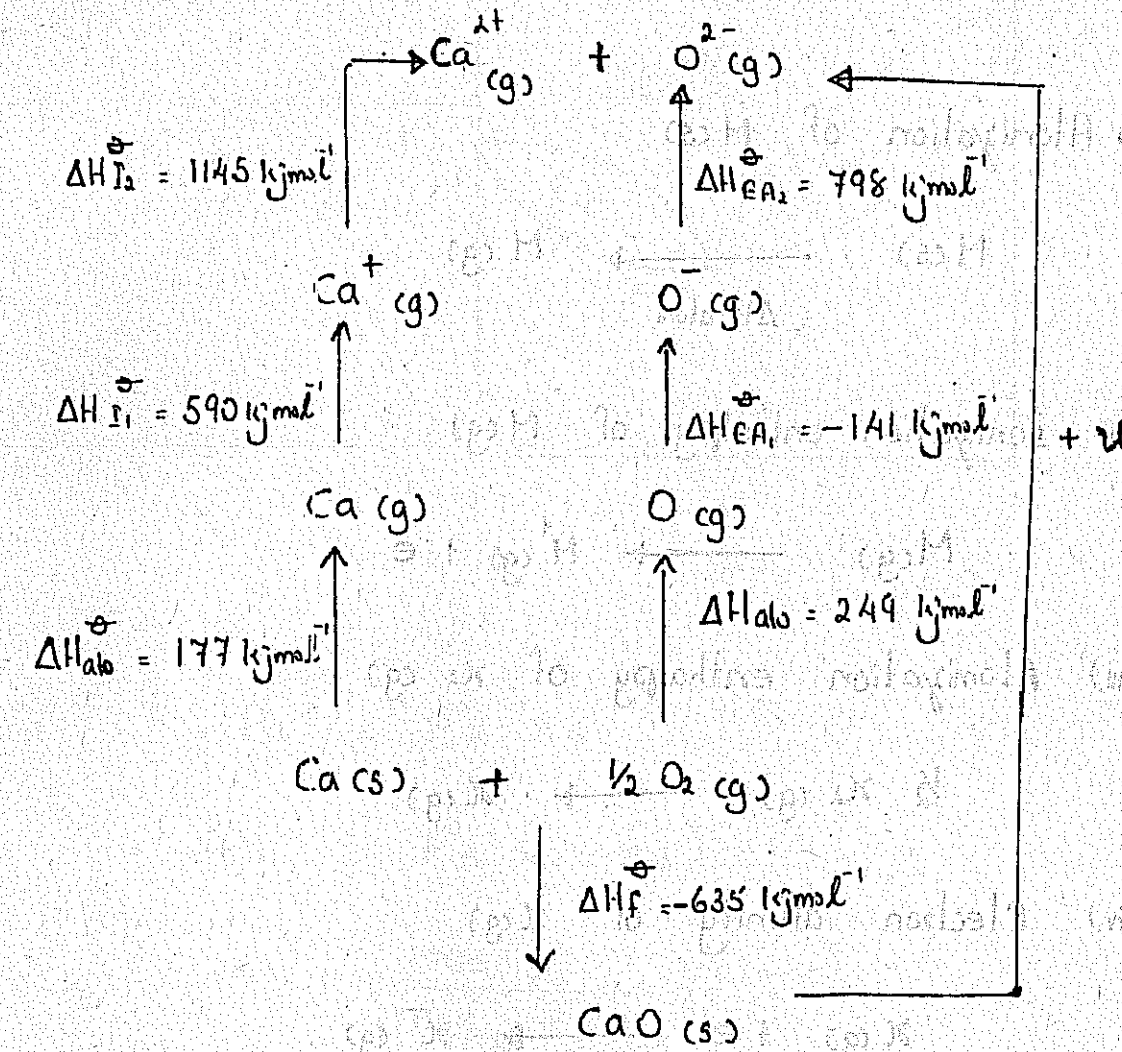
iv) Electron affinity of $X_{(g)}$

$$X_{(g)} + e \longrightarrow X^-_{(g)}$$

v) Enthalpy of formation of MX

$$M_{(s)} + \tfrac{1}{2} X_{2(g)} \longrightarrow MX_{(s)}$$

b)



$177\ kjmol^{-1} + 590\ kjmol^{-1} + 1145\ kjmol^{-1} + 249\ kjmol^{-1} + (-141\ kjmol^{-1}) + 798\ kjmol^{-1}$

$=$

$-635\ kjmol^{-1} + U$

$U = 3453\ kjmol^{-1}$

iii) Ca - $1s^2\ 2s^2\ 2p^6\ 3s^2\ 3p^6\ 4s^2$

During the 1st ionization process, 1 electron can easily removed from 4s orbital.

But in the second stage effective nuclear charge on 4s' electron is large and therefore need more energy to remove the 2nd electron.

iv). Mg - $1s^2\ 2s^2\ 2p^6\ 3s^2$

Ca - $1s^2\ 2s^2\ 2p^6\ 3s^2\ 3p^6\ 4s^2$

Both Mg & Ca belong to Group II, Ca has higher atomic radius than that of Mg. Therefore Mg nucleus attract 3s electron more towards it and need more energy to remove that. Therefore 1st ionization energy of Mg is a larger value compared to that of Ca.