# **PUBLIC EXAMINATION 2019**

# CHEMISTRY

# **QUESTION TYPE B**

1.B)First Order

2.A)Diethyl ether

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# **QUESTION TYPE A**

3.A)(ii)and (iii)	1. A)Diethyl ether
4.D)(1)-(iii),(2)-(i),(3)-(iv),(4)-(ii)	2. B)First Order
5.D)Pentagonal bipyramidal	3. B)C <sub>6</sub> H <sub>5</sub> NH-NH-C <sub>6</sub> H <sub>5</sub>
6.A)4.90BM	4. D)Pentagonal bipyramidal
7.D)Strong reducing agents	5. D)CH <sub>3</sub> COCH <sub>3</sub>
8.C)dynamic	6. C)C <sub>1</sub> -C <sub>2</sub>
$9.C)C_1-C_2$	7. C)w-PΔV
10.C)ohm <sup>-1</sup> m <sup>-1</sup>	8. A)-OH
11.C)w-P∆V	9. A)(ii)and (iii)
12.A)-OH	10. D)(1)-(iii),(2)-(i),(3)-(iv),(4)-(ii)
13.A)Ionic	11. D)Strong reducing agents
$14.B)C_6H_5NH-NH-C_6H_5$	12. A)Ionic
15.D)CH <sub>3</sub> COCH <sub>3</sub>	13. C)dynamic

14. A)4.90BM

15. C) $ohm^{-1}m^{-1}$ 

**16.**The higher the nuclear charge of

protons in the nucleus, the higher is the ionization energy.

Because of the higher nuclear charge, the electrons are bound with more force and hence higher energywill be required for their removal.

For instance, magnesium has higher nuclearcharge (12 protons) as compared to sodium (11 protons). Hence ionization energyin case of magnesium is higher as compared to sodium.

17. These elements shows the oxidation states of +2, +3, +4, +5 and +6. Out

of these, +4 oxidation state is most common state.

**18.** A small quantity of Radioactive CO<sup>\*</sup><sub>2</sub> containing radioactive oxygen  $O^{18}$  is

mixed with ordinary carbondioxide and the process is carried out.

It has beenfound that oxygen gas evolved along with sugar formation is non-radioactive.

Therefore O<sub>2</sub> produced comes from water and not from carbondioxide. So the

correct mechanism is as follows.

 $6CO^{*}_{2} + 6H_{2}O \rightarrow C_{6}H_{12}O^{*}_{6} + 6O_{2}$ 

**19.** glassy orvitreous state is a condition in which certain substance can exist, lying between the solid and liquid states.

20. Complete reduction of glucose with

concentrated hydriodic acid in the presence of red phosphorous produces

n-hexane as the major product.

This indicates that the six carbon atoms in the glucose molecule form an



This is used as Urinary antiseptic in medicine, in the name of 'Urotropine'.

24.II reaction will go faster then I reaction. This is due to the activation energy is minimum the reaction go faster

 $=\frac{(IP)_{F}+(EA)_{F}}{(EA)_{F}}$ 

25.

Electronegativity of fluorine

$$=\frac{17.4+3.62}{5.6} = \frac{21.02}{5.6} = 3.75$$

26.

Lanthanides		Actinides	
i)	Binding energies of 4f electrons are higher.	i)	Binding energies of 5f electrons are lower.
ii)	Maximum oxidation satate exhibited by lanthanides is +4 e.g. Ce <sup>4+</sup>	ii)	Due to lower binding energies they show higher oxidation states such as +4, +5 and +6. Uranium exhibits +6 oxidation state in UF <sub>6</sub> and UO <sub>2</sub> Cl <sub>2</sub>
iii)	4f electrons have greater shielding effect.	iii)	5f electrons have poor shielding effect.

27. Let 'a' and 'b' be the number of  $\alpha~\beta$  particles emitted during the change

 $_{90}$ Th<sup>232</sup>  $\rightarrow {}_{82}$ pb<sup>208</sup> + a  $_{2}$ He<sup>4</sup> + b  $_{-1}$ e<sup>0</sup>

Comparing the mass numbers,

$$232 = 208 + 4a + b \times 0$$
  

$$4a = 232 - 208$$
  

$$= 24$$
  

$$a = 6$$

Comparing the atomic numbers

 $90 = 82 + 2 \times a + (-1)b$ = 82 + 2a - b 2a-b= 90 - 82 = 8 2(6) - b = 8 b = 12-8 = 4

Number of  $\alpha$  - particle emitted = 6 Number of  $\beta$  - particles emitted = 4

28. 'Q' is defined as the ratio of product of initial concentrations

of products to the product of initial concentrations of reactants under nonequilibrium

conditions. For example, in the equilibrium.

 $aA + bB \quad \overleftarrow{\leftarrow} \rightarrow \ lL + mM$ 

K.RAMALINGAM PG Asst.in Chemistry SHRI RENGA EDUCATIONAL INSTITUTIONS RAYARPALAYAM NAMAKKAL MAIN ROAD 9788855343

Let [A], [B], [L] and [M] be the actual concentrations present before the occurrence of equilibrium. These concentrations are considered as the nonequilibrium concentration conditions and the reaction quotient



'Q' is given as

29. (i) It should have a suitable colour.

(ii) It should be able to fix itself or be capable of being fixed to the fabric.

(iii)It should be fast to light.

30. The standard emf  $E_0$ , of a cell is the standard reduction potential of

right-hand electrode (cathode) minus the standard reduction potential of the

left-hand electrode (anode). That is,

 $E_{ocell} = E_{oright} - E_{o}$  left

= Cathode potential – Anode potential

31. When equal amounts of d-isomer and l-isomer are mixed one gets a

"racemic mixture" and this process is called racemisation. A racemic mixture

becomes optically inactive. Because, in this mixture rotation towards

clockwise direction by the dextro isomers is

compensated by the rotation

towards the anticlockwise direction by the laevo isomers.Eg:d,l lactic acid

32.



(1, 3, 4, 5, 6, pentahydroxy-2-hexanone)

Fructose molecules have 4 chiral centers of their carbon atoms. A maximum of  $2^n$  optical isomers can occur for a molecule, where 'n' is the number of chiral centers. Since there are 4 chiral centers in fructose, it should have  $2^4 = 16$  optical isomers.

#### 33.

Log $k_1/k_2$ =E<sub>a</sub>/2.303 R[1/T<sub>2</sub> -1/T<sub>1</sub>] ∴ $k_1/k_2$ =100000J Mole<sup>-1</sup>/2.303\* 8 314JK<sup>-1</sup>mol<sup>-1</sup> = -0.5745 ∴  $k_1/k_2$  antilog (-0.5745) = 0.2664  $k_2$  =1/0.2664 $k_1$  = 3.75 times  $k_1$ . ∴  $k_2$ , the rate constant at 35°C will be 3.75 times the rate constant at 25°C.

34.a) i) a)  $CO_2$ -sp b) $NO_2$ -sp<sup>2</sup>

c)ClO<sub>2</sub> -sp<sup>3</sup> d)XeF<sub>2</sub> -sp<sup>3</sup>d

ii) "it is impossible to measure simultaneously both the position and velocity (or momentum) of a microscopic particle with absolute

# accuracy or certainty."

Mathematically, uncertainty principle can be put as

$$\Delta x.\Delta p \ge \frac{h}{4\pi}$$

follows.

where,  $\Delta x =$  uncertainity in the position of the particle and  $\Delta p =$  uncertainity in the momentum of the particle.

Or

b)i) One of the familiar characteristic of p-block elements is to show inert paireffect i.e. the tendency of being less availability for ns electron in bonding.The inert pair effect increases down the group with the increase in atomic

number.

ii) 1) Silicones act as excellent insulators for electric motors and other appliances

as they can withstand high temperatures.

2) Straight chain polymers of 20 to 500 units are used as silicone fluids. They

are water repellent because of the organic side group. These polymers are

used in waterproofing textiles, as lubricants and as polish.

3) Silicone rubber retain their elasticity even at low temperatures and resist

chemical attack. They are mixed with paints to make them damp-resistant.

 $(35.a)i)[Ti(H_2O)_6]^{3+}$  is coloured because Ti<sup>3+</sup> one electron present in d orbital The colour of transition metal ions is due to the presence of unpaired electrons in it and the energy gap between two energy levels in the same d-subshell being small. Sc<sup>3+</sup> ions are also colourless because of the absence of d-electrons.

ii) The articles to be plated with chromium are made the cathode in an electrolytic

bath consisting of chromic acid and sulphuric acid while the anode is made

of a plate of lead. During electrolysis chromium deposits on the article

(cathode). Generally the articles are first plated with nickel and then subjected

to chromium plating.

Or

b)i)

- 1) Every metal atom has two types of valencies
  - i) Primary valency or ionisable valency

ii) Secondary valency or non ionisable valency

2) The primary valency corresponds to the oxidation state of the metal ion.

The primary valency of the metal ion is always satisfied by negative ions. 3) Secondary valency corresponds to the coordination number of the metal ion or atom. The secondary valencies may be

satisfied by either negative

ions or neutral molecules.

4) The molecules or ion that satisfy secondary valencies are called ligands.

5) The ligands which satisfy secondary valencies must project in definite

directions in space. So the secondary valencies are directional in nature

whereas the primary valencies are non-directional in nature.

6) The ligands have unshared pair of electrons.

These unshared pair of electrons

are donated to central metal ion or atom in a

compound. Such compounds

are called coordination compounds.

36.a)i) The lattice points in molecular crystals consist of molecules which do not

carry any charge. The forces binding the molecules together are of two types

(i) Dipole-dipole interaction and (ii) Vanderwaal's forces. Dipole-dipole forces

occur in solids which consists of polar molecules e.g., ice. The Vanderwaal's

forces are more general and occur in all kinds of molecular solids.

ii) 1. The heats of vapourisation of ionic crystals are high.

2. The vapour pressure of ionic crystals at ordinary temperature are very low.

3. The melting and boiling points of ionic crystals are very high.

4. Ionic crystals are hard and brittle.

5. Ionic crystals are insulators in the solid state.6. Ionic crystals are soluble in water and also in other polar solvents.

Or

b) Consider a general chemical equilibrium reaction in which the reactants and

products are in gaseous phases,

 $aA + bB + cC + \dots$ them  $K_P = P_L^{\ l} p_M^{\ m} p_N^{\ n} \dots / p_A^{\ a} p_B^{\ b} p_C^{\ c} \dots$ where p is the partial pressure of the respective

gases. In terms of molar

concentrations of reactants and products

 $K_{c} = [L]^{l}[M]^{m}[N]^{n...}/[A]^{a}[B]^{b}[C]^{c...}$ 

For any gaseous component 'i' in a mixture, its partial pressure 'p<sub>i</sub>' is related

to its molar concentration 'C<sub>i</sub>' as

 $C_i = p_i/RT$ , since  $p_i = n_i/V$  RT

Where  $n_i/V=C_i=$  number of moles of i per litre. V = volume in litres.

Substituting, concentration terms by partial pressures,

$$K_{c} = \frac{(p_{L}/RT)^{1}(p_{M}/RT)^{m}(p_{N}/RT)^{n}...}{(p_{A}/RT)^{a}(p_{B}/RT)^{b}(p_{C}/RT)^{c}...}$$

$$=\frac{p_{L}^{1}p_{M}^{m}p_{N}^{n}....}{p_{A}^{a}p_{B}^{b}p_{C}^{e}.....}\left(\frac{1}{RT}\right)^{(1+m+n+...)-(a+b+c+...)}$$

$$= \frac{K_p}{(RT)^{\Delta ng}}$$
 and  $\therefore K_p = K_c (RT)^{\Delta ng}$ 

where  $\Delta n_g =$  total number of stoichiometric moles of gaseous products - total

number of stoichiometric moles of gaseous reactants.

37.a)i) a)Paint	<b>Dispersion</b> Phase	Medium
	Solid	Liquid
b) Froths o	f air Gas	Liquid

ii) A substance which destroys the activity of the catalyst is called a poison

and the process is called catalytic poisoning. Some of the examples are

(i) The platinum catalyst used in the oxidation of SO<sub>2</sub> in contact process

is poisoned by arsenious oxide.

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b)i) A buffer solution is one which maintains its pH fairly constant even

upon the addition of small amounts of acid or base. In other words, a buffer solution resists (or buffers) a change in its pH.

That is, we can add a small amount of an acid or base to a buffer solution

and the pH will change very little.

ii) Ka =  $1.8 \times 10^{-5}$ 

 $pK_a = -\log(1.8 \times 10^{-5}) = 4.7447$ 

 $pH = pK_a + log[salt]/[acid]$ 

... Henderson - Hasselbalch equation

 $= 4.7447 + \log 0.5/0.5.$ 

pH = 4.7447 + log1

= 4.7447

38.a)i) The removal of water from the protonated species is facilitated by

anhydrous zinc chloride. Reactivitiy in this reaction with respect to alcohols

is  $3_0 > 2_0 > 1_0$ . This reaction is used in Lucas Test.

 $\begin{array}{ccc} (CH_3)_3 \text{ COH} & \xrightarrow{\text{con.HCl}} & (CH_3)_3 \text{ C-Cl} + H_2 O \\ & & \text{anhyd. ZnCl}_2 & (Turbidity appears immediately) \\ & & \text{at room temperature} \end{array}$ 

(CH<sub>3</sub>)<sub>2</sub> CHOH 2° alcohol  $\xrightarrow[\text{anhyd. ZnCl}_2]{\text{cm.HCl}} (CH_3)_2 CHCl + H_2O$ 

(Turbidity appears after 5-10 minutes) at room temperature

$$\begin{array}{c} \text{CH}_3\text{CH}_2\text{OH} & \xrightarrow{\text{Conc. HCl}} \\ \xrightarrow{\text{1° alcohol}} \\ \end{array}$$

$$CH_3CH_2Cl + H_2O$$

(No turbidity appears easily) but only on heating

ii) –OH group, –OMe, group increase the reactivity of thebenzene ring with respect to electrophilic attack and is **ortho, para - directing.** With a mixture of con.HNO3 and con. H2SO4 it gives a mixture of ortho and para nitro anisole.

# Con I

# b)i) Reducing property

Formic acid is unique because it contains both an aldehyde group and carboxyl group also. Hence it **can act as a reducing** agent. It reduces Fehling's solution, Tollens reagent and decolourises pink coloured KMnO4 solution.

(a) Formic acid reduces ammoniacal silver nitrate solution (Tollen's reagent) to metallic silver.
HCOOH + Ag2O → H2O + CO2 + 2Ag↓ (metallic silver)
(b) Formic acid reduces Fehling's solution. It reduces blue coloured cupric ions to red coloured cupric ions. HCOO- + 2Cu<sup>2+</sup> + 5OH<sup>-</sup>→ CO3<sup>2-</sup> + Cu2O + 3H2O (blue) (red)
(c) On the otherhand when sodium formate is heated to 360°C it decomposes

to hydrogen and sodium oxalate.

$H-COONa \longrightarrow H_2 + Q$	
H COONa	COONa
Sodium formate	Sodium oxalate

ii) Aniline reacts with chloroform and alcoholic KOH to give an offensive

smelling liquid, phenyl isocyanide.

C6H5NH2 + CHCl3 + 3KOH→ C6H5NC + 3KCl + 3H2O (phenyl isocyanide)