Marking Scheme			
Chemistry Sample Paper (2024-25)	CHE-856	Class: 11 th	
1. (b) 6 unpaired e^{-1}		(1)	
2. (b) He, Na and Ar		(1)	
3. (c) F <cl>Br>l</cl>		(1)	
4. (d) O_2^{2-}		(1)	
5. (b) 18 gm/mol		(1)	
6. (c) q=0		(1)	
7. (a) +2		(1)	
8. (c) < ΔU°		(1)	
9. (c)		(1)	
10. (a) 0		(1)	
11. (a) planar		(1)	
12. (a) electron deficient species		(1)	
13. (b) 6.022x10 ²¹ molecules		(1)	
14. (a) 1		(1)	
15. (a) 2		(1)	
16. (d) Assertion(A) is false but Reasor	n(R) is true	(1)	
17. (a) Assertion(A) and Reason(R) are	correct and (R) is correct ex	planation of	
Assertion(A)		(1)	
18. (a) Assertion(A) and Reason(R) are	correct and (R) is correct exp	planation of	
Assertion(A)		(1)	
19. (a) n = 5		(1)	
(b) 16e			
20. (a) as the volume increases, pressu	are decreases so equilibrium	move in forward	

20. (a) as the volume increases, pressure decreases so equilibrium move in forward direction where number of moles increases. (1)

(b) KC (Reverse) =
$$\frac{1}{8.3x15^{-3}} = 120.48$$

$$21.16H^{+}+Cr_{2}O_{7}^{2^{-}}+6e^{-} \rightarrow 2Cr^{3^{+}}+7H_{2}O$$

$$[C_{2}H_{4}O+H_{2}O \rightarrow C_{2}H_{4}O_{2}+2e^{-}+2H^{+}]3$$

$$\overline{Cr_{2}O_{7}^{2^{-}}+16H^{+}+3C_{2}H_{4}O+3H_{2}O} \rightarrow 2Cr^{3^{+}}+7H_{2}O+3C_{2}H_{4}O_{2}+6H^{+}}$$

$$\overline{Cr_{2}O_{7}^{2^{-}}+3C_{2}H_{4}O_{2}+10H^{+}} \rightarrow 2Cr^{3^{+}}+3C_{2}H_{4}O_{2}+4H_{2}O$$
(2)
22

$$CH_{2} = CH - CH_{2} = CH_{2} - CH_{2} - CH_{2} - CH_{2} = CH_{2$$

23. Buffer solution is the solution which resist change in the pH value of solution when small amount of acid or base is added to it.

Buffer solutions are of two types

(a) Acidic buffer:- It contains equimolar quantities of weak acid and its salt with strong base.

Example acetic acid and sodium acetate

4

(b)Basic Buffer:- It contains equimolar quantities of weak base and its salt with strong acid

Example ammonium hydroxide and ammonium chloride (2)

24.
$$\bigcirc$$
 + CH₃COCl $\xrightarrow{\text{Anhy}}_{\text{AlCl3}}$ \bigcirc + HCl

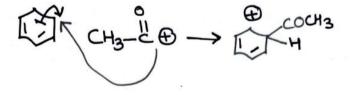
Mechanism

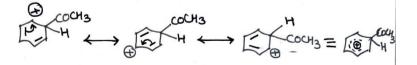
(i) Generation of Electrophile

$$CH_3CO(1 + AICI_3 \rightarrow CH_3CO + AICI_4)$$

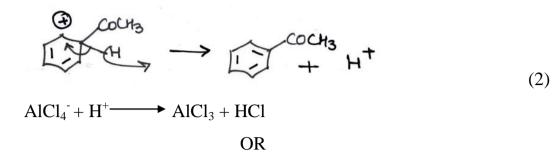
COCH.

(ii) Attack of electrophile to form intermediate carbocation.



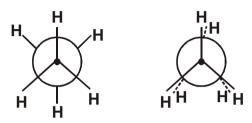


(iii) Loss of proton from carbocation



Newman's Representation

Staggerred



Eclipsed

(1)

Staggerred Conformation is more stable due to less repulsions. (1)

$$25.(a) S < P < O < N \tag{1}$$

$$(b) P < S < N < O \tag{1}$$

(a)
$$3s^2 3p^4$$

Group $\longrightarrow 16^{th}$ group
Block \longrightarrow p-block
Period $\longrightarrow 3^{rd}$ period (1)
(b) $4s^2 \quad 3d^2$
Group $\longrightarrow 4^{th}$ group
Block \longrightarrow d-block
Period $\longrightarrow 4^{th}$ period (1)

SECTION-C

26.(a) 1 mole C_2H_6 contains 2	2 moles of carbon, 3 mole C_2H_6 contains $3x2=6$
moles carbon.	(1

(b) 1 mole C_2H_6 contains 6 mole hydrogen, 3 mole C_2H_6 contains 6x3=18mole hydrogen. (1)

(c) 1 mc	ole C ₂ F	$H_6 = 6.022 \text{ x } 10^{23} \text{ molecules}$	
3 mole $C_2H_6 = 3x6.022 \times 10^{23}$ molecules			
		$= 18.066 \text{ x } 10^{23} \text{ molecules}$	(1)
27. (a)	С	non metal	(1)
(a)	А	alkali metal	(1)
(b)	B alkaline earth metal		(1)
28. ΔG	$= \Delta H - T \Delta S \tag{1}$		
$= 40.63 \times 1000 - 300 \times 108.8$			
	= 79	90 J/mol	(1)
ΔG	$=$ +ve \therefore reaction is not feasible (1)		

$$SO_{2}(g) + \frac{1}{2}O_{2} \rightarrow SO_{3}(g)$$

$$\Delta H = \Sigma \Delta_{f}H^{\circ} \text{ products} - \Sigma \Delta_{f}H^{\circ} \text{ reactants}$$
(1)
$$= \Delta_{f}H^{\circ} SO_{3} - \Delta_{f}H^{\circ} SO_{2}$$

$$= -395.6 - (-296.6)$$
(1)
$$= -395.6 + 296.6$$

$$= -99 \text{ KJ}$$
 (1)

29.(a) AgCl \equiv Cl

$$108+35.5 \quad 35.5$$

$$143.5 \quad 35.5$$
% of Cl = $\frac{35.5}{143.5} \times \frac{\text{Amt.of AgCl formed}}{\text{Wt.of Organic Compound}} \times 100$
% of Cl = $\frac{35.5}{143.5} \times \frac{0.5740}{0.3780} \times 100$
= 37.57%
(2)

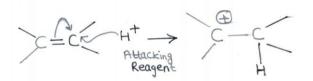
(b) Paper chromatography is a type of partition chromatography which is based upon the differences in the tendencies of substance to distribute between two phases. (1)

OR

- (c) The process of displacement of σ electrons along the saturated carbon chain due to the presence of a polar covalent bond at one end of the chain is called inductive effect (I effect). It is of two types.
 - (i) + I effect \rightarrow substituent has less e⁻ attracting power than H.
 - (ii) I effect \rightarrow substituent has more e⁻ attracting power than H.
 - + I effect example CH_3 , C_2H_5
 - I effect example $-NO_2$, -CN, -X (1¹/₂)

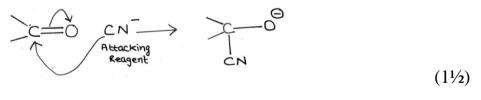
Electromeric effect is the complete transfer of shared pair of πe^{-1} to one of the atom joined by multiple bond. It is of two types

- (i) + E effect
- (ii) E effect
- (i) + E effect is when π e⁻s are transferred to atom to which attacking reagent get attached example



(ii) - E effect is when π bond e⁻ all transferred to atom other than the one to which reagent get attached.

Example



30.(a) NaOH → Na⁺ + OH⁻ 0.01 M 0.01M $K_W = [H^+] [OH^-] = 10^{-14}$ $[H^+] = \frac{10^{-14}}{0.01} \times 100 = 10^{-12}$ (1) pH = - log [H⁺]

$$= -\log 10^{-12}$$

$$= 12$$
(1)
(b) SO₄²⁻
(1)
31.(a) N₂ + 3H₂ \rightarrow 2NH₃

$$\Delta ng = 2 - 4 = -2$$

$$\Delta H = \Delta U + \Delta ng RT$$

$$= -87.425 \times 1000 + (-2) \times 8.314 \times 298$$
(2)
$$= -87425 - 4955.14$$

$$= -77514.7J$$

OR

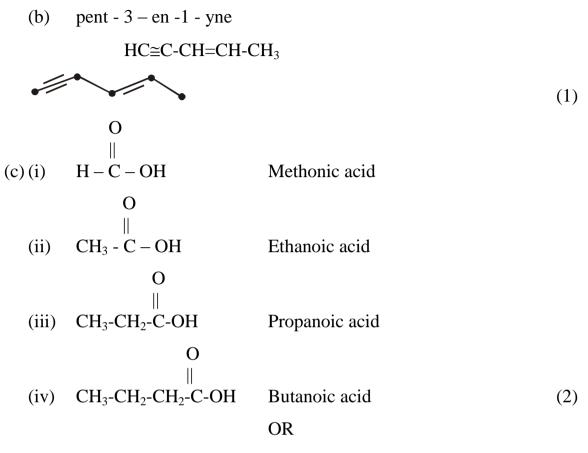
(a)
$$\Delta H = \Delta U$$
 when $\Delta ng=0$
eg. $H_2 + I_2 \rightarrow 2HI$
 $\Delta ng = 2-2=0$
 $\therefore \Delta H = \Delta U + 0RT$
 $\therefore \Delta H = \Delta U$
 $\Delta H = \Delta U$
 $\Delta H = \Delta U$
 $\Delta H = \Delta U + P\Delta V$
when $\Delta V = 0$ then $\Delta H = \Delta U$ (2)

OR

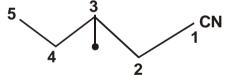
any other relevant example.

$\Delta ng = 1-3 = -2$	$\Delta H = \Delta U - 2RT$	$\Delta H < \Delta U$	
$\Delta ng = 1 - 1 = 0$	$\Delta H = \Delta U {+} O$		
$\Delta ng = 1 - 0 = 1$	$\Delta H = \Delta U {+} RT$	ΔΗ>ΔU	
$\Delta H < \Delta U$ in (i)			(1)
$\Delta H > \Delta U$ in (iii)			(1)
	$\Delta ng = 1 - 1 = 0$ $\Delta ng = 1 - 0 = 1$ $\Delta H < \Delta U \text{ in (i)}$	$\Delta ng = 1 - 1 = 0 \qquad \Delta H = \Delta U + O$ $\Delta ng = 1 - 0 = 1 \qquad \Delta H = \Delta U + RT$ $\Delta H < \Delta U \text{ in (i)}$	$\Delta ng = 1 - 0 = 1$ $\Delta H = \Delta U + RT$ $\Delta H > \Delta U$ $\Delta H < \Delta U$ in (i)

32.(a) H 1 0 2 3 4 5



(i)



3-methyl pentane nitrile (1)

(ii) 1 2 3 4 $CH_3 - CH - CH - CH_3$ | | OH Br

3-Bromobutan-2-01 (1)

SECTION-E

$$33.(a) \Delta V = 45 \text{ x} \frac{2}{100} = 0.9 \tag{1}$$

$$m = \frac{40}{1000}$$

$$\Delta x = \frac{h}{4\pi m \Delta v}$$
(1)

$$= \frac{6.626 \times 10^{-34}}{4 \times 3.14 \times 40 \times 10^{-3} \times 0.9}$$

$$= 1.46 \times 10^{-33} m$$
(1)
Orbit Orbital
(a) It I well defined It is region in space

(a)	It I well defined	it is region in space
	circular path	around nucleus
	around which	where the probability
	e ⁻ revolve	to find e is maximum
(b)	Represent planar	three dimensional motion of electron
	motion	

OR

$$(a)\frac{\lambda_{A}}{\lambda_{B}} = \frac{h}{P_{A}}\frac{P_{B}}{h} = \frac{P_{B}}{P_{A}}$$
(1)

$$P_B = \frac{P_A}{2}$$

(b)

$$\frac{\lambda_A}{\lambda_B} = \frac{P_A}{2.PA} = \frac{1}{2} \tag{1}$$

$$\frac{5x10^{-8}}{\lambda B} = \frac{1}{2} \therefore \lambda B = 10x10^{-8}m$$

= 10⁻⁷m (1)

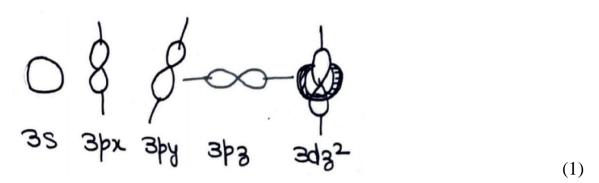
(i) It is obtained when It is obtained when white radiation emitted by white light is passed through solution and the excited substance are analysed with transmitted light is spectroscope analysed through spectroscope

(ii) Emission spectrum Consist of dark lines consist of bright lines in otherwise coloured lines separated continuous spectrum by dark spaces. 34.(a) P(Ground State) $1s^2 2s^2 2p^6 3s^2 3px^1 3py^1 3pz^1$

P(Excited State) $ls^2 2s^2 2p^6 3px^1 3py^1 3pz^1 3dz^{2^1}$

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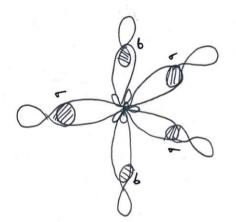
(2)

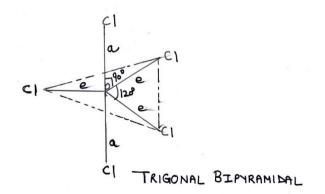


5 orbital hybridise and form 5 new orbitals of same energy and same shape.



5 sp3d hybrid orbitals adopt trigonal bipyramidal arrangement which overlap with 3pz orbital to form 5 σ bonds





(1)

Axial bonds are longer than equatorial bonds due to greater repulsions from other bonds in axial position. (1)

(b) BeH₂ is linear molecule. Therefore the resultant dipole moment of twoBe-H bonds get cancelled giving zero dipole moment.

$$\begin{array}{c} \longleftarrow \\ H - Be - H \\ \mu = 0 \end{array}$$
(2)

(a)
$$O_2 = \sigma ls^2 \sigma^* ls^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2pz^2 \pi 2px^2 = \pi 2py^2 \pi^* 2px^1 = \pi^* 2py^1$$

 $O_2^+ = \sigma ls^2 \sigma^* ls^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2pz^2 \pi 2px^2 = \pi 2py^2 \pi^* 2px^1 = \pi^* 2py^0$
 $O_2^- = \sigma ls^2 \sigma^* ls^2 \sigma 2s^2 \sigma^* 2s^2 \sigma 2pz^2 \pi 2px^2 = \pi 2py^2 \pi^* 2px^2 = \pi^* 2py^1$ (1)

Bond order
$$O_2 = \frac{8-4}{2} = \frac{4}{2} = 2$$

 $O_2^+ = \frac{8-3}{2} = \frac{5}{2} = 2.5$
 $O_2^- = \frac{8-5}{2} = \frac{3}{2} = 1.5$
(1)

Higher is the bond order, more is the stability.

$$O_2^+ > O_2 > O_2^-$$
 (1)

(b) CO₂ has zero dipole moment so CO₂ is linear as two C=O bond moments get cancelled where as H₂O molecule has resultant dipole moment. Two O-H bonds are arranged in angular shape and the bond moment of two O-H bonds give resultant dipole moment. (2)

(i)
$$CH_2 = CH_2 + H_2 \xrightarrow{\text{Ni}} CH_3 - CH_3$$
 (1)
523-573K

(ii)
$$2CH_3CH_2Cl + Hg_2F_2 \longrightarrow 2CH_3CH_2F + Hg_2Cl_2$$
 (1)

(iii)
$$CH_3Br + 2Na + CH_3Br \xrightarrow{Dry} CH_3 - CH_3$$
 (1)
ether

