

 OSDAV Public School, Kaithal Half yearly Exams (2024-25) Class : XI Subject : CHEMISTRY

SET- A

Time: 3 Hrs. M.M. : 70 **General Instructions:-**

All questions are compulsory.

(a) There are 33 questions in this question paper.

(b) SECTION A consists of 16 multiple -choice questions carrying 1 mark each.

(c) SECTION B consists of 5 short answer questions carrying 2 marks each.

(d) SECTION C consists of 7 short answer questions carrying 3 marks each.

(e) SECTION D consists of 2 case - based questions carrying 4 marks.

(f) SECTION E consists of 3 long answer questions carrying 5 marks.

(g) Use of log tables and calculators is not allowed

OSDAV PUBLIC SCHOOL. KAITHAL

Answer Key for Mid Term Chemistry

Class-XI (2024-25)

SET-A

SECTION A

1. a. 2.0 mol of H2SO⁴

2. d. 2

3. a. Na>Mg<Al<Si

4. b. BF³

5. a. Trigonal bipyramidal

6. d. KMnO⁴

7. d. F

8. a. 1

9. d. 1

10. c. NH³

11. c. 1.806×10^{23}

12. b. loses electron to the oxidising agent

13. a. Both A and R are true and R is the correct explanation of A.

14. a. Both A and R are true and R is the correct explanation of A.

15. b. Both A and R are true but R is not the correct explanation of A.

16. a. Both A and R are true and R is the correct explanation of A.

SECTION B

17. a. Before the reaction, the metal centre could be described as $sp²$ hybridized, and trigonal planar in structure. Much of the chemistry of aluminium can be described in terms of the metal filling its vacant p orbital to give a $sp³$ description.

b. Axial bonds are not stronger than equatorial bonds. In fact, axial bonds are generally weaker because they experience more repulsion from nearby atoms, making them longer and less stable compared to equatorial bonds.

18. he ratio of their velocities is:

 $V_{CH4}/V_{O2} = m_{CH4}/m_{O2}$

Now, using the molar masses:

- Molar mass of CH₄= 16 g/mol,
- Molar mass of O2= 32 g/mol.

Taking the ratio of their velocities:

 $32/16=2$

Therefore, the velocity of methane molecule is **twice** the velocity of oxygen molecules for them to have the same de Broglie wavelength.

19. a. The bond angle in H₂O is smaller than in NH₃ because H₂O has two lone pairs on oxygen, while NH₃ has only one lone pair on nitrogen. Lone pairs repel more strongly than bonding pairs, and the additional lone pair in H2O causes greater repulsion, pushing the hydrogen atoms closer together and reducing the bond angle to 104.5°. In contrast, NH₃ experiences less repulsion with only one lone pair, resulting in a slightly larger bond angle of 107°.

 b. Atomic orbitals must have similar energy, proper symmetry, and significant overlap for effective combination into molecular orbitals.

20. **Step 1. First Write the Given Redox Reaction.** $MnO_a + F$ ------- $MnO₂ + I₂$ Step2. Identify Oxidation and Reduction half Reaction. MnO_a ------ $MnO₂$ [Reduction] Γ -------- I_2 [Oxidation] Step3. Balance the atoms undergoing change in the Oxidation number. \therefore MnO₄ -------- MnO₂ [Change of 4 units]. $2r$ ---------- I_2 [Change of 2 units]

Step 4. Multiply 1st equation by 1 and second equation by 2.

 \therefore MnO_a -------- MnO₂ and 4⁻ ---------- 2l₂

Step 5. Now, Add both the equations,

: $4r + MnO_4$ ---------- $I_2 + MnO_2$

Step 6. Now, Balance the charges by adding water and Hydrogen ions.

: $4f + MnO₄ + 2H₂O$ -----------1₂ + MnO₂ + 4OH

21. Magnesium's first ionization energy is higher than sodium's because its outer electron is more tightly bound due to a greater nuclear charge. However, sodium's second ionization energy is much higher than magnesium's because, after losing one electron, sodium attains a stable noble gas configuration. Removing an electron from this stable core requires significantly more energy than removing magnesium's second outer electron.

SECTION- C

22.

23.a. The second electron affinity of oxygen is expected to be positive (or less negative) compared to the first. The first electron affinity is negative because energy is released when an electron is added to a neutral oxygen atom, forming $O₋$. In contrast, the second electron affinity involves adding another electron to the negatively charged $O₋$ ion, which requires energy to overcome the repulsion between the two negatively charged species, making the process endothermic.

b. Fluorine (F) has a less negative electron gain enthalpy than chlorine (Cl), even though fluorine is more electronegative. This is because fluorine's small atomic size results in significant electronelectron repulsion in its compact 2p orbital. When an electron is added to fluorine, the high electron density causes more repulsion, making the process less energetically favourable. On the other hand, chlorine has a larger atomic size and a 3p orbital, where the added electron experiences less repulsion.

c. Period- 3, Group- 15

24.a.

The energy (E) associated with 300 nanometre photon is given by:

$$
=\frac{(6.626\times10^{-34})(3.0\times10^8ms^{-1})}{300\times10^{-9}}
$$

= 6.62×10⁻⁹J

 $E = \frac{hc}{\lambda}$

$$
E = \frac{hc}{\lambda}
$$

=
$$
\frac{(6.626 \times 10^{-34})(3.0 \times 10^8 m s^{-1})}{300 \times 10^{-9}}
$$

=
$$
6.62 \times 10^{-9} J
$$

Now, we will find energy of one mole of photons.

$$
=(6.626\times 10^{-19}J)\times (6.022\times 10^{23}mol^{-1})\\=3.9\times 10^{5}Jmol^{-1}
$$

Now, we will find minimum energy needed to remove a mole of electrons from sodium

 $=3.9\times10^{5}Jmol^{-1}-1.68\times10^{5}Jmol^{-1}$ $=(3.99-1.68)\times10^5 J mol^{-1}$ $= 2.31 \times 10^5 J$ mol⁻¹

We will find the minimum energy for one mole of electron

$$
=\frac{2.31\times10^{5}Jmol^{-1}}{6.022\times10^{23}mol^{-1}}\\=3.84\times10^{-19}J
$$

Now, by using this we will find the wavelength.

$$
\begin{aligned} \lambda &= \frac{hc}{E} \\ &= \frac{(6.626 \times 10^{-34})(3.0 \times 10^8 m s^{-1})}{3.84 \times 10^{-19} J} \\ &= 517 nm \end{aligned}
$$

b. Pauli's Exclusion Principle states that no two electrons in an atom can have identical sets of four quantum numbers, meaning each electron must occupy a unique state, allowing a maximum of two electrons per orbital with opposite spins.

 $25. a.$

100g solution contains $10gNaCl$ $w = 10g, m_{NaCl} = 58.5,$ Volume of solution $=$ $\frac{100}{1.071 \times 1000}$ litre $w_{H_2}o = 100 - 10 = 90g$ $\text{Molarity} = \frac{\text{Wt.of} \text{solute}}{\text{mol.wt.of} \text{solve} \times V_{\text{in}L}}$ $=\frac{10\times1.071\times1000}{58.5\times100}=1.83M$ Wt.of solute $\text{Molality} = \frac{\text{wt}.\text{of solute}}{\text{mol}.\text{wt}.\text{of solute} \times \text{weight of solvent in kg}}$ $=\frac{10\times1000}{58.5\times90}=1.90m$

b. Gay-Lussac's Law of Combining Volumes states that when gases react together at constant temperature and pressure, the volumes of the gaseous reactants and products can be expressed in simple whole number ratios. For example, if two gases react to form a gas, the volume of the product gas relative to the volumes of the reactants will be in a simple ratio, such as 1:1, 2:1, or 3:2, depending on the specific reaction.

26. a. Water (H₂O) is a liquid at room temperature due to strong hydrogen bonding between its molecules, which results from its bent shape and high polarity. In contrast, hydrogen sulfide (H₂S) has weaker dipole-dipole interactions and London dispersion forces, leading to lower boiling and melting points, causing it to exist as a gas at room temperature.

b. BeH₂ has a zero dipole moment because its linear molecular geometry allows the polar Be-H bonds to cancel each other out. Although each Be-H bond is polar due to the difference in electronegativity between beryllium and hydrogen, the symmetry of the linear structure means the dipoles are equal in magnitude and opposite in direction, resulting in no overall dipole moment for the molecule.

c. NaCl does not conduct electricity in the solid state because its ions are held tightly in a rigid crystal lattice, preventing them from moving freely. Electrical conductivity requires the movement of charged particles, which only occurs when NaCl is dissolved in water or melted. In these states, the ions are free to move, allowing the substance to conduct electricity.

27. a.

1 mol (55 + 2 \times 16 = 87 g) of MnO2 reacts completely with 4 mol (4 \times 36.5 = 146 g) of HCl.

Now, 87 g of MnO2 will react with 146 g of HCl.

So, 5.0 g of MnO2 will react with (5 X 146)/87 g of HCl.

5.0 g of MnO2 will react with 8.3 g of HCl.

Therefore, 8.4 g of HCI will react entirely with 5 g of manganese dioxide.

b. The limiting reagent in a chemical reaction is the reactant that is consumed first, thus determining the maximum amount of product that can be formed. Once the limiting reagent is used up, the reaction cannot proceed any further, even if other reactants are still available.

28. a. In quantum mechanics, L represents the principal quantum number, indicating the main energy level of an electron, while l denotes the azimuthal quantum number, which defines the shape of the atomic orbital. L indicates energy levels, and l specifies orbital shapes.

b. De Broglie's principle states that every moving particle or object has an associated wave-like nature. This means that particles such as electrons exhibit both particle and wave characteristics, leading to the concept of wave-particle duality in quantum mechanics.

c. Degenerate orbitals are orbitals that have the same energy level. In an atom, multiple orbitals within a given subshell are degenerate, meaning they are equal in energy. Degeneracy occurs in systems where the electrons experience the same potential energy, allowing them to occupy any of these orbitals without preference.

SECTION-D

29. a. $(n-1)d^{1-10}$ ns⁰⁻²

b. Ionization energy is the energy required to remove an electron from an atom or ion in its gaseous state, forming a positively charged ion. It reflects the atom's tendency to lose an electron and varies across the periodic table.

c. 1) II

2) I

30. a.

Number of spectral lines = $\frac{n(n-1)}{2}$ $=\frac{6(6-1)}{2}$ \mathfrak{D} $=15$

b. Emission spectra are the wavelengths of light emitted by atoms or molecules when electrons transition from higher to lower energy states. These spectra can be continuous or consist of distinct lines, allowing for the identification of elements based on their unique spectral signatures.

c.

$$
\frac{1}{\lambda}=R\Bigg[\frac{1}{n_1^2}-\frac{1}{n_2^2}\Bigg]
$$

 λ is the wavelength

R is the Rydberg Constant which is 1.097×10^{-7} m⁻¹

n₁ is the lower energy level

n₂ is the higher energy level

Putting in the numbers:

$$
\frac{1}{\lambda} = 1.097 \times 10^7 \bigg[\frac{1}{2^2} - \frac{1}{5^2} \bigg]
$$

 $\frac{1}{\lambda} = 1.097 \times 10^7 [0.25 - 0.04] = 0.2304 \times 10^7 \text{ m}^{-1}$

: $\lambda = 4.34 \times 10^{-7}$ m

 $\lambda = 434$ nm

SECTION-E

$31. a.$

Assuming 1 gm. Of gas is burnt The weight % of $C = \frac{3.38 \times 12}{44} \times 100 - 92.18\%$ The weight % of .H. = $\frac{0.69 \times 2}{18}$ = 7.67 % **ISSA AT A WAY CARD ARE THE**

Experical formula of compound = C_1H_1

(ii) Given 10 lit of gas at STP weighs - 11.6 gas

22.44 lit of gas at STP weights = $\frac{22.4 \times 11.6}{10}$ = 25.984

 \therefore Molecular weight of given gas = 25.984

(iii) Molecular formula = n (emperical formula) $n = \frac{\text{Mol wt}}{\text{Emp.wt}} = \frac{25.984}{13} = 2$:. Molecular formula = 2(*CH*) = C_2H_2 .

b.

In the conversion, the 0.94 is multiplied with 2 because the oxygen is diatomic.

Hence, the formula is $Fe₂O₃$.

32. a.

Electronic configuration:

 $\sigma 1s^2$ < σ^*1s^2 < $\sigma 2s^2$ < σ^*2s^2 , $[\pi 2p_x^2 = \pi 2p_y^2]$ < $\sigma 2p_z^2$ < $[\pi^*2p_x = \pi^*2p_y]$ < σ^*2p_z

Let's calculate the bond order of $N_{\overline{2}};$ Bond order $=$ $\frac{Bonding electrons - Anti bonding electrons}{2}$ $=\frac{10-4}{2}$ = 3

 $N₂$ does not have unpaired electrons, hence it is diamagnetic.

 $_b$.</sub>

 $33.a.$

 $\Delta x = 5.8 \times 10^{-4}$ m

```
Speed = v = 500 m/s.
Uncertainty in speed is 0.02 %
\Delta v = 0.02 % of 500 m/s
= 0.02100 \times 500 = 0.1 m/s
According to Heisenberg uncertainty principle,
\Delta x * \Delta p \ge h / 4\piHere \Delta x = uncertainty in position
\Delta p = uncertainty in momentum
h = Planck's constant = 6.63 \times 10^{-34} kg m<sup>2</sup>/s
\Delta p = m^* \Delta vm = mass of electron = 9.1 \times 10^{-31} kg
\Delta v = uncertainty in speed = 0.1 m/s
\Delta p = m \Delta v = 9.1 \times 10^{-31} kg × 0.1 m/s = 9.1×10<sup>-32</sup> kg m/s
\Delta x^* \Delta p \ge h / 4\piPlug all the values;
\Delta x \ge h / 4\pi * \Delta p
```
If $l = 0$ $m = 0$. This is an s orbital If $l = 1$, $m = -1$, 0, +1. This gives the three p orbitals. So $m = 0$ is ok. If $l = 2$ $m = -2, -1, 0, 1, 2$. This gives the five d orbitals. c.

$F^{-1}1s^22s^22p^6$

d. Hund's Rule of Maximum Multiplicity states that for degenerate orbitals (orbitals of the same energy), electrons will occupy these orbitals singly before pairing up. This means that each orbital within a given subshell receives one electron with the same spin direction before any orbital gets a second electron. This arrangement minimizes electron-electron repulsion and maximizes the total spin, resulting in a more stable electron configuration.

b.

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Answer Key for Mid Term Chemistry

Class-XI (2024-25)

SET-B

(only different questions)

SECTION-A

- 1. c. 0.0177
- 2. d. 2
- 3. c. Si,P,C,N
- $4. b. BF₃$
- 5. c. Octahedral
- 6.d. KMnO⁴
- 7. a. Na
- 8. a. 4s orbital
- 9. b. 1
- 10. b. Loses electron to the oxidising agent
- 11. c. 1.806×10^{23}
- 12. c. NH³
- 13. b.
- 14. a.
- 15.a.

16.a.

SECTION- B

17.b. Lone pairs of electrons distort the geometry of molecules because they exert greater repulsive forces on bonded electron pairs. Unlike bonding pairs, lone pairs are located closer to the nucleus, leading to stronger repulsion. For example, in water (H_2O) , the oxygen atom has two lone pairs and two bonding pairs of electrons. According to the ideal tetrahedral arrangement, the bond angle should be 109.5°, but the lone pairs push the hydrogen atoms

closer together, reducing the bond angle to about 104.5°. This distortion results in a bent molecular shape rather than a perfect tetrahedral.

a.

Boron atom in BF₃ is sp² hybridized. The orbital picture of boron in the excited state can

be shown as:

Nitrogen atom in NH₃ is sp³ hybridized. The orbital picture of nitrogen can be represented as:

 $\frac{1}{2p_x} \frac{1}{2p_y} \frac{1}{2p_z}$ $\left| \uparrow \downarrow \right|$

After the reaction has occurred, an adduct F_3B . NH₃ is formed as hybridization of 'B' changes to sp³. However, the hybridization of 'N' remains intact.

18.

For the electron, the de Broglie wavelength is: $\lambda_E = \frac{h}{m_e v_e}$ For the proton, the de Broglie wavelength is: $\lambda_P = \frac{h}{m_p v_p}$

Step 3: Set the wavelengths equal to each other To find when the de Broglie wavelengths are equal, we set them equal: $\lambda_F - \lambda_P$ This gives us: $\frac{\hbar}{m_e v_e} = \frac{\hbar}{m_p v_p}$

Step 4: Cancel Planck's constant

Since h is common on both sides, we can cancel it out: $\frac{1}{m_{e}v_{e}}=\frac{1}{m_{p}v_{p}}% \frac{1}{m_{p}v_{p}}\sum_{i=1}^{p}\left\vert \frac{p_{e}^{2}}{p_{e}^{2}}\right\vert ^{2} \label{v1}%$

Step 5: Rearrange the equation Rearranging gives us:

 $m_p v_p = m_e v_e$

Step 6: Substitute the known masses Substituting the known values of the masses:

- Mass of electron $m_{\rm e} = 9.1095 \times 10^{-31} \, \rm kg$

- Mass of proton $m_p=1.6725\times 10^{-27}\,\mathrm{kg}$

This gives us: 1.6725×10^{-27} $v_p = 9.1095 \times 10^{-31}$ v_e

Step 7: Express v_c in terms of v_p

Rearranging this equation to express v_e in terms of v_p :

 $v_c = \frac{1.6725 \times 10^{-27}}{1}$ $9.1095\times10^{-31}\,v_p$

Step 8: Calculate the ratio of the masses

Calculating the ratio: 1.6725×10^{-27} $\overline{9.1095\times 10^{-31}}\approx 1836$ Thus, we have: $v_e = 1836 v_p$

19. a. Both NH $_3$ (ammonia) and H $_2$ O (water) have distorted tetrahedral geometries due to lone pairs of electrons, but the bond angle of water is smaller than that of ammonia. In NH₃, nitrogen has one lone pair and three bonding pairs, creating a bond angle of around 107°. In H₂O, oxygen has two lone pairs and two bonding pairs, resulting in a bond angle of about 104.5°. The reason for the smaller angle in water is that lone pairs repel more strongly than bonding pairs, and in water, the two lone pairs exert even greater repulsion, compressing the bond angle more than in ammonia, which has only one lone pair.

20. $SO_{2(g)} + 2H_2O_{(1)} \rightarrow HSO_{4(aq)}^- + 3H_{(aq)}^+ + 2e_{(aq)}^-$ And the reduction half reaction as: $MnO_{4(aq)}⁺ + 8H_(aq)⁺ + 5e⁻ \rightarrow Mn_(aa)²⁺ + 4H₂O₍₁₎$ Multiplying the oxidation half reaction by 5 and the reduction half reaction by 2, and then by adding them, we have the net balanced redox reaction as: $2MnO_{4(aq)}⁺ 5SO_{s(g)} + 2H₂O₍₁₎ + H_(aq)⁺ \rightarrow 2Mn_(aq)²⁺ + 5HSO_{4(aq)}$

21. The first ionization energy of isotopes is generally the same. This is because ionization energy primarily depends on the number of protons and the arrangement of electrons in an atom, which remain the same for isotopes of an element. Isotopes differ only in the number of neutrons, and since neutrons are neutral particles, they do not significantly affect the attraction between the nucleus and the electrons. Consequently, isotopes of the same element have nearly identical ionization energies.

b. Modern periodic law states that the physical and chemical properties of elements are periodic functions of their atomic numbers. This means that when elements are arranged in order of increasing atomic number (the number of protons in an atom's nucleus), elements with similar properties recur at regular intervals or periods. The atomic number, rather than atomic mass as in Mendeleev's periodic law, is the key determinant in organizing the periodic table, providing a more accurate reflection of element properties.

SECTION- C

23. c. Period- 3

Group- 2

26. a. Ionic bonds form due to the electrostatic attraction between positively and negatively charged ions. This attraction occurs equally in all directions around the ions, meaning the strength of the bond is the same in every direction, making ionic bonds non-directional.

In contrast, covalent bonds involve the sharing of electrons between atoms, and this sharing happens along specific axes between the bonded atoms. The electron density is concentrated between the atoms, resulting in a specific spatial orientation, making covalent bonds directional in nature.

b. KHF₂ exists, but KHCl₂ does not, due to the unique ability of fluoride ions (F^-) to form stable hydrogen bonds, which chloride ions $(Cl⁻)$ cannot. In potassium bifluoride $(KHF₂)$, the fluoride ion forms a strong hydrogen bond with the hydrogen atom, creating a $[HF_2]$ ⁻ ion.

This structure is stabilized because fluoride is small, highly electronegative, and can easily form these strong hydrogen bonds.

Chloride ions, being larger and less electronegative than fluoride, cannot form similarly strong hydrogen bonds. As a result, a compound like KHCl₂, where $[HCl₂]⁻$ would need to form, is not stable and does not exist.

c. AlF₃ (aluminum fluoride) is a high-melting solid, while $SiF₄$ (silicon tetrafluoride) is a gas due to the nature of bonding in these compounds.

In AlF₃, the bond between aluminum and fluorine is highly ionic, meaning the attraction between the oppositely charged ions forms a strong lattice structure. This ionic lattice requires a significant amount of energy to break, resulting in a high melting point and solid state at room temperature.

On the other hand, SiF₄ has covalent bonds between silicon and fluorine, forming discrete molecular units. The intermolecular forces (van der Waals forces) between these SiF₄ molecules are weak, leading to a much lower boiling point, and making SiF4 a gas at room temperature. Thus, the difference in bonding (ionic in AlF_3 and covalent in SiF_4) explains their different physical states.

27.

$$
\begin{array}{r}\n\cdot \text{S} = C = S \\
\cdot \text{S} = 15\% \text{ (w/w)} + 15\
$$

Mass of solvent in kg 150×1000

$$
\Rightarrow \frac{150 \times 1000}{98 \times 850} = 1.80 \text{ m}
$$

SECTION-E

Bond Order- 2, magnetic nature- Paramagnetic

b.

