

AP EAPCET 2026 May 12 Shift 1

Question Paper (Memory-Based) with Solutions

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General Instructions

- (i) The test is of 3 hours duration.
- (ii) This test paper consists of 160 questions. The maximum marks are 160.
- (iii) Physics and Chemistry contains 40 questions each and Mathematics contains 80 questions.
- (iv) Each question carries +1 marks for correct answer and there is no negative marking for wrong answer.

1. Let $f(x) = \int \frac{\sqrt{x}}{(1+x)^2} dx (x \geq 0)$. Then $f(3) - f(1)$ is equal to:

- (A) $-\frac{\pi}{12} + \frac{1}{2} + \frac{\sqrt{3}}{4}$
- (B) $\frac{\pi}{12} + \frac{1}{2} - \frac{\sqrt{3}}{4}$
- (C) $-\frac{\pi}{6} + \frac{1}{2} + \frac{\sqrt{3}}{4}$
- (D) $\frac{\pi}{6} + \frac{1}{2} - \frac{\sqrt{3}}{4}$

Correct Answer: (B) $\frac{\pi}{12} + \frac{1}{2} - \frac{\sqrt{3}}{4}$

Solution: Concept: The expression $f(3) - f(1)$ represents the definite integral of the function from 1 to 3. Specifically, by the Fundamental Theorem of Calculus:

$$f(3) - f(1) = \int_1^3 \frac{\sqrt{x}}{(1+x)^2} dx$$

To solve this, we use a trigonometric substitution to simplify the radical and the denominator. Since we have a term like $(1+x)$, substituting $x = \tan^2 \theta$ is effective because $1 + \tan^2 \theta = \sec^2 \theta$.

Step 1: Substitution and changing limits. Let $x = \tan^2 \theta$. Then:

$$dx = 2 \tan \theta \sec^2 \theta d\theta$$

Now, let's find the new limits for θ :

- When $x = 1$, $\tan^2 \theta = 1 \Rightarrow \tan \theta = 1 \Rightarrow \theta = \frac{\pi}{4}$.
- When $x = 3$, $\tan^2 \theta = 3 \Rightarrow \tan \theta = \sqrt{3} \Rightarrow \theta = \frac{\pi}{3}$.

Step 2: Transforming the integral. Substitute the values into the integral:

$$I = \int_{\pi/4}^{\pi/3} \frac{\sqrt{\tan^2 \theta}}{(1 + \tan^2 \theta)^2} \cdot (2 \tan \theta \sec^2 \theta) d\theta$$

Using the identity $1 + \tan^2 \theta = \sec^2 \theta$:

$$I = \int_{\pi/4}^{\pi/3} \frac{\tan \theta}{(\sec^2 \theta)^2} \cdot 2 \tan \theta \sec^2 \theta d\theta$$

$$I = \int_{\pi/4}^{\pi/3} \frac{2 \tan^2 \theta \sec^2 \theta}{\sec^4 \theta} d\theta = 2 \int_{\pi/4}^{\pi/3} \frac{\tan^2 \theta}{\sec^2 \theta} d\theta$$

Convert to sine and cosine:

$$I = 2 \int_{\pi/4}^{\pi/3} \frac{\sin^2 \theta / \cos^2 \theta}{1 / \cos^2 \theta} d\theta = 2 \int_{\pi/4}^{\pi/3} \sin^2 \theta d\theta$$

Step 3: Integrating and applying limits. Use the power-reduction identity $\sin^2 \theta = \frac{1 - \cos 2\theta}{2}$:

$$I = 2 \int_{\pi/4}^{\pi/3} \frac{1 - \cos 2\theta}{2} d\theta = \int_{\pi/4}^{\pi/3} (1 - \cos 2\theta) d\theta$$

$$I = \left[\theta - \frac{\sin 2\theta}{2} \right]_{\pi/4}^{\pi/3}$$

Evaluate at the upper limit ($\pi/3$): $\frac{\pi}{3} - \frac{\sin(2\pi/3)}{2} = \frac{\pi}{3} - \frac{\sqrt{3}/2}{2} = \frac{\pi}{3} - \frac{\sqrt{3}}{4}$.

Evaluate at the lower limit ($\pi/4$): $\frac{\pi}{4} - \frac{\sin(2\pi/4)}{2} = \frac{\pi}{4} - \frac{\sin(\pi/2)}{2} = \frac{\pi}{4} - \frac{1}{2}$.

Subtracting the two:

$$I = \left(\frac{\pi}{3} - \frac{\sqrt{3}}{4} \right) - \left(\frac{\pi}{4} - \frac{1}{2} \right)$$

$$I = \frac{\pi}{3} - \frac{\pi}{4} + \frac{1}{2} - \frac{\sqrt{3}}{4} = \frac{\pi}{12} + \frac{1}{2} - \frac{\sqrt{3}}{4}$$

Quick Tip: When you see \sqrt{x} and $(1+x)$ in the same integral, the substitution $x = \tan^2 \theta$ is usually the fastest way to collapse the denominator using Pythagorean identities.

2. Gas is being pumped into a spherical balloon at the rate of $30 \text{ ft}^3/\text{min}$. Then the rate at which the radius increases when it reaches the value 15 ft is:

- (A) $\frac{1}{30\pi}$ ft/min
- (B) $\frac{1}{15\pi}$ ft/min
- (C) $\frac{1}{20\pi}$ ft/min
- (D) $\frac{1}{25\pi}$ ft/min

Correct Answer: (A) $\frac{1}{30\pi}$ ft/min

Solution:

Concept: This is a related rates problem. We are given the rate of change of volume (dV/dt) and need to find the rate of change of the radius (dr/dt). The volume V of a sphere is given by:

$$V = \frac{4}{3}\pi r^3$$

Step 1: Differentiating the volume formula. We differentiate both sides of the volume equation with respect to time t using the chain rule:

$$\frac{dV}{dt} = \frac{4}{3}\pi \cdot \frac{d}{dt}(r^3)$$

$$\frac{dV}{dt} = \frac{4}{3}\pi \cdot 3r^2 \frac{dr}{dt}$$

$$\frac{dV}{dt} = 4\pi r^2 \frac{dr}{dt}$$

Step 2: Plugging in given values.

We are given:

- $\frac{dV}{dt} = 30 \text{ ft}^3/\text{min}$
- $r = 15 \text{ ft}$

Substitute these into our differentiated equation:

$$30 = 4\pi(15)^2 \frac{dr}{dt}$$

$$30 = 4\pi(225) \frac{dr}{dt}$$

$$30 = 900\pi \frac{dr}{dt}$$

Step 3: Solving for dr/dt .

$$\frac{dr}{dt} = \frac{30}{900\pi}$$

Simplify the fraction by dividing both numerator and denominator by 30:

$$\frac{dr}{dt} = \frac{1}{30\pi} \text{ ft/min}$$

Quick Tip: In related rates, always identify your "knowns" and your "unknowns" before differentiating. Remember that $\frac{dV}{dt} = (\text{Surface Area}) \times \frac{dr}{dt}$ for a sphere.

3. Let the function $f(x)$ be defined as $f(x) = \frac{x-|x|}{x}$, then:

- (A) the function is continuous everywhere
- (B) the function is not continuous
- (C) the function is continuous when $x < 0$
- (D) the function is continuous for all x except zero

Correct Answer: (D) the function is continuous for all x except zero

Solution:

Concept: A function is continuous where it is defined and the limit equals the function value. First, we must redefine $f(x)$ by removing the absolute value sign based on the domain of x .

Step 1: Redefining the function for different intervals. Recall the definition of $|x|$:

$$|x| = \begin{cases} x & \text{if } x > 0 \\ -x & \text{if } x < 0 \end{cases}$$

Note that at $x = 0$, the denominator of $f(x)$ is zero, so the function is undefined at $x = 0$.

- For $x > 0$: $f(x) = \frac{x-x}{x} = \frac{0}{x} = 0$.
- For $x < 0$: $f(x) = \frac{x-(-x)}{x} = \frac{x+x}{x} = \frac{2x}{x} = 2$.

Step 2: Analyzing continuity.

The function is a constant 0 for all $x \in (0, \infty)$, which is continuous. The function is a constant 2 for all $x \in (-\infty, 0)$, which is also continuous. However, at $x = 0$, the function is not defined. Furthermore, the left-hand limit (2) is not equal to the right-hand limit (0), showing a jump discontinuity.

Step 3: Conclusion.

Since the function is defined and constant in the intervals $(-\infty, 0)$ and $(0, \infty)$, it is continuous on those intervals. The only point of concern is $x = 0$, where the function fails to exist. Thus, the function is continuous for all x except zero.

Quick Tip: Always check the domain first. If a function's denominator is zero at a point, it cannot be continuous there, regardless of whether the limit exists.

4. If the direction ratios of two lines are given by $l + m + n = 0$ and $mn - 2ln + lm = 0$, then the angle between the lines is:

- (A) $\frac{\pi}{4}$
- (B) $\frac{\pi}{3}$
- (C) $\frac{\pi}{2}$
- (D) 0

Correct Answer: (C) $\frac{\pi}{2}$

Solution:

Concept: The angle θ between two lines with direction ratios (l_1, m_1, n_1) and (l_2, m_2, n_2) is found using:

$$\cos \theta = \frac{|l_1 l_2 + m_1 m_2 + n_1 n_2|}{\sqrt{l_1^2 + m_1^2 + n_1^2} \sqrt{l_2^2 + m_2^2 + n_2^2}}$$

If the dot product $l_1 l_2 + m_1 m_2 + n_1 n_2 = 0$, the lines are perpendicular ($\theta = \pi/2$).

Step 1: Substituting one variable.

From $l + m + n = 0$, we have $n = -(l + m)$. Substitute this into the second equation $mn - 2ln + lm = 0$:

$$m(-(l + m)) - 2l(-(l + m)) + lm = 0$$

$$-ml - m^2 + 2l^2 + 2lm + lm = 0$$

$$2l^2 + 2lm - m^2 = 0$$

Step 2: Forming a quadratic in (l/m) .

Divide the equation by m^2 :

$$2\left(\frac{l}{m}\right)^2 + 2\left(\frac{l}{m}\right) - 1 = 0$$

Let the two roots (direction ratios of the two lines) be $\frac{l_1}{m_1}$ and $\frac{l_2}{m_2}$.

From the properties of quadratic equations:

Product of roots: $\frac{l_1 l_2}{m_1 m_2} = \frac{c}{a} = \frac{-1}{2} \Rightarrow 2l_1 l_2 = -m_1 m_2 \Rightarrow 2l_1 l_2 + m_1 m_2 = 0$.

Step 3: Finding the final condition.

We also have $n_1 = -(l_1 + m_1)$ and $n_2 = -(l_2 + m_2)$.

Product $n_1 n_2 = (l_1 + m_1)(l_2 + m_2) = l_1 l_2 + l_1 m_2 + l_2 m_1 + m_1 m_2$.

Using the sum of roots $\frac{l_1}{m_1} + \frac{l_2}{m_2} = -\frac{b}{a} = -1$, we get $l_1 m_2 + l_2 m_1 = -m_1 m_2$.

Substitute this back: $n_1 n_2 = l_1 l_2 - m_1 m_2 + m_1 m_2 = l_1 l_2$.

Now calculate $l_1 l_2 + m_1 m_2 + n_1 n_2$:

Using $m_1 m_2 = -2l_1 l_2$:

$$l_1 l_2 + (-2l_1 l_2) + l_1 l_2 = 0$$

Since the sum of the products of the direction ratios is zero, the lines are perpendicular.

Quick Tip: In problems involving homogeneous equations in l, m, n , always aim to reduce the system to a quadratic in terms of a ratio like l/m or m/n .

5. The equation of the normal to the curve $y = \log_e x$ at the point $P(1, 0)$ is ____.

- (A) $2x + y = 2$
- (B) $x - 2y = 1$
- (C) $x - y = 1$
- (D) $x + y = 1$

Correct Answer: (D) $x + y = 1$

Solution:

Concept: The slope of the tangent to a curve $y = f(x)$ at point (x_1, y_1) is $m_t = f'(x_1)$. The slope of the normal is the negative reciprocal: $m_n = -1/m_t$. The equation of the line is then $y - y_1 = m_n(x - x_1)$.

Step 1: Finding the slope of the tangent.

The given curve is $y = \ln x$.

Differentiating with respect to x :

$$\frac{dy}{dx} = \frac{1}{x}$$

At the point $P(1, 0)$, the slope of the tangent m_t is:

$$m_t = \left. \frac{dy}{dx} \right|_{x=1} = \frac{1}{1} = 1$$

Step 2: Finding the slope of the normal.

Since the normal is perpendicular to the tangent:

$$m_n = -\frac{1}{m_t} = -\frac{1}{1} = -1$$

Step 3: Equation of the normal line.

Using the point-slope form with point $(1, 0)$ and slope $m_n = -1$:

$$y - 0 = -1(x - 1)$$

$$y = -x + 1$$

$$x + y = 1$$

Quick Tip: For $y = \ln x$, the tangent at $(1, 0)$ is always $y = x - 1$. Swapping the coefficients and changing the sign of one (perpendicular line property) gives the normal $x + y = k$.

6. The radius in the hydrogen atom in the ground state is 0.53 \AA . The radius of Li^{2+} ion (atomic number = 3) in a similar state is

- (a) 0.53 \AA
- (b) 1.06 \AA
- (c) 0.17 \AA
- (d) 0.265 \AA

Correct Answer: (c) 0.17 \AA

Solution:

Concept: According to Bohr's model of the atom, the radius of the n^{th} orbit for a hydrogen-like species (single-electron systems like H , He^+ , Li^{2+} , etc.) is given by the formula:

$$r_n = a_0 \frac{n^2}{Z}$$

Where:

- r_n is the radius of the n^{th} orbit.

- a_0 is the Bohr radius (radius of H-atom in ground state, $\approx 0.529\text{\AA}$ or 0.53\AA).
- n is the principal quantum number (orbit number).
- Z is the atomic number of the element.

Step 1: Identify the given values for Hydrogen and Lithium. For the Hydrogen atom in the ground state:

- Principal quantum number (n) = 1 (since it is the ground state).
- Atomic number (Z_H) = 1.
- Radius (r_H) = 0.53\AA .

For the Li^{+2} ion in a similar state:

- Principal quantum number (n) = 1 (as it is in a "similar state" i.e., ground state).
- Atomic number for Lithium (Z_{Li}) = 3.

Step 2: Establishing the relationship between radii. Using the formula $r \propto \frac{n^2}{Z}$, and since n is constant ($n = 1$) for both cases, we can state that:

$$r \propto \frac{1}{Z}$$

Therefore, the ratio of the radius of the Li^{+2} ion to the radius of the Hydrogen atom is:

$$\frac{r_{Li^{+2}}}{r_H} = \frac{Z_H}{Z_{Li}}$$

Step 3: Calculating the final radius.

Substitute the known values into the ratio:

$$\frac{r_{Li^{+2}}}{0.53} = \frac{1}{3}$$

Now, solve for $r_{Li^{+2}}$:

$$r_{Li^{+2}} = \frac{0.53}{3}$$

$$r_{Li^{+2}} = 0.17666... \text{Å}$$

Rounding to two decimal places as per the options provided:

$$r_{Li^{+2}} \approx 0.17 \text{Å}$$

Quick Tip: To quickly solve Bohr radius problems, remember that as the nuclear charge (Z) increases, the nucleus pulls the electron closer, making the radius smaller. Since Lithium has 3 protons ($Z = 3$) compared to Hydrogen's 1, its ground state radius must be exactly one-third of Hydrogen's.

7. The energy of the second bohr orbit of the hydrogen atom is -328 kJ mol^{-1} ; hence the energy of the fourth bohr orbit would be:

- (a) -41 kJ/mol
- (b) -82 kJ/mol
- (c) -164 kJ/mol
- (d) -1312 kJ/mol

Correct Answer: (b) -82 kJ/mol

Solution:

Concept: According to Bohr's theory of the hydrogen atom, the energy of an electron in the n^{th} orbit (E_n) is inversely proportional to the square of the principal quantum number (n). The mathematical expression for the energy of an orbit is:

$$E_n = -\frac{k \cdot Z^2}{n^2}$$

Where:

- E_n is the energy of the n^{th} orbit.
- Z is the atomic number (for Hydrogen, $Z = 1$).
- n is the principal quantum number (orbit number).
- k is a constant representing the energy of the ground state.

From this formula, we can derive a proportionality relationship for a specific element:

$$E_n \propto \frac{1}{n^2}$$

Step 1: Establishing the ratio between two orbits. Let E_2 be the energy of the second orbit ($n_1 = 2$) and E_4 be the energy of the fourth orbit ($n_2 = 4$). Based on the inverse-square relationship:

$$\frac{E_4}{E_2} = \frac{n_1^2}{n_2^2}$$

Substituting the orbit numbers:

$$\frac{E_4}{E_2} = \frac{2^2}{4^2} = \frac{4}{16} = \frac{1}{4}$$

Step 2: Substituting the given value. We are given that the energy of the second Bohr orbit (E_2) is -328 kJ mol^{-1} . Using the relationship derived in Step 1:

$$E_4 = E_2 \times \frac{1}{4}$$

$$E_4 = -328 \times \frac{1}{4}$$

Step 3: Final Calculation. Perform the division:

$$E_4 = -\frac{328}{4}$$

$$E_4 = -82 \text{ kJ mol}^{-1}$$

This calculation shows that as the electron moves further from the nucleus (to a higher orbit), the energy level increases (becomes less negative).

Quick Tip: To solve these quickly: if the orbit number doubles (from 2 to 4), the energy becomes $\frac{1}{2^2} = \frac{1}{4}$ of its previous value. Conversely, if the orbit number triples, the energy becomes $\frac{1}{9}$ of the original value. Always remember that energy levels get closer together as n increases.

8. The IUPAC name of an element with atomic number 119 is:

- (a) Ununennium
- (b) Unnilennium
- (c) Unununium
- (d) ununoctium

Correct Answer: (a) Ununennium

Solution:

Concept: For elements with atomic numbers greater than 100, IUPAC established a systematic nomenclature based directly on the digits of the atomic number.

This ensures a temporary name is available before a permanent name (usually honoring a scientist or place) is officially adopted.

The names are derived using numerical roots for each digit:

The name is constructed by putting the roots together in order of the digits and adding the

suffix "-ium" at the end.

Step 1: Breaking down the atomic number digits. The given atomic number is 119. We identify the roots for each individual digit:

- First digit: 1 → root is "un"
- Second digit: 1 → root is "un"
- Third digit: 9 → root is "enn"

Step 2: Assembling the roots. Combine the roots in the order of the digits:

un + un + enn

This gives us the base string: "ununenn".

Step 3: Applying the suffix and final naming. Add the mandatory IUPAC suffix "-ium" to the end of the root string:

ununenn + ium = Ununennium

The symbol for the element is derived from the first letter of each root (capitalizing the first one): Uue.

Note on spelling: If a root ends in 'enn' and the suffix is 'ium', we keep all three 'n's (enn-ium). If a root ends in 'bi' or 'tri', we drop one 'i' (e.g., bi-ium becomes bium).

Quick Tip: To never forget the roots: 0 is Nil (nothing), 1 is Un (uni-), 2 is Bi (bi-), and 9 is Enn (think of 'enneagon' for a 9-sided shape). Just string them together and add "-ium"!

9. Which of the following orders is not correct for the given property?

- (a) $\text{Li} < \text{Na} < \text{K}$ – metallic radius
- (b) $\text{Br} < \text{F} < \text{Cl}$ – electron gain enthalpy
- (c) $\text{C} < \text{N} < \text{O}$ – first ionisation enthalpy
- (d) $\text{Mg}^{+2} < \text{Na}^+ < \text{F}^-$ – ionic radius

Correct Answer: (c) $\text{C} < \text{N} < \text{O}$ – first ionisation enthalpy

Solution: Concept: Periodic properties follow specific trends across periods and down groups. However, there are significant exceptions caused by electronic configurations (half-filled and fully-filled stability) and electron-electron repulsions.

- **Atomic/Metallic Radius:** Increases down a group due to the addition of new shells.
- **Electron Gain Enthalpy:** Generally becomes more negative across a period. For halogens, $\text{Cl} > \text{F}$ due to inter-electronic repulsion in the small F atom.
- **Ionisation Enthalpy:** Generally increases across a period but decreases if an atom has a stable configuration.
- **Ionic Radius:** In isoelectronic species, radius decreases as nuclear charge (Z) increases.

Step 1: Evaluating Option (a): Metallic Radius Li, Na, and K are Alkali metals (Group 1). As we move down the group from Lithium to Potassium, the number of energy shells increases ($n = 2$ for Li, $n = 3$ for Na, $n = 4$ for K). This outweighs the increase in nuclear charge, resulting in an increase in metallic radius. Order: $\text{Li} < \text{Na} < \text{K}$. (Correct)

Step 2: Evaluating Option (b): Electron Gain Enthalpy Usually, Electron Gain Enthalpy decreases down a group.

However, Fluorine (F) has a very small size. When an electron is added to its $2p$ orbital, it faces significant inter-electronic repulsion. Chlorine (Cl) is larger, allowing the electron to be added more easily with a higher release of energy. Order of magnitude: $\text{Br} < \text{F} < \text{Cl}$. (Correct)

Step 3: Evaluating Option (c): First Ionisation Enthalpy Across the second period (C, N, O), ionisation enthalpy generally increases. Let's look at the electronic configurations:

- $\text{C}(Z = 6) : 1s^2 2s^2 2p^2$
- $\text{N}(Z = 7) : 1s^2 2s^2 2p^3$ (Exactly half-filled p -subshell)
- $\text{O}(Z = 8) : 1s^2 2s^2 2p^4$

Nitrogen has a half-filled $2p$ subshell, which is exceptionally stable. Removing an electron from Nitrogen requires more energy than removing one from Oxygen, where electron-electron repulsion in the paired $2p$ orbital makes removal easier. Correct Order: $\text{C} < \text{O} < \text{N}$. The given order $\text{C} < \text{N} < \text{O}$ is Incorrect.

Step 4: Evaluating Option (d): Ionic Radius Mg^{+2} , Na^+ , and F^- all have 10 electrons (isoelectronic).

- Mg^{+2} : $Z = 12$ (Strongest nuclear pull)

- Na^+ : $Z = 11$
- F^- : $Z = 9$ (Weakest nuclear pull)

As Z increases, the radius decreases. Order: $\text{Mg}^{+2} < \text{Na}^+ < \text{F}^-$. (Correct)

Quick Tip: Always watch out for the "N vs O" and "Be vs B" traps in Ionisation Enthalpy! Half-filled and full-filled configurations jump ahead of the next element in the period.

10. Identify the correct set of molecules with zero dipole moment:

- (a) $\text{CO}_2, \text{NH}_3, \text{H}_2\text{O}$
- (b) $\text{NH}_3, \text{NF}_3, \text{BF}_3$
- (c) $\text{PF}_3, \text{NH}_3, \text{CH}_4$
- (d) $\text{CH}_4, \text{BF}_3, \text{CO}_2$

Correct Answer: (d) $\text{CH}_4, \text{BF}_3, \text{CO}_2$

Solution:

Concept: The dipole moment (μ) of a molecule depends on both the polarity of the individual bonds and the geometric shape (symmetry) of the molecule. A molecule has a zero dipole moment if it is highly symmetrical, such that the individual bond dipoles cancel each other out as vectors.

- **Polar Molecules:** Contain lone pairs on the central atom or have asymmetrical surrounding atoms (e.g., $\text{NH}_3, \text{H}_2\text{O}, \text{NF}_3$).
- **Non-Polar Molecules:** Symmetrical shapes where bond dipoles cancel (e.g., Linear, Trigonal Planar, Tetrahedral with identical atoms).

Step 1: Analyzing molecules in the options. Let's examine the symmetry and dipole cancellation for each unique molecule mentioned:

- **CO_2 :** It has a linear geometry ($\text{O}=\text{C}=\text{O}$). The two $\text{C}=\text{O}$ bond dipoles are equal in magnitude but opposite in direction. They cancel out perfectly. $\mu = 0$.
- **BF_3 :** It has a trigonal planar geometry. The three $\text{B}-\text{F}$ bond dipoles are oriented at 120° to each other. Their vector sum is zero. $\mu = 0$.

- **CH₄**: It has a tetrahedral geometry. All four C-H bonds are identical and symmetrically arranged. The vector sum of these four dipoles is zero. $\mu = 0$.

Step 2: Identifying molecules with non-zero dipole moments. Now, let's see why the other sets are incorrect:

- **NH₃ and NF₃**: These have pyramidal geometry due to the presence of a lone pair on Nitrogen. The lone pair contributes to the dipole and prevents the bond dipoles from canceling. $\mu \neq 0$.
- **H₂O**: It has a bent (V-shape) geometry because of two lone pairs on Oxygen. The dipoles do not cancel. $\mu \neq 0$.
- **PF₃**: Similar to NH₃, it is pyramidal with a lone pair. $\mu \neq 0$.

Step 3: Final Selection.

Comparing our findings with the options:

- (a) Contains NH₃ and H₂O ($\mu \neq 0$).
- (b) Contains NH₃ and NF₃ ($\mu \neq 0$).
- (c) Contains PF₃ and NH₃ ($\mu \neq 0$).
- (d) Contains CH₄, BF₃, and CO₂. All three are symmetrical and have zero dipole moment.

Quick Tip: A quick shortcut: If the central atom has no lone pairs and all the surrounding atoms are the same, the molecule will almost always have a zero dipole moment due to its perfect symmetry!

11. The potential energy of a simple harmonic oscillator when the particle is half way to its end point is (where E is the total energy):

- (A) $\frac{2}{3}E$
- (B) $\frac{1}{8}E$
- (C) $\frac{1}{4}E$
- (D) $\frac{1}{2}E$

Correct Answer: (C) $\frac{1}{4}E$

Solution:

Concept: In Simple Harmonic Motion (SHM), the total energy E of the oscillator is constant and is the sum of its kinetic energy and potential energy. The potential energy (U) at a displacement x is given by:

$$U = \frac{1}{2}kx^2$$

The total energy (E) occurs when the displacement is at its maximum (amplitude A), where kinetic energy is zero:

$$E = \frac{1}{2}kA^2$$

Step 1: Defining the displacement.

The problem states the particle is "half way to its end point." The end point of an oscillator is its amplitude A . Therefore, the displacement x is:

$$x = \frac{A}{2}$$

Step 2: Calculating Potential Energy at that point. Substitute $x = \frac{A}{2}$ into the potential energy formula:

$$U = \frac{1}{2}k\left(\frac{A}{2}\right)^2$$

$$U = \frac{1}{2}k\left(\frac{A^2}{4}\right)$$

$$U = \frac{1}{4}\left(\frac{1}{2}kA^2\right)$$

Step 3: Relating to Total Energy. Since we know that $E = \frac{1}{2}kA^2$, we can substitute E into our expression for U :

$$U = \frac{1}{4}E$$

Quick Tip: Remember that Potential Energy in SHM is proportional to the square of displacement ($U \propto x^2$). If the distance is halved, the energy becomes $(1/2)^2 = 1/4$ of the maximum energy.

12. A current loop of magnetic moment 0.5 Am^2 is placed in a magnetic field of 0.4 T . If the loop is rotated from an angle of 0° to 180° , what is the change in potential energy?

- (A) -0.4 J
- (B) 0.4 J
- (C) -0.2 J
- (D) 0.2 J

Correct Answer: (B) 0.4 J

Solution:

Concept: The potential energy (U) of a magnetic dipole (current loop) in a uniform magnetic field B is given by the scalar product:

$$U = -\vec{M} \cdot \vec{B} = -MB \cos \theta$$

where M is the magnetic moment and θ is the angle between the magnetic moment and the magnetic field.

Step 1: Calculating Initial Potential Energy. Initially, the angle $\theta_1 = 0^\circ$.

$$U_{\text{initial}} = -MB \cos(0^\circ)$$

Since $\cos(0^\circ) = 1$:

$$U_{\text{initial}} = -MB$$

Substituting values: $U_{\text{initial}} = -(0.5)(0.4) = -0.2 \text{ J}$.

Step 2: Calculating Final Potential Energy. Finally, the loop is rotated to $\theta_2 = 180^\circ$.

$$U_{\text{final}} = -MB \cos(180^\circ)$$

Since $\cos(180^\circ) = -1$:

$$U_{\text{final}} = -MB(-1) = +MB$$

Substituting values: $U_{\text{final}} = +(0.5)(0.4) = +0.2\text{J}$.

Step 3: Calculating the Change in Potential Energy. The change in potential energy (ΔU) is the difference between the final and initial states:

$$\Delta U = U_{\text{final}} - U_{\text{initial}}$$

$$\Delta U = 0.2\text{J} - (-0.2\text{J})$$

$$\Delta U = 0.2 + 0.2 = 0.4\text{J}$$

Quick Tip: The work done in rotating a dipole from θ_1 to θ_2 is always $\Delta U = MB(\cos \theta_1 - \cos \theta_2)$. For a 180° rotation from the stable equilibrium (0°), the work is always $2MB$.

13. Twenty seven drops of same size are charged at 220 V each. They combine to form a bigger drop. Calculate the potential of the bigger drop.

- (A) 1320 V
- (B) 1520 V
- (C) 1980 V
- (D) 660 V

Correct Answer: (C) 1980 V

Solution:

Concept: When small charged droplets coalesce into a single large drop, two physical quantities remain conserved: Total Volume and Total Charge. The electric potential V of a spherical drop of radius r and charge q is given by:

$$V = \frac{kq}{r}$$

By finding the relationship between the radius and charge of the small drops versus the large drop, we can determine the new potential.

Step 1: Relation between radii using Volume Conservation. Let the radius of each small drop be r and the radius of the big drop be R . The volume of 27 small drops equals the volume of 1 big drop:

$$27 \times \left(\frac{4}{3} \pi r^3 \right) = \frac{4}{3} \pi R^3$$

$$27r^3 = R^3$$

Taking the cube root on both sides:

$$R = 3r$$

Step 2: Relation between charges using Charge Conservation. Let the charge on each small drop be q and the charge on the big drop be Q .

$$Q = 27q$$

Step 3: Calculating the potential of the bigger drop. Potential of a small drop (v): $v = \frac{kq}{r} = 220 \text{ V}$. Potential of the big drop (V):

$$V = \frac{kQ}{R}$$

Substitute $Q = 27q$ and $R = 3r$:

$$V = \frac{k(27q)}{3r} = 9 \left(\frac{kq}{r} \right)$$

$$V = 9 \times v$$

$$V = 9 \times 220 = 1980 \text{ V}$$

Quick Tip: For n identical droplets combining, the potential of the big drop is always $V_{big} = n^{2/3} \times V_{small}$. Here, $27^{2/3} = (3^3)^{2/3} = 3^2 = 9$.

14. Two polaroids are oriented with their principal planes making an angle of 60° . Then the percentage of incident unpolarized light which passes through the system is:

- (A) 100%
- (B) 50%
- (C) 12.5%
- (D) 37.5%

Correct Answer: (C) 12.5%

Solution:

Concept: This problem involves two stages of light transmission:

1. Unpolarized light passing through the first polaroid (Polarizer): The intensity is always halved.
2. Polarized light passing through the second polaroid (Analyzer): The intensity follows Malus's Law: $I = I_0 \cos^2 \theta$.

Step 1: Transmission through the first polaroid. Let the incident unpolarized light intensity be I_{in} . After passing through the first polaroid, the light becomes plane-polarized, and its intensity I_1 is:

$$I_1 = \frac{I_{in}}{2}$$

Step 2: Transmission through the second polaroid (Malus's Law). The polarized light I_1 now hits the second polaroid at an angle $\theta = 60^\circ$. The resulting intensity I_2 is:

$$I_2 = I_1 \cos^2(60^\circ)$$

Substitute the value of $\cos(60^\circ) = \frac{1}{2}$:

$$I_2 = I_1 \left(\frac{1}{2}\right)^2 = \frac{I_1}{4}$$

Step 3: Calculating final percentage. Substitute I_1 from Step 1:

$$I_2 = \left(\frac{I_{in}}{2}\right) \times \frac{1}{4} = \frac{I_{in}}{8}$$

To find the percentage:

$$\text{Percentage} = \frac{I_2}{I_{in}} \times 100 = \frac{1}{8} \times 100 = 12.5\%$$

Quick Tip: Always remember: First polaroid \rightarrow divide by 2. Second polaroid \rightarrow multiply by $\cos^2 \theta$. For 60° , $\cos^2(60^\circ)$ is 0.25 or $1/4$. Half of $1/4$ is $1/8$, which is 12.5%.

15. The equation of path of a projectile moving in x-y plane is given by $y = x - \frac{x^2}{25}$, then the initial speed and maximum height of particle will be (take $g = 10 \text{ m/s}^2$):

- (A) $4\sqrt{10} \text{ m/s}$ and $\frac{29}{4} \text{ m}$
- (B) $25\sqrt{10} \text{ m/s}$ and $\frac{21}{4} \text{ m}$
- (C) $2\sqrt{10} \text{ m/s}$ and $\frac{31}{4} \text{ m}$
- (D) $5\sqrt{10} \text{ m/s}$ and $\frac{25}{4} \text{ m}$

Correct Answer: (D) $5\sqrt{10} \text{ m/s}$ and $\frac{25}{4} \text{ m}$

Solution:

Concept: The standard trajectory equation for a projectile launched with initial velocity u at an angle θ with the horizontal is:

$$y = x \tan \theta - \frac{gx^2}{2u^2 \cos^2 \theta}$$

By comparing the given equation $y = x - \frac{x^2}{25}$ with this standard form, we can extract the values for the launch angle and the initial velocity. Once these are found, the maximum height H_{max} can be calculated using the formula:

$$H_{max} = \frac{u^2 \sin^2 \theta}{2g}$$

Step 1: Comparing the equations to find the launch angle. Comparing $y = x - \frac{x^2}{25}$ with $y = x \tan \theta - \frac{gx^2}{2u^2 \cos^2 \theta}$: The coefficient of x gives:

$$\tan \theta = 1 \implies \theta = 45^\circ$$

At $\theta = 45^\circ$, we have:

$$\sin 45^\circ = \frac{1}{\sqrt{2}}, \quad \cos 45^\circ = \frac{1}{\sqrt{2}}, \quad \text{and} \quad \cos^2 45^\circ = \frac{1}{2}$$

Step 2: Finding the initial speed (u). The coefficient of x^2 gives:

$$\frac{g}{2u^2 \cos^2 \theta} = \frac{1}{25}$$

Substitute $g = 10$ and $\cos^2 \theta = 1/2$:

$$\frac{10}{2u^2(1/2)} = \frac{1}{25} \implies \frac{10}{u^2} = \frac{1}{25}$$

$$u^2 = 250 \implies u = \sqrt{25 \times 10} = 5\sqrt{10} \text{ m/s}$$

Step 3: Calculating the maximum height (H_{max}). Now, substitute $u^2 = 250$ and $\sin^2 \theta = 1/2$ into the maximum height formula:

$$H_{max} = \frac{u^2 \sin^2 \theta}{2g} = \frac{250 \times (1/2)}{2 \times 10}$$

$$H_{max} = \frac{125}{20}$$

Divide both by 5:

$$H_{max} = \frac{25}{4} \text{ m}$$

Thus, the initial speed is $5\sqrt{10}$ m/s and the maximum height is $\frac{25}{4}$ m.

Quick Tip: To find the maximum height quickly from a trajectory equation $y = ax - bx^2$, you can use calculus. The maximum height occurs at the vertex where $dy/dx = 0$. Here, $1 - (2x/25) = 0 \implies x = 12.5$. Plugging $x = 12.5$ into the equation gives $y = 12.5 - (12.5)^2/25 = 12.5 - 6.25 = 6.25$, which is exactly $25/4$.