

DPP No. 11 & 22

Topic : Mole Concept

Hints & Solutions

DPP No. # 11

- Mass of a neutron = 1.675×10^{-24} g mass of a proton = 1.672×10^{-24}
- Carbon is ${}_6\text{C}^{12}$ and silicon is ${}_{14}\text{Si}^{28}$.
- No change by doubling mass of electrons, however by reducing mass of neutron to half total atomic mass becomes $6 + 3$ instead of $6 + 6$. Thus reduced by 25%.
- ${}_Z\text{X}^A$, $A = N + P$
- $\text{NO}_3^- = 7 + 8 \times 3 + 1 = 32$
- fraction = $\frac{\text{vol. of nucleus}}{\text{vol. of atom}} = \frac{\frac{4}{3}\pi(10^{-13})^3}{\frac{4}{3}\pi(10^{-8})^3} = 10^{-15}$.
- $2(p + n) + 3p = 140 \quad \therefore 7x = 140 \quad \therefore x = 20$
 $\therefore p = e = n = 20 \quad \therefore \text{Total number of nucleons} = n + p = 40$
 $\therefore \text{Element} = \text{Calcium}$

DPP No. # 12

- $\text{PE} = -\frac{kZe^2}{r}$.
- * Isotopes have same atomic number but different mass number.
- Isobars have same mass number.
- * Isotones have same number of neutrons.
- Each has 10 electrons.
 In $\text{CH}_3^+ = 6 + 3 - 1 = 8 \text{ e}$
 In $\text{H}_3\text{O}^+ = 3 + 8 - 1 = 10 \text{ e}$
- * Isoelectronic species have same number of electrons.
- $\frac{R_1}{R_2} = \left(\frac{A_1}{A_2}\right)^{1/3}$

$$\Rightarrow \frac{1}{2} = \left(\frac{A_1}{A_2} \right)^{1/3}$$

$$\Rightarrow \frac{A_1}{A_2} = \frac{1}{8}$$

ratio of atomic mass number.

8.
$$r_0 = \frac{4KZe^2}{M_0v_0^2}$$

$$\Rightarrow 2r_0 = \frac{4KZe^2}{M_0v'^2}$$

$$\Rightarrow r_0v_0^2 = 2r_0v'^2$$

$$\Rightarrow v' = \frac{v_0}{\sqrt{2}}$$

9. Given
$$R = \frac{4KZe^2}{M_0v_0^2}$$

$$\therefore R' = \frac{4KZe^2}{M_0\left(\frac{v_0}{2}\right)^2} = 4R$$

$$\therefore \% \text{ error} = \frac{4R - R}{R} \times 100 = 300 \%$$

10. Use
$$R = \frac{4Kze^2}{m_\alpha v_\alpha^2}$$

11. Definition

12. In one second, wave can travel distance = $v \times \lambda = 10 \times 2.5 \text{ m} = 25 \text{ m}$
In 40 seconds, it will travel = $25 \times 40 \text{ m} = \mathbf{1000 \text{ m}}$.

DPP No. # 13

1. Maximum wave length will correspond to minimum frequency as $\lambda \propto \frac{1}{\nu}$, and that is given for red light in the spectrum.

$$\lambda_{\text{max.}} = \frac{C}{\nu_{\text{min.}}} = \frac{3 \times 10^8 \text{ m/s}}{4 \times 10^{14} \text{ s}^{-1}} = 750 \times 10^{-9} \text{ m.}$$

$$\Rightarrow 7500 \text{ \AA.}$$

2.
$$\lambda = \frac{C}{\nu} = \frac{3 \times 10^8 \text{ m/s}}{1200 \times 10^3 \text{ s}^{-1}} = 250 \text{ m} = 0.25 \text{ km.}$$

$$\bar{\nu} = \text{Wave no.} = \frac{1}{\lambda} = \frac{1 \text{ km}}{0.25 \text{ km}} = 4 \text{ wave per km.}$$

3. (a) $R = R_0 A^{1/3} \quad \therefore \frac{4}{3} \pi R^3 = \frac{4}{3} \pi R_0^3 A$
 $\therefore V \propto A \quad \therefore n = 1$

(b)
$$\bar{\nu} = \frac{\nu}{c} = \frac{7.5 \times 10^{14}}{3 \times 10^8} = 2.5 \times 10^6 \text{ m}^{-1}$$

$$4. \quad \frac{E_1}{E_2} = \frac{\lambda_2}{\lambda_1} = \frac{6000}{3000} = 2.$$

$$5. \quad \text{Use } E = \frac{nhc}{\lambda}, \quad \text{Here } n \text{ is number of photons.}$$

$$6. \quad \text{Photon absorbed} = \frac{hc}{300 \times 10^{-9}} = \frac{6.6 \times 10^{-34} \times 3 \times 10^8}{300 \times 10^{-9}} = 6.6 \times 10^{-19} \text{ Joule}$$

$$\text{One re-emitted photon energy} = \frac{hc}{500 \times 10^{-9}} = 3.96 \times 10^{-19} \text{ Joule}$$

$$\text{other photon have energy} = 6.6 \times 10^{-19} - 3.93 \times 10^{-19} = 2.65 \times 10^{-19} \text{ Joule.}$$

$$7. \quad \text{Use } E = \frac{nhc}{\lambda}$$

$$60 \times 60 = \frac{n \times 6.64 \times 10^{-34} \times 3 \times 10^8}{620 \times 10^{-9}}$$

$$n = 1.125 \times 10^{22}$$

$$8. \quad \text{Energy of one photon} = \frac{12400}{6200} = 2 \text{ eV} = 2 \times 96 = 192 \text{ KJ mol}^{-1}$$

$$\therefore \% \text{ of energy of photon converted to K.E. of A atoms} = \frac{192 - 144}{192} \times 100 = \frac{48}{192} \times 100 = 25\%$$

$$9. \quad E_{\text{emitted}} = \frac{50}{100} \times E_{\text{absorbed}}$$

$$\text{No. of emitted photons} \times \text{Energy of emitted photon} = \frac{50}{100} \times \text{No. of absorbed photon} \times \text{Energy of absorbed photon.}$$

$$\therefore 5x \times \frac{12400}{5000} = \frac{50}{100} \times 8x \times \frac{12400}{\lambda(\text{\AA})}$$

$$\lambda(\text{\AA}) = 4000 \text{ \AA}$$

DPP No. # 14

$$1. \quad \text{For I experiment, } hv_1 = W + KE_{\text{max1}} \quad \dots (1)$$

$$\text{For II experiment, } hv_2 = W + KE_{\text{max2}} \quad \dots (2)$$

$$\text{here, } v_2 = 2v_1 \text{ and } KE_{\text{max2}} = 3 KE_{\text{max1}}$$

$$\therefore 2hv_1 = W + 3 KE_{\text{max1}} \quad \dots (3)$$

$$\text{From (1) and (3): } hv_1 = 2KE_{\text{max1}} \quad \text{or} \quad h\left(\frac{v_2}{2}\right) = 2\left(\frac{KE_{\text{max2}}}{3}\right)$$

\therefore % of incident energy converted into max KE in II experiment

$$= \frac{KE_{\text{max2}}}{hv_2} \times 100 = \frac{3}{4} \times 100 = 75\%.$$

2. The maximum KE of photoelectron is corresponding to maximum stopping = 18.6 eV

$$\therefore E_{\text{incident}} = W + KE_{\text{max}}$$

$$\frac{12400}{400} \text{ eV} = W + 18.6 \text{ eV}$$

$$W = 12.4 \text{ eV}$$

$$\therefore \lambda_0 = \frac{12400}{12.4} \text{ \AA} = 1000 \text{ \AA}$$

3. Only for Single electron species.

$$4. \frac{1}{\lambda_1} = RZ^2 \left[\frac{1}{n_C} - \frac{1}{n_A} \right] \dots\dots (1)$$

$$\frac{1}{\lambda_1} = RZ^2 \left[\frac{1}{n_C} - \frac{1}{n_B} \right] \dots\dots (ii)$$

$$\frac{1}{\lambda_3} = RZ^2 \left[\frac{1}{n_B} - \frac{1}{n_A} \right]$$

$$\frac{1}{\lambda_3} = \frac{1}{\lambda_1} - \frac{1}{\lambda_2} = \frac{\lambda_2 - \lambda_1}{\lambda_1 \lambda_2} = \frac{1}{3000}$$

$$\lambda_3 = 3000 \text{ \AA}.$$

5. For $r = 0.52 \text{ \AA} \times \frac{12}{1}$

For $L^{2+} r_1 = 0.529 \times \frac{12}{3}$

$$\frac{r}{r_1} = 3 \quad \Rightarrow \quad r_1 = \frac{r}{3}$$

6. $r_4 - r_3 = 7 \times r_1$

7. Use $V_n = 2.185 \times 10^8 \left(\frac{Z}{n} \right) \text{ cm/sec.}$

8. $PE = - \frac{KZe^2}{r}.$

9. $\frac{nh}{2\pi} = \frac{2h}{\pi} \quad \Rightarrow n = 4,$

$$P.E. = 2(T.E.) = 2 \left(-13.6 \times \frac{2^2}{4^2} \right) = -6.8 \text{ eV.}$$

10. (A) $E_n^{-y} \propto r_n / Z$

$$\left(\frac{Z^2}{n^2} \times 13.6 \text{ eV} \right)^{-y} \propto \frac{1}{Z} \left(\frac{n^2}{Z} \times 0.529 \text{ \AA} \right)$$

$$y = 1$$

(B) $l_n \propto n^x \Rightarrow \frac{nh}{2\pi} \propto n^x \Rightarrow x = 1$

(C) Potential energy = 2 (total energy)

(D) $T_n \propto \frac{n^3}{Z^2} \Rightarrow t = -2 \quad \Rightarrow \quad m = -3.$

DPP No. # 15

1. $E_{I \text{ Photon}} + E_{II \text{ Photon}} = E_{\text{Single Photon}}$

$$hc \bar{\nu}_1 + hc \bar{\nu}_2 = \frac{hc}{\lambda}$$

$$\therefore \lambda = \frac{1}{\bar{v}_1 + \bar{v}_2} = \frac{1}{5.25 \times 10^8 + 7.25 \times 10^8} = \frac{1}{12.5 \times 10^8} = 8 \times 10^{-10} \text{ m} = 8 \text{ \AA}$$

2. Use : $E_1 - E_2 / E_2 - E_3$

$$3. \frac{r_1}{r_2} = \frac{1}{4} \quad \Rightarrow \quad \frac{r_3}{r_4} = \frac{9}{16} \quad \Rightarrow \quad \frac{r_2}{r_4} = \frac{1}{4}$$

So corresponding energy of ratio $\frac{1}{4}$ is $E_2 - E_1$ and $E_4 - E_2$.

$$4. \frac{R_1}{R_2} = \frac{4}{9} = \frac{n_1^2}{n_2^2}, \text{ hence } \frac{n_1}{n_2} = \frac{2}{3}. \text{ So, } \frac{f_1}{f_2} = \frac{n_2^3}{n_1^3} = \frac{27}{8}.$$

5. Electrostatic force of attraction $F \propto \frac{Z^3}{n^4}$

$$\therefore \frac{(F_{n=3})_{\text{He}^+}}{(F_{n=2})_{\text{Li}^{2+}}} = \frac{2^3/3^4}{3^3/2^4} = \left(\frac{2}{3}\right)^7 = \left(\frac{3}{2}\right)^{-7} \quad \therefore x = -7 \quad F = \frac{kZe^2}{R^2}$$

6. Order of energy \rightarrow Violet $>$ Blue $>$ yellow $>$ red
 Order of energy $\rightarrow E_{2 \rightarrow 1} > E_{5 \rightarrow 2} > E_{6 \rightarrow 3} > E_{4 \rightarrow 3}$
 \therefore Violet ($2 \rightarrow 1$), Blue ($5 \rightarrow 2$), yellow ($6 \rightarrow 3$), Red ($4 \rightarrow 3$)

7.* BE for ($n = 3$) = $1.51 Z^2 = 12 \text{ eV}$ (given)

$$\therefore Z^2 = 12/1.51$$

$$\text{I Excitation potential} = 10.2 Z^2 = 10.2 \times (12/1.51) = 81 \text{ V}$$

$$\text{II Excitation potential} = 12.09 Z^2 = 12.09 \times (12/1.51) = 96 \text{ eV}$$

$$\text{Ionisation potential} = 13.6 Z^2 = 13.6 (12/1.51) = 108 \text{ V}$$

$$\text{BE of } (n = 2) = 3.4 Z^2 = 3.4 \times (12/1.51) = 27 \text{ eV}$$

8. Let the given transition for both the species is $n_1 \rightarrow n_2$

$$\text{Then } X_{\text{cm}}^{-1} = R \times 2^2 \left[\frac{1}{n_1^2} - \frac{1}{n_2^2} \right] \text{ For He}^+ \quad \dots (i)$$

$$\text{and (wave no.) Be}^{3+} = R \times 4^2 \left[\frac{1}{n_1^2} - \frac{1}{n_2^2} \right] \text{ For Be}^{3+} \quad \dots (ii)$$

From eq. (i) and (ii) (wave no.) $\text{Be}^{3+} = 4x \text{ cm}^{-1}$.

DPP No. # 16

$$1. \lambda = \frac{h}{p} = \frac{h}{mv} = \frac{h}{\sqrt{2KE}}$$

$$\text{but } v^2 = \frac{2KE}{m} \quad \text{therefore } \lambda = \frac{hm}{2KE}$$

$$2. \text{ Use } C = v\lambda \quad \Rightarrow \quad \bar{u} = \frac{1}{\lambda}$$

$$3. \text{ K.E.}_{\text{proton}} = 1 + (1)(3) = 4 \text{ eV} \therefore \lambda_p = \frac{h}{\sqrt{2m_p(\text{KE})_p}} \quad \& \quad \text{KE}_{\alpha\text{-particle}} = 20 - (2)(2) = 16 \text{ eV} \therefore \lambda_\alpha = \frac{h}{\sqrt{2m_\alpha(\text{KE})_\alpha}}$$

$$\therefore \frac{\lambda_p}{\lambda_\alpha} = \sqrt{\frac{m_\alpha(\text{KE})_\alpha}{m_p(\text{KE})_p}} = \sqrt{\frac{4 \times 16}{1 \times 4}} = \frac{4}{1}$$

$$4.* \quad 4.25 = (W_0)_A + (\text{K.E.})_A$$

$$4.70 = (W_0)_B + (\text{K.E.})_A - 1.5$$

$$\text{So } (W_0)_B - (W_0)_A = 0.45 + 1.5 = 1.95$$

$$\text{Now, } \lambda_B = 2\lambda_A$$

$$\frac{h}{\sqrt{2m(\text{K.E})_B}} = \frac{2h}{\sqrt{2m(\text{K.E})_A}}$$

$$\text{So } (\text{K.E})_A = 4 (\text{K.E})_B$$

$$4.25 - (W_0)_A = 4 [4.7 - (W_0)_B]$$

$$4(W_0)_B - (W_0)_A = 14.55$$

$$\text{So } (W_0)_B = 4.2\text{eV}$$

$$\text{So } (W_0)_A = 2.25\text{ eV}$$

$$(\text{K.E.})_A = 2\text{eV}$$

$$(\text{K.E})_B = 0.5\text{eV}$$

5. number of revolutions per second

$$= \frac{v}{2\pi r} = \frac{2.18 \times 10^6 \left(\frac{Z}{n}\right)}{2 \times 3.14 \times 0.529 \times \left(\frac{n^2}{Z}\right) \times 10^{-10}} = \frac{2.18 \times 10^6 \left(\frac{1}{2}\right)}{2 \times 3.14 \times 0.529 \times \left(\frac{2^2}{1}\right) \times 10^{-10}}$$

$$\text{Number of revolution in } 10^{-8} \text{ second} = \frac{2.18 \times 10^6 \left(\frac{1}{2}\right)}{2 \times 3.14 \times 0.529 \times \left(\frac{2^2}{1}\right) \times 10^{-10}} \times 10^{-8} = 8.2 \times 10^6.$$

6. The ionisation energy of He^+ is 19.6×10^{-18} J.

$$\therefore \text{Energy of the first orbit of } \text{He}^+ (Z = 2) = 19.6 \times 10^{-18} \text{ J.}$$

$$\therefore \text{Energy of the first orbit of } \text{H}^+ (Z = 1) = \frac{19.6 \times 10^{-18}}{4} \text{ J}$$

$$\therefore \text{Energy of the first orbit of } \text{Li}^{2+} (Z = 3) = \frac{19.6 \times 10^{-18}}{4} \times 9 = 4.41 \times 10^{-17} \text{ J.}$$

7. (A) Transition $n \rightarrow 6$ to $n \rightarrow \infty$ For Li^{2+} sample
 (B) Transition $n \rightarrow 1$ to $n \rightarrow 2$ For H-atom sample
 (C) Transition $n \rightarrow 1$ to $n \rightarrow 3$ For He^+ sample
 (D) Transition $n \rightarrow 1$ to $n \rightarrow \infty$ For H-atom sample

8. $\Delta E = \frac{12400}{1026} = 12.09 \text{ eV.}$

$$\text{So, } \Delta E = E_3 - E_1.$$

Hence, induced radiations will be correspond to following energy transition

$$E_3 \rightarrow E_1, E_3 \rightarrow E_2 \text{ and } E_2 \rightarrow E_1.$$

9. $-13.6 \frac{Z^2}{n^2} = 4R = 4 \times 2.2 \times 10^{-18} \text{ J.}$

$$Z^2 = \frac{4 \times 2.2 \times 10^{-18} \text{ J}}{13.6 \times 1.6 \times 10^{-19}} = 4; \quad Z = 2.$$

$$r = 0.529 \frac{n^2}{Z} \times 10^{-10} \text{ m.} \quad r = 0.529 \times 10^{-10} \times \frac{1}{2} = 2.645 \times 10^{-11} \text{ m.}$$

DPP No. # 17

1. $Z = 2 \quad n_1 = 2 \quad n_2 = \infty$

$$\bar{v} = R(2)^2 \left(\frac{1}{2^2} - \frac{1}{\infty^2} \right) = R$$

2. $h\nu_1 = 13.6 \text{ eV}$
 $h\nu_2 = 13.6 \times 2^2 \text{ eV}$
 $h\nu_3 = 13.6 \times 2^2 \times \frac{3}{4} \text{ eV}$
 $\Rightarrow h\nu_2 = h\nu_1 + h\nu_3$
 $\Rightarrow \nu_2 = \nu_1 + \nu_3$

3. (i) Series limit of Lyman series $\Rightarrow n = \infty$ to $n = 1$.
(ii) Series limit of Balmer series $\Rightarrow n = \infty$ to $n = 2$.
 $E_{n=2 \text{ to } n=1} = E_{n=\infty \text{ to } n=1} - E_{n=\infty \text{ to } n=2}$

$$\frac{hc}{\lambda} = \frac{hc}{\lambda_1} - \frac{hc}{\lambda_2}$$

$$\frac{1}{\lambda} = \frac{1}{\lambda_1} - \frac{1}{\lambda_2} \quad \Rightarrow \quad \lambda = \frac{\lambda_1 \lambda_2}{\lambda_2 - \lambda_1}$$

4. Both the photons will not be absorbed by the electron of H-atom as the energy levels are quantised.

$$\text{Sum of energies of both photons} = \frac{12400}{1240} + \frac{12400}{2000} = 10 + 6.2 = 16.2 \text{ eV} > (IE)_H$$

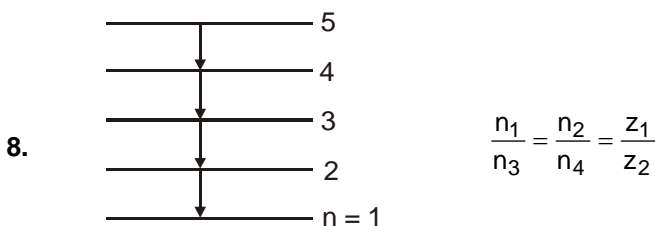
5. (A) Only first four spectral lines belonging to Balmer series in hydrogen spectrum lie in visible region.
(B) If a light of frequency ν falls on a metal surface having work functional $h\nu_0$, photoelectric effect will take place only if $\nu \geq \nu_0$, since ν_0 is the minimum frequency required for photoelectric effect.

6. $\frac{\Delta n(\Delta n + 1)}{2} = 15$
 $\Rightarrow \Delta n = 5 \quad \Rightarrow n - 2 = 5 \quad \Rightarrow n = 7$

7. $\bar{\nu}_1 = R \times 3^2 \left(\frac{1}{2^2} - \frac{1}{3^2} \right) = \frac{5R}{4}$

$$\bar{\nu}_2 = R \times \frac{3^2}{3^2} = R$$

$$\bar{\nu}_2 - \bar{\nu}_1 = \frac{5R}{4} - R = \frac{R}{4}$$



Clearly 2nd lowest energy is $4 \rightarrow 3$ transition
hence transition is Li^{2+} having same energy is $9 \rightarrow 12$

9. (A) $6 \rightarrow 3 \quad \Delta n = 3$
 $\therefore \text{no. of lines} = \frac{3(3+1)}{2} = 6.$ All lines are in infrared region
(B) $7 \rightarrow 3 \quad \Delta n = 4$

$$\therefore \text{no. of lines} = \frac{4(4+1)}{2} = 10. \quad \text{All lines are in infrared region}$$

$$(C) 5 \rightarrow 2 \quad \Delta n = 3$$

$$\text{no. of lines} = \frac{3(3+1)}{2} = 6. \quad \text{All lines are in visible region}$$

$$(D) 6 \rightarrow 2 \quad \Delta n = 4$$

$$\text{no. of lines} = \frac{4(4+1)}{2} = 10. \quad \text{All lines are in visible region.}$$

$$10. \quad \nu = R_c Z^2 \left[\frac{1}{n_1^2} - \frac{1}{n_2^2} \right]$$

$$\text{For } 2 \rightarrow 1 \text{ transition in H-atom sample, } \nu = R_c(1)^2 \left[\frac{1}{1^2} - \frac{1}{2^2} \right] = \frac{3R_c}{4}$$

$$\therefore (H)_{2 \rightarrow 1} = (He^+)_{4 \rightarrow 2} = (Li^{2+})_{6 \rightarrow 3}$$

Thus, given photon is not emitted from $8 \rightarrow 3$ transition in He^+ ion sample.

DPP No. # 18

1. Total spectral lines obtained from H-atom = 6
Total spectral lines obtained from He^+ -ion = 6
One line is common between them so total number of lines are 11.

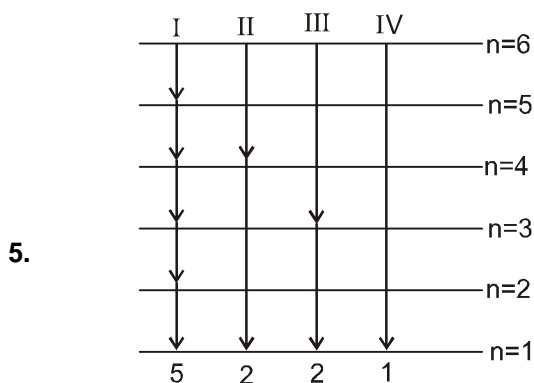
2. Balmer series lines lies in visible region.

3. $n_2 \rightarrow n_1$, max different spectral lines = 10
 $\therefore \Delta n = n_2 - n_1 = 4$

$$\therefore \text{change in angular momentum} = (\Delta n) \frac{h}{2\pi} = 4 \left(\frac{h}{2\pi} \right) = 8 \left(\frac{h}{4\pi} \right)$$

$$\therefore y = 8$$

4. $(Li^{2+})_{12 \rightarrow 3} = (H)_{4 \rightarrow 1}$
 \therefore No. of lines in infrared region = 1 ($4 \rightarrow 3$) paschen series



\therefore Minimum number of atoms required = 4

6. Let excited state be n.

Case - I : There is a transition to first excited state i.e. 2nd level from n^{th} level.

$$10.20 + 17.00 = 13.6 Z^2 \left[\frac{1}{2^2} - \frac{1}{n^2} \right] \quad \dots (1)$$

Case - II : There is a transition to second excited state i.e. 3rd level from n^{th} level.

$$4.25 + 5.95 = 13.6 Z^2 \left[\frac{1}{3^2} - \frac{1}{n^2} \right] \quad \dots (2)$$

on dividing (1) to (2), we have n^{th} level is = 6.

So, excited state is 5th.

So, $n = 5$.

7. $n\lambda = 2\pi r \Rightarrow$ so $\lambda = \frac{2\pi r}{3} = \frac{2\pi}{3} \times (53 \text{ pm}) \times \frac{9}{2} = 5\text{\AA}$

8. Use formula $2\pi r_n = n\lambda$
We can't apply Bohr radius formula for Be^{2+}
 $2\pi r_n = n\lambda$ | ~~vd km~~; ~~kd ja~~

9. $\frac{4}{n} = \frac{6}{3} \Rightarrow n = 2$ electron is present in 2nd orbit of Be^{3+} . (Be^{3+} d s f) r h d {k em i l f r by ~~DMW~~

$$2\pi r = 2\lambda \Rightarrow \lambda = \pi r \Rightarrow r = 0.529 \times 10^{-10} \times \frac{2^2}{4} = 0.529 \times 10^{-10} = 0.529\text{\AA}.$$

DPP No. # 19

- Only Spin quantum number (s) is not derived from Schrodinger wave equation.
- number of electrons in subshells = $2(2l + 1)$
- For $n = 8$ to $n = 6$, energy difference is minimum and $\lambda \propto \frac{1}{\text{Energy}}$
- S_1 : Photoelectric effect can be explained on the basis of particle nature of electromagnetic radiations.
 S_2 : $n = 2, \ell = 1 \therefore$ 2p-orbital \therefore dumb-bell shaped.
 S_3 : d_{xy} orbital has its lobes directed at an angle of 45° from X-axis and Y-axis. So, it has zero probability of finding electrons along X-axis and Y-axis.
- S_1 : Angular momentum = $mvr = n \left(\frac{h}{2\pi} \right) \therefore$ Angular momentum $\propto n$.
 S_2 : An orbital can only accommodate 2 electrons with opposite spin.
 S_3 : s-orbital is non-directional in nature, rest all orbitals are directional.

DPP No. # 20

- $n = 4, m = -3 \therefore$ only possible value of ℓ is 3.
 \therefore Orbital angular momentum = $\sqrt{\ell(\ell+1)} \frac{h}{2\pi} = \frac{2\sqrt{3}h}{2\pi} = \frac{\sqrt{3}h}{\pi}$.
- $Z = 26 \rightarrow [\text{Ar}]4s^2 3d^6$
 $\sqrt{n(n+2)} = \sqrt{24} \Rightarrow n = 4$
In d orbital number of unpaired electron = 4, but element have charge so 4s electron have to be removed hence $n+ = 2$.
- V ($Z = 23$), $[\text{Ar}] 4s^2 3d^3$ unpaired electron = 3 ;
Cr ($Z = 24$), $[\text{Ar}] 4s^1 3d^5$ unpaired electron = 6
Mn ($Z = 25$), $[\text{Ar}] 4s^2 3d^5$ unpaired electron = 5
- For $n = 4, \ell \neq 4$, for $\ell = 3, m \neq 4$

5. Total spin = 3 $\Rightarrow \frac{n}{2} = 3 \Rightarrow n = 6$

i.e. magnetic moment = $\sqrt{n(n+2)} = \sqrt{6(6+2)} = \sqrt{48}$ B.M.

6. $25\text{Mn} - [\text{Ar}] 3d^5 4s^2$



Given $\sqrt{n(n+2)} = \sqrt{15} \Rightarrow n = 3$

Hence to have '3' unpaired electrons Mn must be in '+4' state.

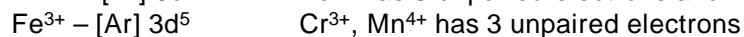
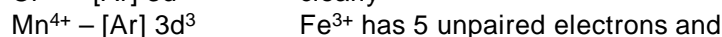
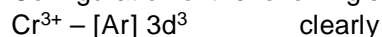
7. Magnetic moment = $\sqrt{n(n+2)}$

8. Orbital angular momentum of electron

= $\sqrt{\ell(\ell+1)} \frac{h}{2\pi} \Rightarrow \sqrt{\ell(\ell+1)} \frac{h}{2\pi} = \sqrt{3} \frac{h}{\pi} \Rightarrow \ell = 3$

\therefore number of orientations = $2\ell + 1 = 2 \times 3 + 1 = 7$

9. Configuration of the following elements



10. Maximum possible number of electrons in an atom with $(n + \ell = 7) = 7s(2) + 6p(6) + 5d(10) + 4f(14) = 32$

11. total spin = $\pm 1/2 \times \text{No. of Unpaired } e^-$

DPP No. # 21

1. Definition

2. $\lambda = \frac{12.3}{\sqrt{v}}$

3. ${}_{16}\text{S}^{32} = e^- = 16$

$x^{+2} = e^- = 16$

($\because A = Z + N$)

4. (B) has same number of electrons i.e., 18.

$[\text{NH}_3 \rightarrow \text{BH}_3] = 10 + 8 = 18.$

5. E.C. $\rightarrow 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 3d^1, 4s^2$

6. (a) $\text{Co}^{3+} : 1s^2 2s^2 2p^6 3s^2 3p^6 3d^6 \therefore$ 4 unpaired electrons $\therefore \mu = \sqrt{4(4+2)} = \sqrt{24} = 4.9$ BM

(b) Number of radial nodes = $n - \ell - 1$

Number of radial nodes in 3p orbital = $3 - 1 - 1 = 1$

(c) Number of electrons with $(m = 0)$ in $\text{Mn}^{2+} (1s^2 2s^2 2p^6 3s^2 3p^6 3d^5)$ ion = $1s(2) + 2s(2) + 2p(2) + 3s(2) + 3p(2) + 3d(1) = 11$

(d) Orbital angular momentum for the unpaired electron in V^{4+} lies in 3d orbital. $\therefore \ell = 2$

\therefore Orbital angular momentum = $\sqrt{\ell(\ell+1)} \frac{h}{2\pi} = \frac{\sqrt{6} h}{2\pi}$

7. (a) $x + e^- \rightarrow x^-$

energy released = $E.A_1 = 30.87$ eV/atom

Let no. of moles of X be a

$\therefore a \times N_A \times 30.87 = 4 \times N_A \times 4.526 + 4 \times N_A \times 13.6 + 4 \times N_A \times 12.75 \Rightarrow a = 4$ moles.

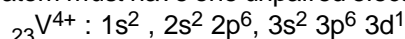
8. Number of unpaired electron are given by

$$\text{Magnetic moment} = \sqrt{n(n+2)} \text{ B.M.}$$

where n is number of unpaired electrons

$$\text{or } 1.73 = \sqrt{n(n+2)} \quad \text{or } 1.73 \times 1.73 = n^2 + 2n \quad \therefore n = 1$$

Now Vanadium atom must have one unpaired electron and thus its configuration is



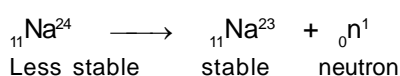
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8. A and D are isotopes. B, C and D are isobars.

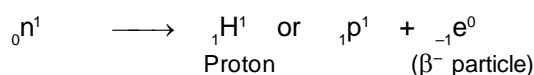
9. (i) p,s (ii) q,r (iii) p,s (iv) q,r

10. Isotopic (${}_{11}\text{Na}^{24}$) is less stable than ${}_{11}\text{Na}^{23}$ because it show radioactive decays. (Less stability of Na^{24} w.r.t.

Na^{23} is also based upon $\frac{13}{11}\left(\frac{n}{p}\right)$. Higher value of $\frac{n}{p}$, higher will be unstability so it is disintegrated to attain the stability).



This neutron on decomposition to give proton and β^- particle (${}_{-1}^0\text{e}^0$)



Hence, isotopic sodium is changed into sodium by means of emission of β^- emission.

11. (i) The atomic mass of an element reduces by 4 and atomic number by 2 on emission of an α -particle.
 (ii) The atomic mass of an element remains unchanged and atomic number increases by 1 on emission of a β -particle.

Thus change in atomic mass on emission of 8α -particles will be $8 \times 4 = 32$

New atomic mass = old atomic mass - 32 = 238 - 32 = 206

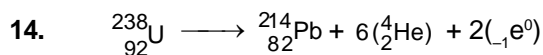
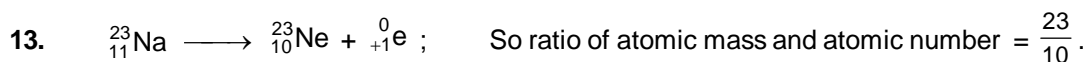
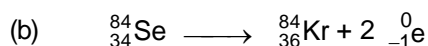
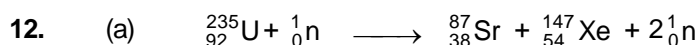
Similarly change in atomic number on emission of 8α -particle will be : $8 \times 2 = 16$

i.e., New atomic number = old atomic number - 16 = 92 - 16 = 76

On emission of 6β -particles the atomic mass remains unchanged thus, atomic mass of the new element will be 206.

The atomic number increases by 6 unit thus new atomic number will be $76 + 6 = 82$

Thus, the equation looks like : ${}_{92}\text{X}^{238} \xrightarrow[-6\beta]{-8\alpha} {}_{82}\text{Y}^{206}$



$\alpha = 6, \beta = 2$

Total = 8

