Exercise-1

> Marked questions are recommended for Revision.

PART - I : OBJECTIVE QUESTIONS

1. What possibly can be the ratio of the de Broglie wavelengths for two electrons each having zero initial energy and accelerated through 50 volts and 200 volts? (A) 3 : 10 (B) 10:3 (C) 1:2 (D) 2:1 2. The speed of a proton is one hundredth of the speed of light in vacuum. What is its de-Broglie wavelength? Assume that one mole of protons has a mass equal to one gram $[h = 6.626 \times 10^{-27} \text{ erg}]$ secl: (A) 13.31 × 10⁻⁷ Å (B) 1.33 × 10⁻³ Å (D) 1.31 × 10⁻² Å (C) 13.13 × 10⁻⁵ Å An a-particle is accelerated through a potential difference of V volts from rest. The de-Broglie's 3.2 wavelength associated with it is (A) $\sqrt{\frac{150}{V}} Å$ (B) <u>0.286</u> Å (C) $\frac{0.101}{\sqrt{V}}$ Å (D) $\frac{0.983}{\sqrt{N}}$ Å The uncertainity in position and velocity of a particle are 10^{-10} m and 5.27×10^{-24} ms⁻¹ respectively. 4.2 Calculate the mass of the particle (h = 6.625×10^{-34} Joule sec.) (A) 0.099 Kg (B) 0.089 Kg (C) 0.99 Kg (D) Can not predict The de Broglie equation suggests that an electron has 5.2 (A) Particle nature (B) Wave nature (C) Particle-wave nature (D) Radiation behaviour It the uncertainity in position of a moving particle is 0 then find out ΔP 6. (A) 0 (B) 1 (C) ∞ (D) Can not predict 7. Which of the following has least de Broglie λ (A) e⁻ (B) p (C) CO₂ (D) SO₂ The orbital with zero orbital angular momentum is : 8. (A) s (B) p (C) d (D) f Which of the following is electronic configuration of Cu^{2+} (Z = 29)? 9. (B) [Ar]4s² 3d¹⁰ 4p¹ (A) [Ar]4s¹ 3d⁸ (C) [Ar]4s¹ 3d¹⁰ (D) [Ar] 3d9 Spin magnetic moment of Xⁿ⁺ (Z = 26) is $\sqrt{24}$ B.M. Hence number of unpaired electrons and value of n 10.2 respectively are : (A) 4, 2 (B) 2, 4 (C) 3, 1 (D) 0, 2 Which of the following ions has the maximum number of unpaired d-electrons? 11. (A) Zn²⁺ (B) Fe²⁺ (C) Ni3+ (D) Cu+ The total spin resulting from a d⁷ configuration is : 12. (A) 1 (B) 2 (C) 5/2 (D) 3/2 13.2 Given is the electronic configuration of element X : Κ L Μ Ν 2 8 2 11 The number of electrons present with $\Box = 2$ in an atom of element X is : (B) 6 (D) 4 (A) 3 (C) 5

14.	The possible value of \Box and m for the last electron in the Cl ⁻ ion are : (A) 1 and 2 (B) 2 and +1 (C) 3 and -1 (D) 1 and -1								
15.2	For an electron, with n = 3 has only one radial node. The orbital angular momentum of the electron w								
	(A) 0								
16.	The correct set of quantum no. for the unpaired electron of chlorine.								
	n I m n I m (A) 2 1 0 (B) 2 1 1 (C) 3 1 1 (D) 3 0 0								
17.	 Which of the following quantum number has not been derived from Schrodinger wave equation : (A) Principal quantum number (n) (B) Subsidiary quantum number (□) (C) Magnetic quantum number (m) (D) Spin quantum number (s) 								
18.	The orbital angular momentum corresponding to $n = 4$ and $m = -3$ is :								
	(A) 0								
19.*	The correct representation of electronic configuration of Nitrogen atom is : (A) $1 \downarrow$ $1 \downarrow$ $1 \downarrow$ $1 \downarrow$ 1 1 (B) $1 \downarrow$ $1 \downarrow$ $1 \downarrow$ $1 \downarrow$ $1 \downarrow$ 1 (C) $1 \downarrow$ $2 s$ $2 p$ (D) $1 \downarrow$ $2 s$ $2 p$ (D) $1 \downarrow$ $2 s$ $2 p$								
20.*	 Which of the following statements is/are INCORRECT : (A) The value of magnetic quantum number (m) cannot exceed the value of principal quantum number (n) for the same electron. (B) If the electronic configuration of ₆C is written as 1s⁶, then Aufbau's principle has been violated. 								
	(C) The $+\frac{1}{2}$ and $-\frac{1}{2}$ values of spin quantum number denote clockwise and anticlockwise spin o								
	electrons on its axis respectively. (D) The maximum number of electrons in a particular subshell, for which value of azimuthal quantum number is \Box , is given by (4 \Box +2).								
	PART - II : SUBJECTIVE QUESTIONS								
I .	How many unpaired electrons are there in Ni ⁺² ion if the atomic number of Ni is 28.								
2.	Write the electronic configuration of the element having atomic number 56.								
3.	Given below are the sets of quantum numbers for given orbitals. Name these orbitals.(a) $n = 3$ (b) $n = 5$ (c) $n = 4$ $\Box = 1$ $\Box = 2$ $\Box = 1$								
	(d) $n = 2$ (e) $n = 4$ $\Box = 0$ $\Box = 2$								
4.24	Point out the angular momentum of an electron in, (a) 4s orbital (b) 3p orbital (c) 4 th orbit (according to Bohr model)								

5.2	Which of the follow	ring sets of quantun	n numbers are	impossible for e	electrons?	Explain why	in each
	case.						

Set	n		m	S
(i)	1	0	1	$+\frac{1}{2}$
(ii)	3	0	0	$-\frac{1}{2}$
(iii)	1	2	2	$+\frac{1}{2}$
(iv)	4	3	-3	$+\frac{1}{2}$
(v)	5	2	1	$-\frac{1}{2}$
(v)	3	2	1	0

6. Find the total spin and spin magnetic moment of following ion.

(ii) Cu⁺

(i) Fe⁺³

Exercise-2

> Marked Questions may have for Revision Questions.

OBJECTIVE QUESTIONS

- **1.** The Uncertainity in the momentum of an electron is 1.0×10^{-5} kg m s⁻¹. The Uncertainity in its position will be: (h = 6.626×10^{-34} Js) (A) 1.05×10^{-28} m (B) 1.05×10^{-26} m (C) 5.27×10^{-30} m (D) 5.25×10^{-28} m
- 2. A ball weight 25 g moves with a velocity of 6.6×10^4 cm/sec then find out the de Broglie wavelength. (A) 0.4×10^{-33} cm (B) 0.4×10^{-31} cm (C) 0.4×10^{-30} cm (D) 0.4×10^{20} cm
- 3. Calculate the uncertainity in velocity of a cricket ball of mass 150 g if the uncertainity in its position is of the order of 1 Å (h = 6.6×10^{-34} Kg m² s⁻¹) (A) 3.499×10^{-24} ms⁻¹ (B) 3.499×10^{-21} ms⁻¹ (C) 3.499×10^{-20} ms⁻¹ (D) 3.499×10^{-30} ms⁻¹
- 4. Which orbital is non-directional. (A) s (B) p (C) d (D) All
- For which orbital angular probability distribution is maximum at an angle of 45° to the axial direction (A) d_{x²-v²}
 (B) d_{z²}
 (C) d_{xy}
 (D) P_x
- 6. ▲ If n and □ are respectively the principal and azimuthal quantum numbers, then the expression for calculating the total number of electrons in any orbit is -

(A)
$$\sum_{\ell=1}^{\ell=n} 2(2\ell+1)$$
 (B) $\sum_{\ell=1}^{\ell=n-1} 2(2\ell+1)$ (C) $\sum_{\ell=0}^{\ell=n+1} 2(2\ell+1)$ (D) $\sum_{\ell=0}^{\ell=n-1} 2(2\ell+1)$

7. The quantum numbers $+\frac{1}{2}$ and $-\frac{1}{2}$ for the electron spin represent :

(A) Rotation of the electron in clockwise and anticlockwise direction respectively.

(B) Rotation of the electron in anticlockwise and clockwise direction respectively.

- (C) Magnetic moment of the electron pointing up and down respectively,
- (D) Two quantum mechanical spin states which have no classical analogue.

8.	(A) 0, 0, √6 □, √2 □	f the orbital angular mon	(B) 1, 1, √4 □, √2 □	the orbitals 1s, 3s, 3d and 2p -					
	(C) 0, 1, √6 □, √3 □		(D) 0, 0, √20 □, √6						
9.	After np orbitals are fill (A) (n + 1) s	ed, the next orbital filled (B) (n + 2) p	will be : (C) (n + 1) d	(D) (n + 2) s					
10.2	Which of the above sta	atement (s) is/are <u>false.</u>							
	I. Orbital angular mor	nentum of the electron h	naving n = 5 and having	value of the azimuthal quantum					
	number as lowest for t	his principle quantum nu	imber is $\frac{h}{\pi}$.						
	II. If $n = 3$, $\Box = 0$, $m = 0$, for the last valence shell electron, then the possible atomic number may be or 13.								
	III. Total spin of electro	ons for the atom 25Mn is	$\pm \frac{7}{2}$.						
	IV. Spin magnetic mor		2						
	(A) I, II and III	(B) II and III only	(C) I and IV only	(D) None of these					
11.24	In case of $d_{x^2-y^2}$ orbita	ıl							
	(A) Probability of findir(B) Probability of findir(C) Probability of findir	ng the electron along x-a ng the electron along y-a ng the electron is maxim ng the electron is zero in	xis is zero. um along x and y-axis.						
12.	Which of the following (A) The value of □ may (C) The value of s may	y be 2.	t for an electron of quan (B) The value of □ ma (D) All of these	tum numbers n = 4 and m = 2 ? ay be 3.					
13. 🕿	(A) The electronic cont(B) The magnetic quar	ntum number may have	d) ⁵ (4s) ¹ . (Atomic number negative values.	of Cr = 24)					
14. 🙇	Two unpaired electron (A) Principle quantum (C) Magnetic quantum	number	n are different with respe (B) Azimuthul quantur (D) Spin quantum nur	n number					
15. 🙇	Number of electron ha	ving the quantum numb	ers n = 4, \Box = 0, s = $-\frac{1}{2}$	in Zn ⁺² ion is/are :					
	(A) 1	(B) 0	(C) 2	(D) 5					
16.2a	Spin angular momentu	-	in sodium (Atomic No. =	11) is					
	(A) $\frac{\sqrt{3}}{2}$	(B) 0.866 h/2π	$(C) - \frac{\sqrt{3}}{2} \frac{h}{2\pi}$	(D) None of these					
17.	(A) $\frac{1}{2}$ (B) 0.866 h/2 π (C) $-\frac{1}{2} \frac{1}{2\pi}$ (D) None of these Statement-1 : For n = 3, \Box may be 0, 1 and 2 and 'm' may be 0, \pm 1 and \pm 2. Statement-2 : For each value of n, there are 0 to (n - 1) possible values of \Box ; for each value of \Box , there are 0 to \pm values of m. (1) Statement-1 is true, statement-2 is true; Statement-2 is a correct explanation for statement-1. (2) Statement-1 is true, statement-2 is true; Statement-2 is NOT a correct explanation for statement-1. (3) Statement-1 is true, statement-2 is false. (4) Statement-1 is false, statement-2 is true.								

- **18.** Statement-1 : The possible number of electrons in a subshell is $(4 \square + 2)$
 - Statement-2: The possible number of orientations of a sub-shell are $(2\Box + 1)$
 - (1) Statement-1 is true, statement-2 is true; Statement-2 is a correct explanation for statement-1.
 - (2) Statement-1 is true, statement-2 is true; Statement-2 is NOT a correct explanation for statement-1.
 - (3) Statement-1 is true, statement-2 is false.
 - (4) Statement-1 is false, statement-2 is true.

Exercise-3

* Marked Questions may have more than one correct option.

PART - I : JEE (ADVANCED) / IIT-JEE PROBLEMS (PREVIOUS YEARS)

1. The orbital angular momentum of an electron in 2s-orbital is :

(2) zero

 $(1) + \frac{1}{2} \frac{h}{2\pi}$

(3)
$$\frac{h}{2\pi}$$

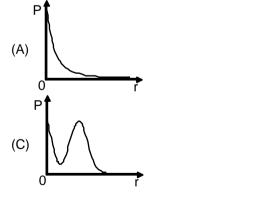
(4) $\sqrt{2} \frac{h}{2\pi}$

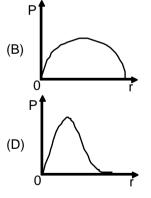
[JEE 1996]

- 2. A compound of Vanadium has a magnetic moment of 1.73 BM. Work out the electronic configuration of the Vandium ion in the compound. [JEE 1997]
- 3. Find the orbital angular momentum of electron if it is in 2p orbital of H in terms of $\frac{h}{2\pi}$.

[JEE 2005, 4/144]

- 4. The maximum number of electrons that can have principal quantum number, n = 3, and spin quantum number, $m_s = -1/2$, is [JEE 2011, 4/180]
- 5. In an atom, the total number of electrons having quantum numbers n = 4, $|m_{\Box}| = 1$ and $m_s = -1/2$ is [JEE(ADVANCED)-2014, 3/120]
- Not considering the electronic spin, the degeneracy of the second excited state (n = 3) of H atom is 9, while the degeneracy of the second excited state of H⁻ is : [JEE(ADVANCED)-2015, 4/168]
- 7. P is the probability of finding the 1s electron of hydrogen atom in a spherical shell of infinitesimal thickness, dr, at a distance r from the nucleus. The volume of this shell is $4\pi r^2 dr$. The qualitative sketch of the dependence of P on r is: [JEE(Advanced) 2016, 3/124]





	Answer Q.8, Q.9 and Q.1 of the following table.	0 by appropriately n	natching the inform	ation given in the three columns					
	The wave function, $\psi_{n, l}$,	m, is a mathematical	function whose val	ue depends upon spherical polar					
	coordinates (r, θ , ϕ) of the electron and characterized by the quantum numbers <i>n</i> , <i>I</i> and <i>m</i> _{<i>l</i>} . Here distance from nucleus, θ is colatitude and ϕ is azimuth. In the mathematical functions given in Table. Z is atomic number and ϕ is Pabr radius.								
	Table, Z is atomic number and a₀ is Bohr radius.Column 1Column 2Column 3								
	(I) 1s orbital	(i) $\psi_{n, l, m_l} \propto \left(\frac{1}{2} \right)^{1/2}$	$\frac{z}{a_{o}}\right)^{\frac{3}{2}}e^{-\left(\frac{zr}{a_{o}}\right)}$	(P) $\psi_{n, l, m_l}(r) = 0$					
	(II) 2s orbital	(ii) One radial n	ode	(Q) Probability density at nucleus $\propto \frac{1}{a_o^3}$					
	(III) 2p _z orbital	(iii) $\psi_{n, l, m_l} \propto \left(\frac{1}{2} \right)^{-1}$	$\frac{Z}{a_o} \int_{0}^{\frac{5}{2}} r e^{-\left(\frac{Zr}{2a_o}\right)} cos\theta$	(R) Probability density is maximum at nucleus					
	(IV) 3d _z ² orbital	(iv) xy-plane is a	a nodal plane	(S) Energy needed to excite electron from n = 2 state to n = 4 state is $\frac{27}{32}$ times the energy needed to excite electron from n = 2 state to n = 6 state					
8.	For He⁺ ion, the only INCO (A) (I) (i) (S) (B	RRECT combination) (II) (ii) (Q)	is (C) (I) (iii) (R)	[JEE(Advanced) 2017, 3/122] (D) (I) (i) (R)					
9.	-	olumn 1, the only C() (I) (ii) (S)	DRRECT combinatio (C) (IV) (iv) (R)	n for any hydrogen-like species is [JEE(Advanced) 2017, 3/122] (D) (III) (iii) (P)					
10.	For hydrogen atom, the on			[JEE(Advanced) 2017, 3/122] (D) (I) (i) (S)					
	PART - II : JEE (M	AIN) / AIEEE P	ROBLEMS (P	REVIOUS YEARS)					
		JEE(MAIN) OFFL	INE PROBLEMS						
1.	Which of the following ions (1) Mn ⁺² (2	agnetic moment? (3) Ti ⁺²	[AIEEE 2002, 3/225] (4) Cr ⁺² .						
2.	Uncertainity in position of m.sec ⁻¹) is: (plank's constation (1) 2.1×10^{-18} (2)	Hence, Uncertainity in velocity (in [AIEEE 2002, 3/225] (4) 5.0 × 10 ⁻²⁴							
3.	approximately (planck's o		-	ring with a velocity of 10 m/s is [AIEEE 2003, 3/225] (4) 10 ⁻²⁵ m					
4.	The numbers of d-electron	s retained in Fe ²⁺ (ato	omic number Fe = 26) ion is [AIEEE 2003, 3/225]					

5.	The orbital angular momentum for an electro	n revolving in an orbit	
	momentum for an s-electron will be given by		[AIEEE 2003, 3/225]
	(1) $+\frac{1}{2} \cdot \frac{h}{2\pi}$ (2) Zero	(3) $\frac{h}{2\pi}$	(4) $\sqrt{2} \cdot \frac{h}{2\pi}$
6.	Which of the following set a of quantum numbe	rs is correct for an electr	on in 4f orbital? [AIEEE 2004, 3/225]
	(1) n = 4, l =3, m = +4, s = +1/2 (3) n = 4, l = 3, m = +1, s = +1/2	(2) n = 4, l = 4, m = -4, (4) n = 3, l=2, m =-2, s	, s = -1/2
7.	Consider the ground state of Cr atom (Z = 24) numbers, \Box = 1 and 2 are, respectively (1) 12 and 4 (2) 12 and 5	. The numbers of electro (3) 16 and 4	ons with the azimuthal quantum [AIEEE 2004, 3/225] (4) 16 and 5
8.	In a multi-electron atom, which of the followin have the same energy in the absence of magne (i) $n = 1$, $l = 0$, $m = 0$ (ii) $n = 2$, $l = 0$, (iv) $n = 3$, $l = 2$, $m = 1$ (v) $n = 3$, $l = 2$, (1) (iv) and (v) (2) (iii) and (iv)	etic and electric field ? m = 0 (iii) n = m = 0	[AIEEE 2005, 3/225] = 2, l = 1, m = 1
9.	 (1) (1) and (1) (2) (1) and (1) (3) 3p orbital is lower in energy than 3d orbital (4) 3s orbital is lower in energy than 3p orbital 	the hydrogen atom is co ergy	
10.	Uncertainity in the position of an electron (mas Accurate upto 0.001%, will be : (h = 6.63×10^{-3} (1) 19.2 × 10 ⁻² m (2) 5.76 × 10 ⁻² m	³⁴ J-s)	[AIEEE 2006, 3/165]
11.	The 'spin-only' magnetic moment [in units of B (Atomic number : Ni = 28) (1) 2.84 (2) 4.90	ohr magneton (μ _β)] of Ni (3) 0	i ²⁺ in aqueous solution would be [AIEEE 2006, 3/165] (4) 1.73
12.	Which of the following set of quantum numbers	represents the highest e	energy of an atom ? [AIEEE 2008, 3/105]
	(1) n = 3, l = 0, m = 0, s = $+\frac{1}{2}$	(2) n = 3, l = 1, m =1, s	
	(3) n = 3, l = 2, m = 1, s = $+\frac{1}{2}$	(4) n = 4, l = 0, m = 0, s	$s = +\frac{1}{2}$
13.	Calculate the wavelength (in nanometer) assoproton = 1.67×10^{-27} kg and h = 6.63×10^{-34} J-(1) 0.40 nm (2) 2.5 nm		ving at 1.0 × 10 ³ m s ⁻¹ (Mass of [AIEEE 2009, 4/144] (4) 0.032 nm
14.	In an atom, an electron is moving with a speed which the position of the electron can be locate \times 10 ⁻³¹ kg):	d is (h = 6.6 × 10 ^{−34} kg n	
	(1) 5.10 × 10 ^{−3} m (3) 3.83 × 10 ^{−3} m	(2) 1.92 × 10 ^{−3} m (4) 1.52 × 10 ^{−4} m	
15.	The electrons identified by quantum numbers n (a) $n = 4$, $\Box = 1$ (b) $n = 4$, $\Box = 0$ can be placed in order of increasing energy as	(c) n = 3, □ = 2 :	[AIEEE 2012, 4/120] (d) n = 3, □ = 1
	(1) (c) < (d) < (b) < (a) (3) (b) < (d) < (a) < (c)	(2) (d) < (b) < (c) < (a) (4) (a) < (c) < (b) < (d)	

Qua									
16.				of rubidium atom (Z = 37) is : [JEE(Main)2014, 4/120]					
	(1) 5, 0, 0, + 1 /2	(2) 5, 1, 0, $+\frac{1}{2}$	(3) 5, 1, 1, + <mark>1</mark>	(4) 5, 0, 1, $+\frac{1}{2}$					
17.	potential differen	ce V esu. If e and m are c	harge and mass of an e	veen two charged plates kept at a electron, respectively, then the value					
	of $\frac{h}{\lambda}$ (where λ is	wavelength associated wit	th electron wave) is give	n by: [JEE(Main) 2016, 4/120]					
	(1) 2meV	(2) √meV	(3) √2meV	(4) meV					
18.	(Planck's Const.		ass of electron = 9.1097	I × 10 ⁻³¹ kg; charge of electron e = ¹ m ⁻³ A ²) [JEE(Main) 2017, 4/120] (4) 1.65 Å					
		JEE(MAIN) (ONLINE PROBLEMS						
1.	-	vavelength of a particle of n	[JEE(M	ain) 2014 Online (12-04-14), 4/120]					
	(1) 10 ⁻³³ m	(2) 10 ⁻³⁵ m	(3) 10 ⁻³¹ m	(4) 10 ⁻²⁵ m					
2.		uantum number $n = 6$, the c	[JEE(M	ain) 2015 Online (10-04-15), 4/120]					
	(1) ns→np→(n (3) ns→(n – 2)f-	, , ,	(2) ns \rightarrow (n – 1)d \rightarrow	() I					
		• • • •	(4) ns \rightarrow (n – 2)f \rightarrow						
3.	At temperatuere T, the average kinetic energy of any particle is $\frac{3}{2}$ KT. The de Broglie wavelength								
	(1) Visible photo (2) Thermal proto (3) Thermal proto	follows the order : [JEE(Main) 2015 Online (11-04-15), 4/120] (1) Visible photon > Thermal neutron > Thermal electron (2) Thermal proton > Thermal electron > Visible photon (3) Thermal proton > Visible photon > Thermal electron (4) Visible photon > Thermal electron > Thermal neutron							
4.	The total numbe	r of orbitals associated with							
	(1) 5	(2) 20	[JEE(Mai) (3) 25	n) 2016 Online (09-04-16), 4/120] (4) 10					
5.	Aqueous solution	n of which salt will not conta		e electronic configuration 1s ² 2s ² 2p ⁶ 3s ² 3p ⁶ ? [JEE(Main) 2016 Online (10-04-16), 4/120]					
	(1) NaCl	(2) CaI ₂	(3) NaF	(4) KBr					
6.	The de-Broglie's	wavelength of electron pre		'H' atom is : ain) 2018 Online (15-04-18), 4/120]					
	(1) 0.529 Å	(2) 2π × 0.529 Å	(3) $\frac{0.529}{2\pi}$ Å	(4) 4 × 0.529 Å					
7.	Which of the fo orbitals?	Which of the following combination of statements is true regarding the interpretation of the atomic orbitals? [JEE(Main) 2019 Online (09-01-19), 4/120]							
	(a) An electron in an orbital of high angular momentum stays away from the nucleus than an								
	 electron in the orbitals of lower angular momentum. (b) For a given value of the principal quantum number, the size of the orbit is inversely proportionate to the azimuthal quantum number. 								
	(c) Accordin	ig to wave mechanics, the	ground state angular mo	mentum is equal to $\frac{h}{2\pi}$.					
	(d) The plot			2π show peak shifting towards higher					
	value. (1) (b), (c)	(2) (a), (c)	(3) (a), (d)	(4) (a), (b)					

Qua 8.		de Brog	lie wave	elength	of the electron in	n n th Bohr d						
	(a₀ is l (1) 0.7		ius), the	en the va (2) 0.	alue of n/z is: 40	(3) 1.0	[JEE((4) 1. 50	ine (12-0	01-19), 4	ŀ/120j
		ISW	ers									
		EXE	RCIS	E - 1		16.	(B)	17.	(A)	18.	(A)	
	PART - I							EXE	RCIS	E - 3		
1.	(D)	2.	(B)	3.	(C)			F	PART -	• 1		
4.	(A)	5.	(C)	6.	(C)	1.	(B)					
7.	(D)	8.	(A)	9.	(D)	2.		unpaire			0	
10.	(A)	11.	(B)	12.	(D)		V V+	= 3d ³ = 3d ³	-		3 4	
13.	(A)	14.	(D)	15.	(C)		V ²⁺	= 3d ³			3	
	. ,						V ³⁺ V ⁴⁺	= 3d ² = 3d ¹			2 1	
16.	(C)	17.	(D)	18.	(D)		-		-			
19.*	(AB)	20.*	(BC)			3.	√2 .($\left(\frac{11}{2\pi}\right)$	4.	9	5.	6
	PART - II					_	_		_		_	
1.	2					6.	3	(C)	7.	(D)	8.	
2.	1s² 2s² 2p ⁶ 3s² 3p ⁶ 4s² 3d ¹⁰ 4p ⁶ 5s² 4d ¹⁰ 5p ⁶ 6s²				9.	(A)		10.	(D)			
3.	3p, 5d	, 4p, 2s,	4d					P	ART -	II		
4.	(a) 0,	(b) <u> </u>	n,	(c) $\frac{2I}{\pi}$	<u>1</u>	JEE(MAIN) OFFLINE PROBLEMS						
	(u) 0,	(5) √2	<u>2</u> π',	(σ) π		1.	(1)	2.	(1)	3.	(1)	
5.	•	sible se , and (vi	•	uantum	numbers are	4.	(4)	5.	(2)	6.	(3)	
6.	(i) + 5	/2 or – :	5/2, spir	n magne	tic moment =	7.	(2)	8.	(1)	9.	(1)	
	· · ·	B.M. (ii)	-	-		10.	(3)	11.	(1)	12.	(3)	
		EXE	RCIS	E - 2		13.	(1)	14.	(2)	15.	(2)	
1.	(C)	2.	(A)	3.	(A)	16.	(1)	17.	(3)	18.	(3)	
4.	(A)	5.	(C)	6.	(D)		JEE	(MAIN) (ONLINE	PROBL	EMS	
 7.	(A) (D)	8.	(C) (A)	9.	(A)	1.	(1)	2.	(4)	3.	(4)	
						4.	(3)	5.	(3)	6.	(2)	
10. 3	(A) (D)	11. 14	(C)	12. 15	(D) (B)	7.		answer	was (3),	but corr	ect ans	wer is
3.	(D)	14.	(C)	15.	(B)		(2).					

8. (1)

HINTS & SOLUTIONS

EXERCISE - 1

- 1. $\frac{\lambda_1}{\lambda_2} = \sqrt{\frac{V_2}{V_1}} = \sqrt{\frac{200}{50}} = \frac{2}{1}$.
- **2.** $\lambda = \frac{h}{mv} = 1.33 \times 10^{-3} \text{ Å}$
- **3.** For an α particle, $\lambda = \frac{0.101}{\sqrt{V}}$ Å.
- 4. $\Delta X \ \Delta P \cong \frac{h}{4\pi}$ $m(\Delta X \ \Delta V) = \frac{h}{4\pi} \implies m = 0.099 \text{ Kg}$
- 5. An electron has particle and wave nature both.
- 6. $\Delta X \ \Delta P \ge \frac{h}{4\pi}$ $\Delta X \to 0 \Rightarrow \Delta P \to \infty$
- 7. $\lambda = \frac{h}{mv} \Rightarrow \lambda \propto \frac{1}{m}$
- 8. Orbital angular momentum = $\sqrt{\ell} (\ell + 1) \frac{h}{2\pi} = 0$. ∴ □ = 0 (s orbital).
- **9.** Cu : 1s²2s²2p⁶3s²3p⁶3d¹⁰4s¹.
 - \therefore Cu²⁺: 1s²2s²2p⁶3s²3p⁶3d⁹ or [Ar]3d⁹.
- **10.** Magnetic moment = $\sqrt{n(n+2)} = \sqrt{24}$ B.M.
 - ∴ No. of unpaired electron = 4. X₂₆ : 1s² 2s²2p⁶3s²3p⁶3d⁶4s².
 To get 4 unpaired electrons, outermost configuration will be 3d⁶.
 ∴ No. of electrons last 2 (from 4a²)
 - \therefore No. of electrons lost = 2 (from 4s²).
 - ∴ n = 2.
- In Zn²⁺: [Ar] 3d¹⁰ (0 unpaired electrons).
 Fe²⁺: [Ar] 3d⁶ (4 unpaired electrons) maximum.
 Ni³⁺: [Ar] 3d⁷ (3 unpaired electrons).
 Cu⁺: [Ar] 3d¹⁰ (0 unpaired electrons).

12. d⁷ : 3 unpaired electrons.

$$\therefore$$
 Total spin = $\pm \frac{n}{2} = \pm \frac{3}{2}$

- **13.** X_{23} : $1s^2 2s^2 2p^6 3s^2 3p^6 3d^3 4s^2$. No. of electron with $\Box = 2$ are 3 (3d³).
- 14. Cl₁₇⁻ : [Ne] 3s² 3p⁶. Last electron enters 3p orbital.
 ∴ □ = 1 and m = 1, 0, -1.
- **15.** Number of radial nodes = $n \Box 1 = 1$, n = 3. $\therefore \qquad \Box = 1$. Orbital angular momentum
- $= \sqrt{\ell \, (\ell+1)} \, \frac{h}{2\pi} \, = \, \sqrt{2} \, \frac{h}{2\pi} \, .$
- Cl₁₇ : [Ne] 3s² 3p⁵.
 Unpaired electron is in 3p orbital.
 ∴ n = 3, □ = 1, m = 1, 0, -1.
- **17.** Only Spin quantum number (s) is not derived from Schrodinger wave equation.
- **18.** n = 4, m = -3 ∴ only possible value of □ is 3. ∴ Orbital angular momentum = $\sqrt{\ell(\ell+1)} \frac{h}{2\pi} = \frac{2\sqrt{3}h}{2\pi} = \frac{\sqrt{3}h}{\pi}$.
- **19.** Only (A) and (B) arrangements follow Hund's rule.
- **20.** (B) If the electronic configuration of ₆C is written as 1s⁶, then Pauli exclusion principle has been violated.
 - (D) The $+\frac{1}{2}$ and $-\frac{1}{2}$ values of spin quantum number denote two quantum mechanical spin states, which have no classical analogue.

PART - II

1. Ni Atomic No : 28 Ni : [Ar] $3d^8 4a^2$; Ni²⁺ [Ar] $3d^8 4s^0$

No. of unpairecd electron = 2

Quantum Number & Electronic Configuration 2. Atomic No. 56 Electronic configuration: 1s² 2s² 2p⁶ 3s² 3p⁶ 4s² 3d¹⁰ 4p⁶ 5s² 4d¹⁰ 5p⁶ 6s². 3. (a) n = 3 , □ = 1 \Rightarrow 3p (b) n = 5 , □ = 2 \Rightarrow 5d (c) n = 4 , □ = 1 \Rightarrow 4p (d) n = 2 , □ = 0 \Rightarrow 2s \Rightarrow 4d (e) n = 4 , □ = 2 Orbital angular momentum = $\sqrt{\ell(\ell+1)} \frac{h}{2}$ 4. For 4s orbital, $\Box = 0$ Angular momentum = $\sqrt{0(0+1)} \frac{h}{2\pi} = 0$ For 3p orbital, $\Box = 1$ Angular momentum = $\sqrt{1(1+1)} \frac{h}{2\pi} = \frac{h}{\sqrt{2\pi}}$. *:*.. For 4th orbit, Angular momentum $=\frac{nh}{2\pi}=\frac{4h}{2\pi}=\frac{2h}{\pi}.$

- 5. (i) $\Box = 0 \implies m = 0 \ (m \neq 1)$ (iii) $n = 1 \implies \Box = 0 \ (\Box \neq 2)$ (vi) $s = + 1/2 \ or - 1/2 \ (s \neq 0)$
- 6. (i) $_{26}Fe^{3+}$: $1s^2 2s^2 2p^6 3s^2 3p^6 3d^5$

It contains 5 unpaired electrons \therefore n = 5

 $\therefore \text{ Total spin} = \pm \frac{n}{2} = \pm \frac{5}{2}$

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

$$=\sqrt{5(5+2)} = \sqrt{35}$$
 BM.

(ii) 29Cu⁺ : 1s² 2s² 2p⁶ 3s² 3p⁶ 3d¹⁰ It contains 0 unpaired electron

- \therefore Total spin = 0.
- \therefore Spin magnetic Moment = 0.

EXERCISE - 2

- 1. $\Delta p.\Delta x = \frac{h}{4\pi}$ $\Rightarrow \Delta x = \frac{6.62 \times 10^{-34}}{4 \times 3.14 \times 1 \times 10^{-5}} = 5.27 \times 10^{-30} \text{ m.}$
- **2.** $\lambda = \frac{h}{mv} = 0.4 \times 10^{-33} \text{ cm}$

3.
$$\Delta x \cdot \Delta p \approx \frac{h}{4\pi} \Rightarrow \Delta v = 3.499 \times 10^{-24} \text{ ms}^{-1}$$

- 4. s orbital is spherical so non-directional.
- The lobes of d_{xy} orbital are at an angle of 45° with X and Y axis. So along the lobes, angular probability distribution is maximum.
- 6. Total number of electrons in an orbital=2(2□+1).
 The value of □ varies from 0 to n 1.
 ∴ Total numbers of electrons in any orbit

$$= \sum_{\ell = 0}^{\ell = n - 1} 2(2\ell + 1) \, .$$

7. Spin quantum number does not comes from Schrodinger equation.

 $s = +\frac{1}{2}$ and $-\frac{1}{2}$ have been assigned arbitrarily.

- 8. For 1s, 3s, 3d and 2p orbital, $\Box = 0, 0, 2, 1$ respectively. Orbital angular momentum = $\sqrt{\ell (\ell + 1)} \Box$.
- **9.** After np orbital, (n + 1) s orbital is filled.
- **10.** I : For n = 5, I_{min} = 0. ∴ Orbital angular momentum $= \sqrt{\ell(\ell + 1)} \hbar = 0.$ (False) II : Outermost electronic configuration $= 3s^1 \text{ or } 3s^2.$ ∴ Possible atomic number = 11or 12 (False). III : Mn₂₅ = [Ar] 3d⁵ 4s². ∴ 5 unpaired electrons.
 - \therefore Total spin = $\pm \frac{5}{2}$ (False).

IV : Inert gases have no unpaired electrons. ∴ spin magnetic moment = 0 (True).

- **11.** The lobes of $d_{x^2-y^2}$ orbital are alligned along X and Y axis. Therefore the probability of finding the electron is maximum along x and y-axis.
- 12. n = 4, m = 2 Value of □ = 0 to (n - 1) but m = 2.
 ∴ □ = 2 or 3 only
 - $\Box = 2 \text{ or } 3 \text{ only}$ Value of s may be +1/2 or - 1/2.

- **13.** (A) ₂₄Cr : [Ar]3d⁵4s¹
 - (B) m = $-\Box$ to $+\Box$ through zero.
 - (C) ${}_{47}$ Ag : $1s^22s^22p^63s^23p^64s^23d^{10}4p^65s^14d^{10}$. Since only one unpaired electron is present.
 - \therefore 23 electrons have spin of one type and 24 of the opposite type.
- **14.** Two unpaired electrons present in carbon atom are in different orbitals. So they have different magnetic quantum number.
- **15.** Electronic configuration of Zn^{2+} ion is $1s^2 2s^2 2p^6$ $3s^2 3p^6 3d^{10}$ so no electron in 4s orbital.

16.
$$\sqrt{s(s+1)} \frac{h}{2\pi} = \sqrt{\frac{1}{2} (\frac{1}{2}+1)} \frac{h}{2\pi}$$

- $=\frac{\sqrt{3}}{2}\frac{h}{2\pi}=0.866\frac{h}{2\pi}$
- **17.** For principle quantum number n $\Box = 0$ to (n - 1) and $m = -\Box$ to \Box including zero.
- 18. Factual.

EXERCISE - 3

PART - I

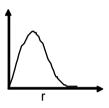
- 1. Orbital angular momentum = $\sqrt{\ell(\ell+1)} \frac{h}{2\pi} = 0$ (since $\Box = 0$ for s orbital).
- For 2p, □ = 1
 ∴ Orbital angular momentum

$$= \sqrt{\ell (\ell+1)} \frac{h}{2\pi} = \sqrt{2} \cdot \left(\frac{h}{2\pi}\right).$$

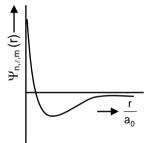
So, electrons with spin quantum number = $-\frac{1}{2}$ will be 1 + 3 + 5 = 9.

5. n = 4, m₀ = 1, -1 Hence \Box can be = 3, 2, 1 i.e. H_f ; 2 orbitals H_d ; 2 orbitals H_p ; 2 orbitals Hence total of 6 orbitals, and we want $m_s = -\frac{1}{2}$, that is only one kind of spin. So, 6 electrons.

- 6. Energy order of orbitals of H is decided by only principle quantum number (n) while energy order of H⁻ is decided by (n + □) rule: Electronic configuration of 'H⁻' is 1s² its Energy order is decided by n+□ rule. H⁻ = 1s²2s⁰2p⁰ Its 2nd excited state is 2p and degenery 2p is '3'.
- 7. For 1s electron in Hatom, plot of radial $4\pi r^2 R^2$ probability function $(4\pi r^2 R^2)$ V/s r is as shown :



- 8. s-orbital is non directional so wave function will be independent of $\cos \theta$.
- **9.** For 2s orbital no. of radial nodes = $n \Box 1 = 1$.



10. For 1s orbital Ψ should be independent of θ , also it does not contain any radial node.

$$\frac{\mathsf{E}_4 - \mathsf{E}_2}{\mathsf{E}_6 - \mathsf{E}_2} = \frac{\frac{\mathsf{E}_1}{16} - \frac{\mathsf{E}_1}{4}}{\frac{\mathsf{E}_1}{36} - \frac{\mathsf{E}_1}{4}} = \frac{-\frac{3\mathsf{E}_1}{16}}{-\frac{8\mathsf{E}_1}{36}} = \frac{3 \times 36}{8 \times 16} = \frac{27}{32}$$

PART - II JEE(MAIN) OFFLINE PROBLEMS

 Mn²⁺ has the maximum number of unpaired electrons (5) andtherefore has maximum moment.

2.
$$\Delta x.\Delta v = \frac{h}{4\pi m}$$
$$\Delta v = \frac{6.6 \times 10^{-34}}{4 \times 3.14 \times 25 \times 10^{-5}}$$
$$\therefore \quad \Delta v = 2.1 \times 10^{-18} \text{ ms}^{-1}.$$

- 3. $\lambda = \frac{h}{mv} = \frac{6.63 \times 10^{-34} \times 1000}{60 \times 10}$ = 11.05 x 10⁻³⁴ = 1.105 x 10⁻³³ metres.
- 4. ${}_{26}Fe = 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 3d^6, 4s^2$ $Fe^{++} = 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 3d^6$ The number of d -electrons retained in Fe²⁺ = 6. Therefore, (4) is correct option.
- The value of □ (azimuthal quantum number) for s -electron is equal to zero.

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

Substituting the value of I for s-electron

$$=\sqrt{0(0+1)} \cdot \frac{h}{2\pi} = 0$$

- 6. For 4f orbital electrons, n = 4 $\Box = 3 \text{ (because } \underset{0 \ 1 \ 2 \ 3}{\overset{\text{s p d } f}{\underset{1 \ 2 \ 3}}}$ m = + 3, + 2, + 1, 0, -1, -2, -3 s = + 1/2.
- 7. ${}_{24}Cr \rightarrow 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 3d^5, 4s^1$ $\Box = 1, \Box = 1, \Box = 2$ (we know for p, $\Box = 1$ and for d, $\Box = 2$). For $\Box = 1$, total number of electrons = 12

For $\Box = 2$, total number of electron = 5.

8. The electron having same principle quantum number and azimuthal quantum number will be the same energy in absence of magnetic and electric field.
(iv) n = 3, l = 2, m = 1
(v) n = 3, l = 2, m = 0

have same n and I value.

- **9.** For hydrogen the energy order of orbital is 1s < 2s = 2p < 3s = 3p = 3d < 4s = 4p = 4d = 4f
- 10. According to Heisenberg's uncertainity principle

$$\Delta x \times \Delta p = \frac{h}{4\pi}$$
$$\Delta x \times (m.\Delta v) = \frac{h}{4\pi}$$
$$\Rightarrow \Delta x = \frac{h}{4\pi m.\Delta v}$$
Here, $\Delta v = \frac{0.001}{100} \times 300 = 3 \times 10^{-3} \text{ ms}^{-1}$

- $\Delta x = \frac{6.63 \times 10^{-34}}{4 \times 3.14 \times 9.1 \times 10^{-31} \times 3 \times 10^{-3}}$ $= 1.29 \times 10^{-2} \text{m.}$ 11. $_{28}\text{Ni} \rightarrow [\text{Ar}]3d^8 4s^2$ $\boxed{11111111} \qquad \boxed{11} \qquad$
- energy. n + l = 3 + 0 = 3
 - n + l = 3 + 1 = 4 n + l = 3 + 2 = 5 (highest energy) n + l = 4 + 0 = 4
- **13.** As $\lambda = \frac{h}{mv} = \frac{6.63 \times 10^{-34}}{1.67 \times 10^{-27} \times 1 \times 10^3}$ = 3.97 × 10⁻¹⁰ M = 0.397 × 10⁻⁹ M = ~ 0.40 nm.
- 14. $\Delta x \times \Delta P = \frac{h}{4\pi}$ $\Delta x \times [m\Delta v] = \frac{h}{4\pi}$ $\Delta v = \frac{600 \times 0.005}{100} = 0.03$ So $\Delta x [9.1 \times 10^{-31} \times 0.03] = \frac{6.6 \times 10^{-34}}{4 \times 3.14}$ $\Delta x = \frac{6.6 \times 10^{-34}}{4 \times 3.14 \times 9.1 \times 0.03 \times 10^{-31}} = 1.92 \times 10^{-3} M.$
- 15. (a) 4 p (b) 4 s (c) 3 d (d) 3 p
 Acc. to (n + □) rule, increasing order of energy (d) < (b) < (c) < (a)
- **16.** Z = 37. Rb is in fifth period. [Kr]5s¹ is its configuration. So n = 5, l = 0, m = 0, s = $+\frac{1}{2}$ or $-\frac{1}{2}$ **17.** K.E. = eV

$$\Rightarrow \lambda = \frac{h}{\sqrt{2meV}}$$
$$\Rightarrow \frac{h}{\lambda} = \sqrt{2meV}$$

18. R = 0.529 $\frac{n^2}{z}$ Å = 0.529 $\frac{2^2}{1}$ Å = 2.12 Å

JEE(MAIN) ONLINE PROBLEMS

- 2. Following Aufbau principle for filling electrons.
- **3.** De-broglie wavelength (for particles) = $\frac{h}{\sqrt{2m \text{ KE}}}$

As temperature is same, KE is same.

So,
$$\lambda \propto \frac{1}{\sqrt{m}}$$
.

Hence λ_{db} (electron) > λ_{db} (neutron)

4. n = 5

Possible subshell are

- \Rightarrow 5s, 5p, 5d, 5f, 5g
- ∴ Total number of orbital = 1 + 3 + 5 + 7 + 9 = 25
- 5. NaF: Na⁺ = $1s^22s^22p^6$ F⁻ = $1s^22s^22p^6$

6.
$$2\pi r = n\lambda$$
 $\lambda = \frac{2\pi r}{n} = \frac{2\pi \times 0.529 \text{ Å}}{1}$

7.
$$mvr = \frac{nh}{2\pi}$$

According to wave mechanics, the ground state angular momentum is equal to $\frac{h}{2\pi}$.

8.
$$2\pi r = n\lambda$$

$$2\pi a_0 \frac{n^2}{Z} = n\lambda$$

$$2\pi a_0 \frac{n^2}{Z} = n1.5\pi a_0$$

$$\frac{n}{Z} = \frac{1.5}{2} = \frac{3}{4} = 0.75$$