THE SOLID STATE

IMPORTANT FORMULAE

1. Density of unit cell (d) =
$$\frac{\text{Mass of unit cell}}{\text{Volume of unit cell}} = \frac{z \times M}{a^3 \times N_4}$$

Table 1.1: Different Parameters of Cubic System

Unit cell	No. of atoms per unit cell	Distance between nearest neighbour (d)	C.N.	Radius (r)
Simple cubic	1	а	6	<u>a</u> 2
Face-centred cubic	4	$\frac{a}{\sqrt{2}}$	12	$\frac{a}{2\sqrt{2}}$
Body-centred cubic	2	$\frac{\sqrt{3}}{2}a$	8	$\frac{\sqrt{3}}{4}a$

Volume occupied by atoms in unit cell (v) $\times 100$ 3. Packing efficiency = Total volume of the unit cell (V)

Table 1.2: Packing efficiency of different crystals

S.No.	Crystal system	Packing efficiency
(<i>i</i>)	Simple cubic	52.4%
(ii)	Body-centred cubic	68%
(iii)	Face-centred cubic	74%
(iv)	Hexagonal close-packed	74%

Radius of the cation $= r^+$ 4. Radius ratio = Radius of the anion

Table 1.3: Structural arrangement of different radius ratios of ionic solids

Radius ratio (r^+/r^-)	Possible coordination number	Structural arrangement	Examples
0.155 - 0.225	3	Trigonal planar	$\mathrm{B_2O_3}$
0.225 - 0.414	4	Tetrahedral	ZnS, SiO ₄ ^{4–}
0.414 - 0.732	6	Octahedral	NaC1
0.732 - 1.0	8	Body-centred cubic	CsCl

- 5. If R is the radius of the spheres in the close packed arrangement, then
 - (i) Radius of octahedral void, r = 0.414 R
 - (ii) Radius of tetrahedral void, r = 0.225 R
- 6. In a close packed arrangement:
 - (i) Number of octahedral voids = Number of atoms present in the close packed arrangement.
 - (ii) Number of tetrahedral voids = $2 \times$ Number of atoms present in the close packed arrangement.

MULTIPLE CHOICE QUESTIONS

Choose and write the correct option in the following questions.

1.	. Which of the following conditions favours the existence of a substance in the solid state?					
	(a) High temperature		(b) Low temperature	[NCERT Exemplar]		
	(c) High thermal ener		(d) Weak cohesive fo			
9		Desired to the second s				
2.	(a) Definite and chara		stic of a crystalline solid?	[NCERT Exemplar]		
	(b) Isotropic nature.	eteristic freat of rusio.	11.			
	1224 N	ally repeated pattern	of arrangement of constit	uent particles in the entire		
	crystal.	un, repetited pittern	or arrangement or consti	de particies in the chare		
	(d) A true solid					
3.	Which of the followin	g is true about the va	alue of refractive index of	f quartz glass?		
				[NCERT Exemplar]		
	(a) Same in all direction	ons	(b) Different in differ	rent directions		
	(c) Cannot be measure	ed	(d) Always zero			
4.	Which of the followin	g statement is not tr	ue about amorphous solic	ls? [NCERT Exemplar]		
	(a) On heating they m	nay become crystalline	e at certain temperature.			
	(b) They may become	crystalline on keepin	g for long time.			
	(c) Amorphous solids	can be moulded by h	eating.			
	(d) They are anisotrop	pic in nature.				
5.			ids is due to	The state of the s		
	(a) a regular arrangement of constituent particles observed over a short distance in the crystal					
	lattice. (b) a regular arrangement of constituent particles observed over a long distance in the crystal					
	(b) a regular arranger lattice.	nent of constituent p	oarticles observed over a i	ong distance in the crystar		
	(c) same arrangement	of constituent partic	les in different directions.			
	(d) different arrangen	nent of constituent pa	rticles in different directi	ons.		
6.	Which one of the follo	owing is a covalent c	rystal?			
	(a) Rock salt	(b) Ice	(c) Quartz	(d) Dry ice		
7.	Which of the followin	g solids is not an ele				
	(i) Mg (s)		(ii) TiO (s)			
	$(iii) I_2(s)$	715 715	(iv) $\mathbf{H}_2\mathbf{O}(s)$	7 15 2**5 2***5 1 7** 5		
	(a) (i) only	(b) (ii) only	(c) (iii) and (iv)	(d) (ii) , (iii) and (iv)		
8.	The number of atoms		W V 200	(1) C		
0	(a) 2	(b) 4	(c) 5	(d) 6		
9.				ions and atom B occupies the face centred points, the		
	formula of the compo		is missing from one of the	re face centifed points, the		
	(a) AB_2	$(b) A_2B_3$	$(c) A_2B_5$	$(d) A_2 B$		
10.	The total number of to	etrahedral voids in t	he face centred unit cell i	is		
	(a) 6	(b) 8	(c) 10	(d) 12		
11.	If Z is the number of a	toms in the unit cell	that represents the closes	st packing sequence—ABC		
	ABC, the number of to	etrahedral voids in t	he unit cell is equal to			
	(a) Z	(b) 2Z	$(c) \frac{Z}{9}$	$(d) \frac{Z}{4}$		

12.	In which pair most ef	ficient p	acking	g is pr	esent?						
	(a) hcp and bcc			9 1) hep a	ind ccp	ĺ			
	(c) bcc and ccp				36	, M 155M		ple cu	bic ce	ell	
13	The fraction of the to	tal volu	ne occ	unied	1000	<i>5</i> /					
10.	(a) $\pi/4$	$(b) \pi/6$	_	upicu		$\pi/(3^{\circ})$	76.23	nt m a		$\pi/(4\sqrt{2})$	
14	A metallic crystal has			aking	20		W.	ontoro	(S) (A)		ittico i
14.	empty space?	the occ	type st	aking	patteri	1. W 11a	ii perc	emage	OI VO	nume of tims to	nuice i
	(a) 68%	(b) 32			N.	26%			N 6	74%	
15.	A solid compound X		Cl str	ucture	. If the	radiu	s of th	e catio	n is	100 pm, the ra	dius o
	the anion (Y ⁻) will be		0 -		. 21.		2		2.35	165 7	
	(a) 275.1 pm	(b) 32	(S) 84 2 6		A1-50.	241.			330 6	165.7 pm	
16.	AB crystallizes in a	The state of the s					The second second	200	'a' e	qual to 387 pr	n. The
	distance between two								(1)	995	
	(a) 250 pm										
17.	If a stands for the edg centred cubic, then the	-			1 22 - 24						
	(a) $\frac{1}{2}a:\frac{\sqrt{3}}{4}a:\frac{1}{2\sqrt{2}}a$	$(b) \frac{1}{2}$	a:√3 a:	$\frac{1}{\sqrt{2}}a$	(4	(c) $\frac{1}{2}a$:	$\frac{\sqrt{3}}{2}a$:	$\frac{\sqrt{2}}{2}a$	(d)	$1a:3a:\sqrt{2a}$	
10	Which of the fellowing		la dos	~ * • • •	عذماذ ماده	Enouale	al dafa	240			
10.	Which of the following							ectr	(4)	728	
10	(a) AgBr								(a)	ZnS	
19.	The type of crystal de	efect is i									
			Na ⁺	CI-	Na ⁺	C1-	Na ⁺	Cl ⁻			
			Cl ⁻	0	CI ⁻	Na ⁺	0	Na ⁺			
			Na ⁺	CI ⁻	0	Cl ⁻	Na ⁺	CI ⁻			
			Cl-	Na ⁺	Cl-	Na ⁺	0	Na ⁺			
	(a) Frenkel defect				(ł) Fren	kel an	d Scho	ttkv o	lefect	
	(c) Interstitial defect				108.1	l) Scho					
20.	Which of the followin	o is ton	abou	thoc	90 X 00	K1			comi	conductors?	
40.	(a) Positive	ig is true	abou	tille C	7 (d)) Neu	- 10 E	-type	Senin	conductors.	
	(c) Negative					.6		n conc	entra	tion of p impu	rity
0.1		* * * * * * * * * * * * * * * * * * *		• • • •	S.N.3	i) bep	ciras o	n conc	ciitia	tion or p impu	Tity
21.	Which among the foll			nisotr		.\ (*!			7.1	C - 1! 1.1	
	(a) Wood	(b) Su				c) Glas			(a)	Sodium chlor	iae
22.	In hcp arrangement,	Constitution Constitution	dinatio	n nun	779-		atom i	S	7.1	1.0	
	(a) 4	(b) 6			3	2) 8			(d)	12	
23.	The number of tetrah	1000 at 1500 1500 1500 1500 1500 1500 1500 150	ids in	a unit			ructur	e			
	(a) 4	(<i>b</i>) 6			(6	c) 8			(d)	10	
24.	The arrangement AB	CABC	refers	to							
	(a) Hexagonal close p	acking			(b)) Tetr	ahedra	al close	pack	ing	
	(c) Cubic close packin	\mathbf{g}			(a	l) Octa	hedra	l close	packi	ng	
25.	Which one of the foll	owing is	an an	orpho	us soli	id?					
	(a) Diamond		aphite			c) Com	mon s	alt	(d)	Glass	
26.	Which of the followin	g is not	the pr	operty	of cry	stallin	e solic	ds?	180 M		
W.	(a) Isotropy	0	F	1					int		
	(c) Strong intermolec	ular for	ces			(b) Sharp melting point(d) A definite and regular geometry					

27.	In a ccp arrangement is	of spheres in three dime	ensions, the coordination	on number of each sphere
	(a) 3	(b) 6	(c) 9	(d) 12
28.	The contribution of a	n atom at the edge centr	re of a unit cell is	
	(a) $\frac{1}{8}$	$(b) \frac{1}{4}$	(c) $\frac{1}{2}$	(d) 1
29.	The closest packing s	equence ABAB repres		
	(a) primitive cubic pa	cking	(b) hexagonal packing	g
12.27	(c) fee packing		(d) bee packing	
30.		hedral void relative to tl	he radius of the sphere $(c) \ 0.414$	s in a close packing is (d) 1.225
21	(a) 1.414	(b) 0.225		(a) 1.223
31.	(a) $0.732 - 1.0$	gement, the radius ratio (b) 0.225 - 0.414	(c) $0.414 - 0.732$	(d) $0.155 - 0.225$
32.		owing is a covalent solid		(, , , , , , , , , , , , , , , , , , ,
	(a) Fe	(b) Diamond	(c) Cu	(d) NaCl
33.	Which one of the follo	owing solids is very soft	in nature?	
	(a) Ionic solid	(b) Covalent solid	(c) Metallic solid	(d) Molecular solid
34.	The existence of a sul	bstance in more than on	e solid modification is	known as
	(a) isomorphism	(b) polymorphism	(c) amorphism	(d) none of these
35.	Words SW World Participant	g compounds is not isot	M. CONTROL SECTION	(1) D' 1
2.0	(a) Rubber	(b) Glass	(c) PVC	(d) Diamond
36.	(a) n-type	nductor is developed by $(b) p$ -type	(c) intrinsic	(d) $n-p$ type
37		g substance will be capa		
01.	(a) Diamond	(b) Graphite	(c) Carborundum	(d) NaCl
38.	* 1	hold iodine molecules		
	(a) London force		(b) dipole-dipole inte	raction
20	(c) Coulombic force		(d) Covalent bond	
39.	Bravais lattices are of (a) 7 types	(b) 10 types	(c) 17 types	(d) 14 types
40.		doped with MgCl ₂ , the		100 N
10.	(a) Interstitial	(b) Schottky	(c) Frenkel	(d) Impurity
41.	Which of the followin	g is a pseudo solid?		
	(a) CaF_2	(b) Glass	(c) NaCl	(d) All of the above
42.	The number of NaCl	molecules in the unit ce	ell of the crystal is	
	(a) 2	(b) 4	(c) 6	(d) 8
43.		kel defects in a crystal _	CONTRACTOR DATE	
	(a) decreases.		(b) increases.	1
4.4	(c) does not change.	41 '4 11 C ' 1	(d) either increase or	decrease.
44.	Number of particles i	n the unit cell of simple (b) 2	(c) 3	(d) 4
15			X. Z	(X)
43.	(a) 1	toms per unit cell of a f	(c) 3	(d) 4
46.		And were	Y	mass is 6.94 g mol^{-1} . The
	edge-length of a unit	cell of $(N_A = 6.02 \times 10^2)$	³ mol ⁻¹) lithium metal	is-
	(a) 527 pm	(b) 264 pm	(c) 154 pm	(d) 352 pm

47	The ionic radii of A+	and R ⁻ ions are 0.98 X	10 ⁻¹⁰ m and 1.81 × 10	0 ⁻¹⁰ m. The coordination
11.	number of each ion in		io mana noi x n	o in the coordination
	(a) 8	(b) 2	(c) 6	(d) 4
48.	The correct statement	regarding defects in cr	ystalline solid is	
	(a) Frenkel defects dec	crease the density of crys	stalline solids.	
	(b) Frenkel defect is a	dislocation defect.		
	(1) (4)	und in halides of alkali i		
		ive no effect on the dens	er en lees sterv	
49.			0 0	th of 361 pm. If there are
	(a) 80 pm	ne unit cell, then the race (b) 108 pm	(c) 40 pm	(d) 128 pm
50			A150 87 5 \$2.	The density of the metal
30.	is 2.72 g cm ⁻³ . The mo	olar mass of the (N, Ava	gadro's constant = 6.05	$2 \times 10^{23} \text{ mol}^{-1}$) metal is
		(b) 20 g mol^{-1}		
51.	Structure of a mixed	oxide is cubic close pac	ked (ccp). The cubic u	nit cell of mixed oxide is
				pied by divalent metal A
		The state of the s		e formula of the oxide is
-0		(b) A_2BO_2		The state of the s
52.	diameter of the metal		lattice. The edge of the	e unit cell is 408 pm. The
		(b) 408 pm	(c) 144 pm	(d) 204 pm
53.	- A - A	dral voids per atom pre	W A1	
	(a) 1	7.		4.
54.				gth of the side of the unit
	The state of the s	om, the atomic radius o	and the second s	
	(a) 151.8 pm	(b) 75.5 pm	(c) 300.5 pm	(d) 240.8 pm
55.			attice with a unit cell le	ength of 334 pm. What is
	the radius of copper a		/ \ 100	/ 1 \ 110
~ ~	(a) 157	(b) 181	(c) 108	(d) 118
50.	semiconductor?	e following elements s	ilicon should be dope	d so as to give p-type of
	(a) Selenium	(b) Boron	(c) Germanium	(d) Arsenic
57.	If NaCl is doped with	10 ⁻⁴ mol% of SrCl ₂ , the	concentration of cation	n vacancies will be
		(b) $6.02 \times 10^{17} \mathrm{mol}^{-1}$		
58.	The fraction of total v	olume occupied by aton	ns present in a simple o	cube is
	(a) $\frac{\pi}{3\sqrt{2}}$	$(b) \frac{\pi}{4\sqrt{2}}$	(c) $\frac{\pi}{4}$	$(d) \frac{\pi}{6}$
	560 KH (1 160 0 HH)	5-730 Extends 11		O.
59.	- 10 AM AM AM AM AM AM AM AM	lour in solid alkali meta		lue to
	(a) Interstitial position	IS	(b) F-centres	
CO	(c) Schottky defect		(d) Frenkel defect	1. '- 426 6 C' 41-4
60.				h is 436.6 pm. Given that number being 6.02 × 10 ²³
	mol ⁻¹ , the density of C		ov allia alla livogadio i	tuniour deing dida in 10
		(b) 42.5 g/cm^3	(c) 0.425 g/cm^3	$(d) 8.25 \text{ g/cm}^3$
61.	A compound formed	by elements X and Y o	crystallises in a cubic s	structure in which the X
		ers of a cube and the Y a	toms are at the face-ce	ntres. The formula of the
	compound is	(h) V V	(c) VV	(d) V V
	$(a) XY_3$	$(b) X_3Y$	(c) XY	$(d) XY_2$

62.						
	surface of the HCP unit cell. The value of $\frac{C}{a}$ is					
	(a) $\sqrt{\frac{2}{3}}$	$(b) \sqrt{\frac{8}{3}}$	(c) $\sqrt{\frac{6}{3}}$	$(d) \sqrt{\frac{7}{3}}$		
63.			of the following stateme	ent given below is correct?		
	(a) Graphic has highe	r than diamond. er electrical conductivity	than diamond			
		er thermal conductivity				
	Section 2 Annual	r C—C bond order than				
64.		atement for cubic close	100 Page 100			
		eighbours of an atom pre	170 A	er is 12.		
	53 M (2007)	he atom packing is 74%. tahedral and tetrahedra		nd 2 respectively.		
	\$3 10	length is $2\sqrt{2}$ times the		1		
65.	· A Company of the Co			hat $r_{\text{Na}^+}/r_{\text{Cl}^-} = 0.55$ and		
	$r_{\text{K}^+}/r_{\text{Cl}^-} = 0.74$. The (a) 1.123	ratio of the side of the u (b) 0.414	nit cell of KCl to that o (c) 0.891	f NaCl is (d) 1.414		
66.	Experimentally it was	s found that a metal oxic	le has formula M _{0.08} O.	Metal M is present as M ²⁺		
	and M ³⁺ in its oxide.	Fraction of the metal w	hich exists as M ³⁺ wou	ld be		
67	(a) 5.08%	(b) 7.01%	(c) 4.08%	(d) 6.05%		
07.	The crystal with meta (a) NaCl	(b) FeO	(c) KCl	(d) ZnO		
68.		0 0	-	The density of the metal		
		olar mass of the metal is				
		tant = $6.02 \times 10^{23} \text{ mol}^{-1}$ (b) 30 g mol^{-1}		(d) 20 g mol^{-1}		
69.	CsCl crystallises in	body-centered cubic lat	93 AM AMERICA	length then which of the		
	following expression		34	$\sqrt{3}$		
	(a) $r_{\text{Cs}^+} + r_{\text{CI}^-} = \sqrt{3} a$	(b) $r_{\text{Cs}^+} + r_{\text{Cl}^-} = 3a$	(c) $r_{\text{Cs}^+} + r_{\text{Cl}^-} = \frac{3a}{2}$	(d) $r_{\text{Cs}^+} + r_{\text{Cl}^-} = \frac{\sqrt{3}}{2}a$		
70.			cubic lattice with a un	it cell edge of 4.29 Å. The		
	radius of sodium met (a) 5.72 Å	(b) 0.93 Å	(c) 1.86 Å	(d) 3.22 Å		
71.				ngth of the unit cell is 'a',		
		between the two atoms i	n the metallic crystal w	70-25		
	(a) $\sqrt{2}a$	(b) $\frac{a}{\sqrt{2}}$	(c) 2a	(d) $2\sqrt{2}a$		
72.	Body centered cubic (a) 8	lattice has a co-ordinati (b) 12	on number of (c) 6	(d) 4		
73.			X 2	0 gcm ⁻³ and cell edge as		
	200 pm is equal to					
7.4	- 10 mm	$(b) 5 \times 10^{24}$	$(c) 1 \times 10^{25}$	$(d) \ 2 \times 10^{25}$		
14.		ottky defects contain in ation and anion vacancie				
	(b) interstitial anions					
	(c) cation vacancies of					
	(d) cation vacancies as	nd interstitial cations.				

75.	Which of the following 2-dimension structure		gonal close packed ar	rangement of spheres in
	(a) ABCABA	(b) ABCABC	(c) ABABA	(d) ABBABB
76.		he packing efficiency in	different types of unit	t cells is
	(a) $fcc < bcc < simple cub$		(b) $fcc > bcc > simple cu$	
	(c) $fcc < bcc > simple cub$	oic	(d) $bcc < fcc > simple cu$	bic
77.	0	hest melting point is ex		
	(a) covalent solids		(b) ionic solids	
	(c) pseudo solids		(d) molecular solids	
78.		-		and atom B occupies the
	formula of the compo		issing from one of the	face-centered points, the
	(a) A_2B	(b) AB ₉	(c) A_2B_3	$(d) A_2 B_5$
79.	27 (3) (372)			crystal of NaCl of mass
	1.00 g is			
	(a) 2.57×10^{21}	$(b) 5.14 \times 10^{21}$	(c) 1.28×10^{21}	$(d) 1.71 \times 10^{21}$
80.	An ionic compound h	as a unit cell consisting	of A ions at the corne	rs of a cube and B ion on
	THE RESERVE AND ADDRESSED TO SERVE AND ADDRES	of the cube. The empir		
	(a) A_3B	(b) AB ₃	(c) A_2B	(d) AB
81.		present in a face-center	red cubic unit cell of a	metal is
	(r is atomic radius)			
	(a) $\frac{20}{3}\pi r^3$	(b) $\frac{24}{3}\pi r^3$	(c) $\frac{12}{3}\pi r^3$	$(d) \ \frac{16}{3} \pi r^3$
82.	The number of unit co	ells that are divided equ	ally in a face-centred o	cubic lattice will be
	(a) 2	(b) 4	(c) 6	(d) 8
83.	A crystalline solid:			
		rom solid to liquid when	heated.	
	(b) has no definite mel	Z NAME CONTROL OF THE PARTY OF	• Those	
		ation of its geometry eas		
0.4	1. 1.	nensional arrangements.		
84.	of atoms present is res		e-centred cubic structur	re, the ratio of the number
	(a) $8:1:6$	(b) $1:2:4$	(c) 4:2:1	$(d) \ 4:2:3$
85.	Na and Mg crystallize	in crystals of bcc and f	30 50	he amount of Na and Mg
		respective crystal unit	-	
	(a) 4 and 2	(b) 9 and 14	(c) 14 and 9	(d) 2 and 4
86.	Ferrous oxide has a c	cubic structure and eac	h unit cell edge is 5.0	A. Assuming the oxide
		ne amount of Fe ²⁺ and C	NO. 0000 0000 0000	
	(a) four Fe^{2+} and four	-	(b) two Fe^{2+} and four	X197
~ -	(c) four Fe ²⁺ and two		(d) three Fe^{2+} and the	ree O
87.		solids shows the Frenke	_	(A) RC1
00	(a) ZnS The total new of weights in	(b) AgBr	(c) AgI	(d) KCl
00.		(b) 3.011×10^{23}		lose-packed structure are
80	N 5	42 10	25 //	hexagonal close packed
03.				mula of the compound is
	(a) C_2A_3	(b) C_3A_2	(c) C_3A_4	(d) C ₄ A ₃

90.	A metal crysta	allises into two	cubic face	s namely face	e centered (fcc) and bod	y centered (bcc),
	and bcc will be		are 3.5 A a	ina 3.0 A resp	becuvery. 11	ie ratio of the	e densities of fo	:c
	(a) 2.1:1	(b) 3	.3:1	(c) 1.2	25 9 : 1	(d) 2.25	59:1	
91.	The total num	ber of atoms p	resent in be	c arrangeme	nt is			
	(a) 2	(b) 4		(c) 6		(d) 1		
92.	The number of	of octahedral v	oids in a un	it cell of ccp s	structure is			
	(a) 2	(b) 3		(c) 4		(d) 6		
93.	The largest vo	oid is						
	(a) triangular			- No. 16	rahedral	(d) octa	hedral	
94.	Close packing		,Ti.;			SOUTH IN		
	(a) fcc	(b) be	CC .	(c) sin	nple cubic	(d) non	e of these	
95.	The formula f	or determinati	on of densi	4. The second se				
	(a) $\frac{a^3N_0}{n\times M}$	(1) -	$n \times M$	$(a) \frac{n}{n}$	$\frac{\times N_0}{\times M}$	a^3	$\times M$	
	(a) $n \times M$	(b) \overline{I}	$\frac{n \times M}{V_0 \times a^3}$	$\frac{(c)}{a^3}$	$\times M$	$(d) \frac{a^3}{N_0}$	$\times n$	
96.	In a crystal co	mposed of n s	pheres, the	total number	of tetrahed	ral voids are		
	(a) n/2	(b) n		(c) 2n		(d) 4n		
97.	Which of the							
	(a) Graphite	72 1990 1997 1997	190 0000	20	dium chlori	de (d) Fe		
98.	The vacant sp	SWEST WAS	ork malane		04	(1) 000		
	(a) 48%	(b) 2		(c) 32		(d) 26%		
99.		gth of the side n the cube will		the distance b	etween the	body center	ed atom and on	le
	15			$\sqrt{\cdot}$	3	$\sqrt{3}$		
	(a) $\frac{2}{\sqrt{3}}a$	(b) -	$\overline{/3}^a$	(c) $\frac{\sqrt{3}}{4}$	-a	$(d) \ \frac{\sqrt{3}}{2}$	a	
nswe	ers							
1. (3. (a)	4. (d)	5. (b)	6. (c)	7. (c)	8. (b)	
9. (11. (b)	12. (b)	13. (b)	14. (b)	15. (c)	16. (d)	
17.		19. (d)	20. (b)	21. (d)	22. (d)	23. (c)	24. (c)	
25.	1.20	27. (d)	28. (b)	29. (b)	30. (c)	31. (b)	32. (b)	
33. (35. (<i>d</i>)	36. (a)	37. (b)	38. (a)	39. (<i>d</i>)	40. (<i>d</i>)	
41. (43. (c)	44. (a)	45. (<i>d</i>)	46. (<i>d</i>)	47. (c)	48. (b)	
49. (50. (a)	51. (<i>d</i>)	52. (a)	53. (a)	54. (<i>a</i>)	55. (<i>d</i>)	56. (b)	
57. (58. (d)	59. (<i>b</i>)	60. (a)	61. (a)	62. (b)	63. (b)	64. (a)	
65. (66. (c)	67. (b)	68. (c)	69. (<i>d</i>)	70. (c)	71. (b)	72. (a)	
73. (b) 74. (a)	75. (c)	76. (<i>b</i>)	77. (<i>b</i>)	78. (<i>d</i>)	79. (a)	80. (<i>d</i>)	

85. (*d*)

93. (b)

84. (*b*)

92. (c)

87. (*d*)

95. (*b*)

86. (*a*)

94. (*a*)

88. (c)

96. (c)

82. (c)

90. (c)

98. (c)

83. (*a*)

91. (*a*)

99. (*d*)

81. (*d*)

89. (c)

97. (b)

CASE-BASED QUESTIONS

1. Read the passage given below and answer the following questions:

Solid is that form of matter which possesses rigidity and hence a definite shape and a definite volume. If intermolecular forces are greater than thermal energy, substances exist as solid. Solids can be classified into two types: Crystalline solids and amorphous solids. Crystalline solids have regular arrangement of particles, definite geometric shapes, sharp melting points and definite heat of fusion. They are anisotropic and undergo clean cleavage. On the other hand,

of t	emperature, no de	finite heat of fusion. Tl	ney are isotropic an	rular shapes, melt over a range d undergo irregular cleavage. e the types of crystalline solids.			
The	following questions	are multiple choice que	stions. Choose the m	ost appropriate answer:			
(i)	the second of the second of the second	ard, electrical insulate mperature. What type		as molten state and melts at			
	(a) Ionic	(b) Molecular	(c) Covalent	(d) Metallic			
(ii)	Which of the follo	wing is a crystalline so	lid?				
	(a) Glass	(b) Sodium chloride	(c) Rubber	(d) Plastic			
		OR					
	The sharp melting point of crystalline solids is due to						
	(a) a regular arrangement of constituent particles observed over a short distance in the crystal lattice.						
	(b) a regular arrang lattice.	gement of constituent p	articles observed ov	er a long distance in the crystal			
	(c) same arrangeme	ent of constituent partic	des in different dire	ctions.			
	(d) different arrang	gement of constituent p	articles in different	directions.			
(iii)	Which of the follo	wing is not a character	istic of a crystalline	e solid?			
	(a) Definite and cha	aracteristic heat of fusic	on.				
	(b) Isotropic nature	2.					
	(c) A regular perio entire crystal.	dically repeated patter	n of arrangement	of constituent particles in the			
	(d) A true solid						

- (iv) Which of the following is true about the value of refractive index of quartz glass?
 - (a) Same in all directions

(b) Different in different directions

(c) Cannot be measured

(d) Always zero

Answers

(i) (c)

(ii) (b) **OR** -(b) (iii) (b)

(iv)(a)

2. Read the passage given below and answer the following questions:

Any departure from a perfectly ordered arrangement of constituent particles is called defect or imperfection. In solids, when the ratio between cations and anions remains the same after defect, it is termed as stoichiometric defects. Schottky and Frenkel defects are categorised into stoichiometric defects. In Schottky defect equal number of cations and anions are missing while in Frenkel defect cations are missing from lattice sites and occupy interstitial sites. In nonstoichiometric defects, the ratio of cations and anions changes as a result of the defect. Metal excess and metal deficiency defects are non-stoichiometric defects.

In these questions (Q. No (i) to (iv)), a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

- (a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
- (b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
- (c) Assertion is correct statement but reason is wrong statement.
- (d) Assertion is wrong statement but reason is correct statement.
- Stoichiometry defects are the point defects that disturb the stoichiometry of (i) Assertion (A): the solid.
 - Stoichiometric defects are also known as thermodynamic defects.
- Schottky and Frenkel defects are stoichiometric defects. (ii) Assertion (A):
 - The ratio between cations and anions remains the same in Schottky and Frenkel defects.
- (iii) Assertion (A): Frenkel defect is not found in pure alkali metal halides.
 - In Frenkel defect, cations are missing from lattice sites.

OR

- **Assertion (A):** Zinc oxide turns yellow on heating.
- Zinc oxide shows metal deficiency defect. Reason (R):
- (iv) Assertion (A): NaCl shows metal excess defect due to anionic vacancies.
 - Sodium loses electron to from Na⁺ ions and the released electrons occupy anionic sites.

Answers

(ii)(a)(iii) (b) **OR** (c) (iv) (a)

3. Read the passage given below and answer the following questions:

A unit cell is the smallest repeating portion of a crystal lattice. The cubic crystal system is composed of these different types of unit cells: Simple cubic, body-centred and face-centred.

In simple cubic unit cell, atoms are only at its corners. So, the total number of atoms in one unit cell is $8 \times \frac{1}{8} = 1$ atom.

In body-centred unit cell, unit cell has an atom at each of its corners and also one atom at its body centre. Thus, total number of atom per unit cell:

(8 corners $\times \frac{1}{9}$ per corner atom) unit cell + 1 body centre atom = 1 + 1 = 2 atoms

In face-centred unit cell, unit cell contains atoms at all the corners and at the centre of all the faces of the cube. Thus, total number of atoms per unit cell:

 $(8 \text{ corners} \times \frac{1}{8} \text{ per corner atom}) + (6 \text{ face centred atoms} \times \frac{1}{2} \text{ per unit cell}) = 1 + 3 = 4 \text{ atoms}.$

In these questions (Q. No (i) to (iv)), a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

- (a) Assertion and reason both are correct statements and reason is correct explanation for assertion.
- (b) Assertion and reason both are correct statements but reason is not correct explanation for assertion.
- (c) Assertion is correct statement but reason is wrong statement.
- (d) Assertion is wrong statement but reason is correct statement.
- (i) Assertion (A): The total number of atoms present in a simple cubic unit cell is one.
 - Simple cubic unit cell has atoms at its corners, each of which is shared between eight adjacent unit cells.

- (ii) Assertion (A): In face-centred unit cell, atoms are at the centre of all faces.
 - Reason (R): Total number of atoms per unit cell is 4.
- (iii) Assertion (A): A crystal lattice is made up of unit cells.
 - (R): Each point in a crystal lattice represents an ion.

OR

- The total number of atoms in body-centred unit cell is 2.
- In body centred unit cell, an atom is at each of its corners and one atom at its body centre.
- (iv) Assertion (A): Bcc arrangement is less closely packed than ccp arrangement.
 - In ccp arrangement, the two atoms at the corners of the unit cell are Reason touching each other whereas in bcc arrangement, they are not touching each other

Answers

(iii) (c) **OR** (a)(ii)(d)(iv)(c)(a)(i)

ASSERTION-REASON QUESTIONS

In the following questions, a statement of assertion followed by a statement of reason is given. Choose the correct answer out of the following choices.

- Assertion and reason both are correct statements and reason is correct explanation for assertion.
- Assertion and reason both are correct statements but reason is not correct explanation for assertion.
- Assertion is correct statement but reason is wrong statement.
- Assertion is wrong statement but reason is correct statement.
- Assertion (A): Graphite is a good conductor of electricity however diamond belongs to the category of insulators.
 - **Reason** (R): Graphite is soft in nature on the other hand diamond is very hard and brittle.
- 2. Assertion (A): The total number of atoms present in a simple cubic unit cell is one.
 - (R): Simple cubic unit cell has atoms at its corners, each of which is shared between eight adjacent unit cells.
- 3. Assertion (A): The coordination number of square close packing in two dimensions is 6.
 - **Reason** (R): In square close packing, each sphere is in contact with four of its neighbours.
- 4. Assertion (A): Total number of octahedral voids present in unit cell of cubic close packing including the one that is present at the body centre, is four.
 - (R): Besides the body centre there is one octahedral void present at the centre of each of the six faces of the unit cell and each of which is shared between two adjacent unit cells.
- **5.** Assertion (A): The packing efficiency is maximum for the fcc structure.
 - **Reason** (R): The coordination number is 12 in fcc structures.
- Assertion (A): Semiconductors are solids with conductivities in the intermediate range from $10^{-6} - 10^4 \text{ ohm}^{-1} \text{ m}^{-1}$.
 - Reason (R): Intermediate conductivity in semiconductor is due to partially filled valence band.
- 7. Assertion (A): Schottky defect is generally shown by the compounds which have high coordination number.
 - (R): In Schottky defect, equal number of cations and anions are missing from the lattice sites

8. Assertion (A): Frenkel defect does not change the density of the solid.

Reason (R): In Frenkel defect smaller ion usually cation is dislocated from its normal site to an interstitial site

9. Assertion (A): In crystalline solids electrical resistance show different values when measured along different directions in the same crystal.

Reason (R): Crystalline solids are isotropic in nature

10. Assertion (A): Total number of octahedral voids present in unit cell of cubic close packing including the one that is present at the body centre, is four.

Reason (R): Besides the body centre there is one octahedral void present at the centre of each of the six faces of the unit cell and each of which is shared between two adjacent unit cells.

Answers

- **1.** (b) **2.** (a) **3.** (d) **4.** (c) **5.** (b) **6.** (c) **7.** (b) **8.** (a)
- **9.** (c) **10.** (c)

HINTS/SOLUTIONS OF SELECTED MCQS

- 1. (b) At low temperature, substance exists in solid state due to low thermal energy and hence decreased molecular motion, which in turn leads to strong intermolecular cohesive forces, i.e., which hold the constituent particles together.
- 7. (c) Iodine is a non-polar molecular solid iodine molecules are held together by London force or dispersion force, this is soft and non conductor for electricity.

Water is a hydrogen bonded molecular solid in which H and O are held together by hydrogen bonding to non-ionic nature, so, it is not an electrical conductor.

9. (c) In a face centered cubic lattice, atom (A) occupies the corner positions. There are 8 corner positions and each position contributes one eighth to the unit cell. Hence, total number of (A) atoms per unit cell = $8 \times 1/8 = 1$.

Atom (B) occupied the face centre positions. There are six face centre positions. One atom of (B) is missing from one of the face centered points. Thus, there are 5 face centre positions that are occupied with (B). Each such position contributes one half to the unit cell. Hence, total number of (B) atoms per unit cell= $5 \times 1/2 = 2.5$

The formula of the compound is $AB_{2.5}$ or A_2B_5

15. (c) For NaCl $\frac{r^+}{r^-} = 0.414$

$$r^{-} = \frac{100}{0.414} = 241.5 \text{pm}$$

16. (d) Distance between two oppositely charge ions for BCC

$$(r^{+} + r^{-}) = \frac{a\sqrt{3}}{2} = \frac{387 \times \sqrt{3}}{2}$$

= 335.15 pm

21. (d) Anisotropy is the property of substances to exhibit variations in electrical and optical properties along different planes of the same crystal. It is seen in crystals, liquid crystals and, less commonly, in liquids. Crystalline solids such as NaCl, BaCl₂, etc, will show anisotropy.

- The **hexagonal** closed packed (hcp) has a **coordination number** of 12 and contains 6 atoms per unit cell.
- (c) In ccp, number of atoms present per unit cell = 1
 - Since in cubic closed packing, the number of tetrahedral voids is equal to twice the number of atoms present per unit cell, hence
 - Number of tetrahedral voids = $2 \times 4 = 8$
- (d) Glass is an amorphous solid due to its short range order and irregular arrangement of constituent particles.
- (a) Crystalline solids are anisotropic in nature means some of their physical properties like electrical conductivity, refractive index etc. are different in different directions in the same crystal.
- (d) A ccp arrangement consists of three repeating layers (ABCABC...) of hexagonally arranged spheres. Spheres in a ccp structure have a coordination number of 12 because they contact six spheres in their layer, three atoms in the layer above and three spheres in the layer below.
- An atom that lies on the edge of a unit cell is shared by four adjacent unit cells, so the 28. contribution of each edge atom is $\frac{1}{4}$
- (b) The closest-packing sequence ABAB... represents hexagonal packing. In every second row, the particles occupy the depressions between the particles of first row. In the third row, the particles are vertically aligned with those in the first row giving ABABAB... arrangement.
- (c) The radius of the sphere that would fit well into an octahedral void in a close-packing is given by r = 0.414R; where r = radius of octahedral void and R = radius of sphere.
- (b) For tetrahedral arrangement, co-ordination number is 4 and radius ratio (r^+/r^-) is 0.225 0.414.
- (b) Covalent solids form crystals which can be viewed as a single giant molecule made up of an almost endless number of covalent bonds.
 - Out of the given solids, diamond is an example of covalent solid while Fe and Cu is a metallic solid, and NaCl is an ionic solid.
- (d) A molecular solid is composed of molecules held together by dispersion or london forces. Its properties are dictated by the weak nature of these intermolecular forces. Molecular solids are soft, have low melting temperatures, and are electrical insulators.
- (b) The existence of a substance in more than one solid modifications is known as polymorphism. It is the ability of a solid material to exist in more than one form or crystal structure.
- (d) Diamond being a crystalline solid is anisotropic in nature. In amorphous solids, there is not long-range order exist between them and therefore the given properties such as electrical conductivity, thermal conductivity, refractive index are identical in all directions and they are isotropic in nature.
- (a) Doping of impurity of higher valent to lower group elements results in formation of n-type of semiconductor. On mixing arsenic with silicon, n-type of semiconductor is obtained.
- (b) In graphite, each carbon is bonded with other 3 other carbons and one valence electron is free. Due to this valence electron, graphite conducts electricity in the solid state.
- (a) Iodine molecules are a class of non-polar molecular solid in which constituents molecules 38. are held together by London or dispersion forces. These solids are soft and are insulators.

- **39.** (d) Bravais Lattice refers to the 14 different ways of arrangement of points in a regular 3-dimensional space.
- **40.** (d) When NaCl is doped with MgCl₂, two Na⁺ are replaced by one Mg²⁺ ion to maintain electrical neutrality. Thus, the cationic vacancies thus produced are equal in number to that of Mg²⁺ ions. Thus, this type of defect is called impurity defect.
- **41.** (b) Glass is an amorphous solid. Like liquids, it has a tendency to flow, though very slowly and due to which, it is considered as pseudo-solid.
- 42. (b) There are a total of four formula units of NaCl per unit cell, i.e., 4Na^+ and 4Cl^- ions.

Na⁺ ions =
$$12 \times \frac{1}{4} + \frac{1}{\text{(body centre)}} = 4$$

(edges) $\frac{1}{\text{(centre)}} = 4$
Cl⁻ ions = $8 \times \frac{1}{8} + \frac{1}{2} \times 6 = 4$
(corners) (face

- 43. (c) Frenkel defect is produced because of missing ions from their normal crystal sites. It is produced when some ions are displaced from their normal sites and occupy interstitial sites. So, it does not affect the density of the crystal.
- 44. (a) The simple cubic has a coordination number of 6 and contains 1 atom per unit cell. This can be calculated as $8 \times \frac{1}{8} = 1$ (Corner atoms)
- **45.** (d) The face-centered cubic (fcc) has a coordination number of 12 and contains 4 atoms per unit cell. This can be calculated as $8 \times \frac{1}{8} + 6 \times \frac{1}{2} = 4$
- 46. (d) For bcc, Z = 2, $\rho = 530 \text{ kgm}^{-3}$ (Corner) (face atoms)

 Atomic mass of Li = 6.94 g mol⁻¹, $N_A = 6.022 \times 10^{23} \text{ mol}^{-1}$ $\rho = 530 \text{ kgm}^{-3} = \frac{530 \times 1000 \text{ g}}{1 \times (100)^3 \text{ cm}} = 0.53 \text{ g cm}^{-3}$

$$\rho = \frac{Z \times M}{N_A \times a^3}$$

$$\Rightarrow 0.53 = \frac{2 \times 6.94}{6.022 \times 10^{23} \times a^3}$$

$$a^3 = \frac{2 \times 6.94}{6.022 \times 10^{23} \times 0.53} = 43.5 \times 10^{-24} \text{ cm}^3$$

$$a = 352 \times 10^{-10} \text{ cm} = 352 \text{ pm}$$

47. (c) Radius ratio $\frac{r^+}{r^-} = \frac{0.98 \times 10^{-10}}{1.81 \times 10^{-10}} = 0.541$

It lies in the range of 0.414 to 0.732 hence, coordination number of each ion will be 6 as the compound will have NaCl type structure, *i.e.*, octahedral arrangement.

48. (b) Frenkel defect is a dislocation defect as smaller ions (usually cations) are dislocated from normal sites to interstitial sites. Frenkel defect is shown by compounds having large difference in the size of cations and anions. In case of alkali metal halides, the size alkali metal ions is large and therefore it cannot be fit into the interstitial site and hence do not show frenkel defect. Also, Schottky defect decreases the density of crystal while Frenkel defect has no effect on the density of crystal.

49. (d) Z = 4, i.e., structure is fcc Hence,

$$r = \frac{a}{2\sqrt{2}} = \frac{361}{2\sqrt{2}} = 127.65 \text{ pm} \approx 128 \text{ pm}$$

(a) Z = 4: for fee structure

$$d = \frac{Z \times M}{N_4 \times a^3}$$

$$M = \frac{d \times N_A \times a^3}{Z} = \frac{2.72 \times 6.02 \times 10^{23} \times (404 \times 10^{-10})^3}{4}$$

$$\Rightarrow$$

$$26.99 \approx 27 \,\mathrm{g \ mol^{-1}}$$
.

(d) Number of atoms in $ccp = 4 = O^{2-}$

Number of tetrahedral voids = $2 \times 4 = 8$

So, number of
$$A^{2+}$$
 ions = $8 \times \frac{1}{4} = 2$

Number of octahedral voids = Number of B^+ ion = 4

So, Ratio,
$$O^{2-}: A^{2+}: B^+$$

Hence, formula of oxide = AB_9O_9

(a) For a face-centered cubic (fcc) structure,

$$r = \frac{a}{2\sqrt{2}}$$
, $a = 408 \text{ pm}$, $r = \frac{408}{2\sqrt{2}}$
= 144 pm

Diameter = $2r = 2 \times 144 = 288 \text{ pm}$

- (a) Number of octahedral voids is same as number of atoms present in a unit cell. The number of atoms in each unit cell of ccp = 4. Therefore, the total number of octahedral voids per atom present in $ccp = \frac{4}{4} = 1$
- (a) Since Li crystallises in bcc crystal atomic radius, $r = \frac{\sqrt{3}a}{4}$ (a = edge length)

$$r = \frac{\sqrt{3}}{4} \times 351 = 151.8 \text{ pm (given a = 351 pm)}$$

Since copper crystallises in a face-centered cubic lattice, atomic radius, $r = \frac{a}{9\sqrt{9}}$ Given (edge length = a = 334 pm)

$$r = \frac{334}{2\sqrt{2}} = 118.10 = 118 \text{ pm}$$

- If silicon or germanium is doped with any of the element of group III (B, Al, Ga, In, Th) of the periodic table, p-type of semiconductor will be obtained.
- (b) As each Sr^{2+} ion introduces one cation vacancy, therefore, concentration of cation vacancies = mol% of $SrCl_2$ added.

Thus, Concentration of cation vacancies = 10^{-4} mol%

$$= \frac{10^{-4}}{100} \times 6.022 \times 10^{23}$$
$$= 6.022 \times 10^{17}$$

The maximum properties of the available volume which may be filled by hard sphere in **58.** simple cubic arrangement is given as $\frac{\pi}{6}$ or 0.52.

(b) F-centres are anionic sites occupied by unpaired electrons. They are responsible for colours.

60. (a)
$$\rho = \frac{Z \times M}{a^3 \times N_4} = \frac{2 \times 213}{(436.6 \times 10^{-10})^3 \times 6.022 \times 10^{23}} = 8.50 \text{ g/cm}^3$$

61. (a) In a unit cell, X atoms at the corners $=\frac{1}{8} \times 8 = 1$ Y atoms at the face centres = $\frac{1}{9} \times 6 = 3$

Ratio of X and Y = 1 : 3. Hence formula is XY_3 .

(b) In HCP unit cell

$$C = 4r\sqrt{\frac{2}{3}}, a = 2r$$

$$\frac{C}{a} = 2\sqrt{\frac{2}{3}} = \sqrt{\frac{8}{3}}$$

- (i) Diamond is the hardest known substance and graphite is soft.
 - (ii) In graphite, each carbon atom is joined to three other C atoms leaving one free valence electron which results in electrical conductance. In diamond, all four valencies of carbon are satisfied due to which diamond is an insulator. Thus, graphite has higher electrical conductivity than diamond.
 - (iii) Diamond has higher thermal conductivity than graphite due to the transfer of thermal vibrations from atom to atom. Diamond has a compact and precisely aligned crystal which helps in the fast movement of heat.
 - (iv) Graphite has a higher C-C bond order than diamond as graphite contains C=C double bond whereas diamond contains C-C single bond only.
- (a) The atom in the middle layers will have 12 nearest neighbours. But the atom in topmost layer has 9 nearest neighbours (6 atoms in the same layer + 3 atoms in the bottom layer).
- $\frac{r_{\text{Na}^+}}{r_{\text{Cl}^-}} = 0.55$ and $\frac{r_{\text{K}^+}}{r_{\text{Cl}^-}} = 0.74$ (a) Given; 65. $\frac{r_{\text{Na}^+}}{r_{\text{Cl}^-}} + 1 = 1.55$ and $\frac{r_{\text{K}^+}}{r_{\text{Cl}^-}} + 1 = 1.74$ $\frac{r_{\text{Na}^{+}} + r_{\text{Cl}^{-}}}{r_{\text{Cl}^{-}}} = 1.55$ $\frac{r_{\text{K}^{+}} + r_{\text{Cl}^{-}}}{r_{\text{Cl}^{-}}} = 1.74$ i.e., ...(i)

...(ii)

Dividing (ii)/(i), $\frac{r_{K^+} + r_{Cl^-}}{r_{K^+} + r_{Cl^-}} = \frac{1.74}{1.55} = 1.123$

(c) The formula $M_{0.98}$ O shows that if there were 100 O-atoms present as O^{2-} ions, then there 66. will be 98 metal atoms present as M^{2+} and M^{3+} .

Let
$$M^{3+} = x$$
, then $M^{2+} = 98 - x$.

As the compound as a whole is neutral, therefore

Total charge on M^{3+} and M^{2+} will be equal to total charge on 100 O^{2-} ions.

or
$$x (+3) + (98 - x) (+2) = 100 \times 2$$
or
$$3x + 196 - 2x = 200$$
or
$$x = 4$$

$$\therefore \qquad \% \text{ of } M^{3+} = \frac{4}{98} \times 100 = 4.08\%$$

and

- The defect that arises due to the missing of a cation from its lattice site and presence of the cation with higher charge in the adjacent site is called metal deficiency defect. Hence, the metals showing variable valency. For example, in FeO, as some Fe may be present as Fe³⁺ and for charge neutrality $3\text{Fe}^{2+} = 2\text{Fe}^{3+}$. Hence, FeO should have metal deficiency defect.
- $\rho = \frac{Z \times M}{a^3 \times N_4}$ 68. Density, (c) $2.72 = \frac{4 \times M}{(540 \times 10^{-10})^3 \times (6.02 \times 10^{23})}$ $M = \frac{2.72 \times (540)^3 \times 6.02 \times 10^{23} \times 10^{-30}}{4}$ or $= 64 \text{ g mol}^{-1}$
- In body-centered cubic (bcc), oppositely charged ions touch each other along the body diagonal.
 - Body diagonal = $2r_{Cs^+} + 2r_{Cl^-}$ But body diagonal = $\sqrt{3}a$ $2(r_{Cs^{+}} + r_{Cl^{-}}) = \sqrt{3}a$ $r_{\text{Cs}^+} + r_{\text{Cl}^-} = \frac{\sqrt{3}}{9}a$ or $r = \frac{\sqrt{3}}{4}a$ (c) For bcc, $=\frac{1.732}{4} \times 4.29 \text{ Å}$ $= 1.86 \,\text{Å}$
- 71. (b) For the fee structure, nearest neighbour distance $(d) \frac{a}{\sqrt{2}}$
- $d = \frac{Z \times M}{a^3 \times N_A}$ or $M = \frac{d \times a^3 \times N_A}{Z}$ **73.** (b) Z = 4for fcc, Now, we know that $n = \frac{m}{M} = \frac{N}{N_0}$ $\frac{100}{M} = \frac{N}{6.022 \times 10^{23}}$ So, $N = \frac{100 \times 6.022 \times 10^{23} \times Z}{d \times a^3 \times N}.$ $\Rightarrow \frac{100 \times 6.022 \times 10^{23} \times 4}{10 \times (200 \times 10^{-10})^3 \times 6.022 \times 10^{23}}$ $\Rightarrow \frac{40}{(200 \times 10^{-10})^3} \text{ or } \frac{40}{8 \times 10^{-24}}$ $\Rightarrow 5 \times 10^{24} \text{ atoms}$
- (a) Schottky defect arises when equal number of cations and anions are missing from the lattice so that electrical neutrality is maintained.
- (c) ABABA describes a hexagonal closed packed arrangement of spheres. In this type of arrangement, the coordination number is 6 and successive rows are arranged in such a way that the crests of the spheres of one row are placed into depression formed between adjacent spheres of the next row.

- 77. (b) Ionic solids have highest melting point due to strong electrostatic forces of attraction.
- **78.** (d) $A \Rightarrow 8 \times \frac{1}{8} = 1$

 $B \Rightarrow 5 \times \frac{1}{2} = \frac{5}{2}$ (Since one atom is missing from one of the face centred point.)

Therefore, formula = $A_1B_{5/2}$ or A_2B_5

79. (a) $n = \frac{m}{M} = \frac{1}{58.5}$ moles $n = \frac{N}{N_0} \text{ or } N = n \times N_0$ $= \frac{1}{58.5} \times 6.022 \times 10^{23} \text{ molecules}$

We know that, there are 4 molecules of NaCl present in a unit cell.

- $\therefore \text{ Number of the unit cell} = \frac{6.022 \times 10^{23}}{58.5 \times 4}$ $= 2.57 \times 10^{21} \text{ unit cells.}$
- 80. (d) As unit cell consists of A ions at the eight corners. Thus, number of A ions = $\frac{1}{8} \times 8 = 1$ Unit cell consists of B ions at the centre of the body. Thus, number of B ion = 1 So, the formula will be AB.
- 81. (d) For fcc, number of atoms = 4

 Volume of single atom = $\frac{4}{\pi}\pi r$

Volume of single atom = $\frac{4}{3}\pi r^3$

Total volume of atoms = $4 \times \frac{4}{3}\pi r^3$ = $\frac{16}{2}\pi r^3$

- **82.** (c) As the F.C.C unit cells consists of 6 faces. Therefore, unit cells is equally shared by 6 different unit cells.
- 83. (a) In crystalline solid there is a regular arrangement of the constituent particles. As the temperature increases the chance that a lattice site may be unoccupied by an ion increases. As the number of defects increases with temperature solid changes abruptly into liquid.
- **84.** (b) For simple cubic cell = $8 \times \frac{1}{8} = 1$

For
$$bcc = 8 \times \frac{1}{8} + 1 = 2$$

$$For fcc = 8 \times \frac{1}{8} + \frac{1}{2} \times 6$$
$$= 4$$

So, the ratio will be 1:2:4

85. (d) For $bcc = 8 \times \frac{1}{8} + 1 = 2$

$$For fcc = 8 \times \frac{1}{8} + \frac{1}{2} \times 6$$
$$= 4$$

Therefore, the number of Na and Mg atoms present in their respective crystals is 2 and 4.

86. (a) $a = 5 \text{ Å} \text{ or } 5 \times 10^{-8} \text{ cm}$

Volume of unit cell = a^3

$$= (5 \times 10^{-8})^{3}$$
$$= 1.25 \times 10^{-22} \text{ cm}^{3}$$

Given,

Density =
$$4 \text{ g/cm}^3$$

The mass of the unit cell =
$$4 \times 1.25 \times 10^{-22}$$

= 5×10^{-22} g

The mass of 1 mol of FeO is = 55.8 + 16 = 71.8 g

So the mass of 1 molecule of FeO will be

$$=\frac{71.8}{6.022\times10^{23}}=1.192\times10^{-22}$$
g

Number of FeO formula unit per unit cell

$$\frac{5 \times 10^{-22}}{1.192 \times 10^{-22}} \approx 4$$

So, the amount of Fe^{2+} and O^{2-} ions in each unit cell will be 4 Fe^{2+} and $4O^{2-}$.

- 87. (d) In KCl, co-ordination number of cation and anion is 6 and 6 respectively. The Frenkel defect is shown by the compounds having low coordination number while Schottky defect is shown by the ionic coordination number.
- 1 mole of compound = 6.022×10^{23} particles 88. (c)

0.5 mole of compound = $0.5 \times 6.022 \times 10^{23}$ particles

No. of tetrahedral voids = $2 \times \text{Number of particles}$

$$= 2 \times 3.011 \times 10^{23}$$
$$= 6.022 \times 10^{23}$$

No. of octahedral voids = Number of particles

$$= 3.011 \times 10^{23}$$

Hence,

Total no. of voids =
$$6.022 \times 10^{23} + 3.011 \times 10^{23}$$

= 9.03×10^{23}

Anions A form hcp So, number of anions (A) = 6 89.

Cations are 75% Octahedral voids

So, number of cations

$$(C) = 6 \times \frac{3}{4}$$
$$= \frac{18}{4} \Rightarrow \frac{9}{9}$$

So, the formula of compound is $C_{9/2} A_6$ or $C_9 A_{12}$ or $C_3 H_4$

90. (c)

$$\rho = \frac{Z \times M}{a^3 \times N_A}$$

for
$$fcc$$
, $Z = 4$, So,
$$\rho = \frac{4 \times N_A}{1 \times 10^{-2}}$$

for
$$bcc$$
, $Z = 2$, So,

$$\rho = \frac{4 \times M}{N_A \times (3.5 \times 10^{-8})^3} \dots (i)$$

$$\rho = \frac{2 \times M}{N_A \times (3 \times 10^{-8})^3} ...(ii)$$

By dividing equation (ii), we get

$$\rho_{fcc} : \rho_{bcc} = 1.259 : 1$$

(a) In bcc unit cell, lattice atoms occupies the corners and body center of a cube.

Each corner has a contribution of $\frac{1}{8}$ and there are 8 corners in a cube.

Effective number of atoms in a cube = $\frac{1}{8} \times 8 = 1$

Also, a body centre will have contribution of 1 and each cube will have 1 body centre.

So effective number of atoms in a cube= 1

Therefore, total number of atoms in bcc unit cell = 1 + 1 = 2

- **92.** (c) The number of octahedral voids in a cubic close packed structure is the same as effective number of atoms. In ccp, effective number of atoms are 4, so there are 4 octahedral voids.
- 93. (b) The radius ratio of the given voids is
 - Triangular 0.155 0.225
 - Cubic 0.732 1
 - \bullet Tetrahedral 0.225 0.414
 - \bullet Octahedral 0.414 0.732

Hence, the largest void is cubic void.

- **94.** (a) Close packing is maximum in the crystal lattice of face-centred cubic lattice with 74% packing efficiency.
- 95. (b) The formula for determination of density of cubic unit cell is

$$\rho = \frac{n \times M}{N_0 \times a^3} \text{ g cm}^{-3}$$

$$\rho = Density$$

n =Number of moles

$$M = Molar mass$$

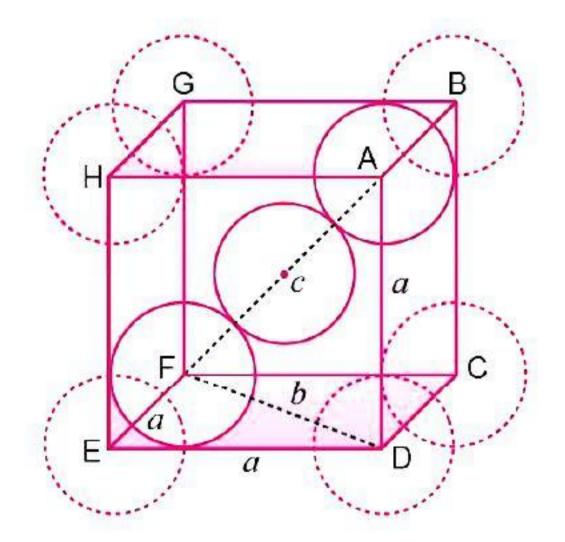
 $N_0 = \text{Avogadro's number}$

$$a = \text{Edge length}$$

- 96. (c) If the number of close-packed spheres is n, then the number of tetrahedral voids is 2n.
- 97. (b) Molecular solids refer to those solids which are composed of molecules held together by the london or dispersion forces, dipole-dipole interactions and hydrogen bonding.

The dry ice (CO₂) is an example of a molecular solid. Graphite is a covalent solid, sodium chloride is a ionic solid and Fe is a metallic solid.

98. (c) **Efficiency of packing in bcc structures:** In this case the atom at the centre is in touch with other two atoms which are diagonally arranged (see below figure). The spheres along the body diagonal are shown with solid boundaries.



In
$$\Delta EFD$$
,

$$b^2 = a^2 + a^2 = 2a^2$$

$$b = \sqrt{2} a$$

In $\triangle AFD$,

$$c^2 = a^2 + b^2 = a^2 + 2a^2 = 3a^2$$

$$\therefore c = \sqrt{3} a$$

The length of the body diagonal c is equal to 4r, r being the radius of the sphere (atom). As all the three spheres along the diagonal touch each other,

$$c = 4r$$

Therefore, $c = 4r = \sqrt{3} a$

$$a = \frac{4r}{\sqrt{3}}$$
 or $r = \frac{\sqrt{3}}{4}a$

As already calculated, the total number of atoms associated with a bcc unit cell is 2, the volume(v) is, therefore,

$$2 \times \frac{4}{3}\pi r^3 = \frac{8}{3}\pi r^3$$

Volume of the unit cell $(V) = a^3 = \left(\frac{4r}{\sqrt{3}}\right)^3 = \frac{64r^3}{3\sqrt{3}}$

Packing efficiency =
$$\frac{v}{V} \times 100 = \frac{(8/3)\pi r^3}{(64/3\sqrt{3}) \times r^3} \times 100 = \frac{\sqrt{3}}{8}\pi \times 100 = 68\%$$

Therefore, 68% of unit cell is occupied by atoms and the rest 32% is empty space.

The distance between the body centered atom and one corner atom is $\frac{\sqrt{3a}}{9}$, *i.e.*, half of the body diagonal 99. body diagonal.