- 1. The number of p-orbital not involved in the hybridisation of middle carbon atom in  $CH_2 = C = CH_2$  is (a) 3 (b) 1 (c) 2 (d) Cannot be predicted
- 2. In the following structure

$$H_{3}C$$
  
 $H_{3}C$   
 $H$ 

how many carbon atom is  $sp^2hybridised$ (a) 1 (b) 2 (c) 3 (d) 4

- 3. The hybridisation of the central atom will change when (a) NH<sub>3</sub> combines with H<sup>+</sup>
  (b) H<sub>3</sub>BO<sub>3</sub> combines with OH<sup>-</sup>
  (c) NH<sub>3</sub> form NH<sub>2</sub><sup>-</sup>
  (d) H<sub>2</sub>O combines with H<sup>+</sup>
- 4. Among the interhalide species

 $IF_2^{\Theta}$ ,  $IF_3$ ,  $IF_4^{\Theta}$  and  $IF_7$ 

(a) All iodine centres are either sp<sup>3</sup>d or sp<sup>3</sup>d<sup>2</sup>hybridised

(b) The minimum angular separation between fluorine atoms is  $60^{\circ}$ 

(c)The anionic species are both isoelectronic and isostructural to XeF2 and XeF4

(d)There is no species having a single lone pair of electrons

- 5. LiF is least soluble among the fluorides of alkali metals, because
  - (a) Smaller size  $Li^+$  impart significant covalent character in LiF
  - (b) The hydration energies of  $Li^{\scriptscriptstyle +}$  and  $F^{\scriptscriptstyle -}$  are quite higher
  - (c) Lattice energy of LiF is quite higher due to the smaller size of Li<sup>+</sup> and F<sup>-</sup>
  - (d) LiF have strong polymeric network in solid
- 6. Which of the following compounds has sp<sup>2</sup>hybridisation ? (a)  $CO_2$  (b)  $SO_2$  (c)  $N_2O$  (d) CO
- 7.  $N(SiH_3)_3$  has -
  - (a) Sp<sup>3</sup>hybridisation, pyramidal shape
  - (b) Sp<sup>2</sup>hybridisation, planar shape
  - (c) Sp<sup>3</sup>hybridisation, tetrahedral shape
  - (d) d sp<sup>2</sup>hybridisation, square planar shape
- **8.** Which of the following angle corresponds to  $sp^2$  hybridisation?

(a)  $90^{\circ}$  (b)  $120^{\circ}$  (c)  $180^{\circ}$  (d)  $109^{\circ}$ 

- 9. On hybridisation of one s and three p-orbitals, we get(a) Four orbitals with tetrahedral orientation(b) Three orbitals with trigonal orientation
  - (c) Two orbitals with linear orientation
  - (d) Two orbitals with perpendicular orientation.
- **10.** Which of the following shown  $dsp^2$  hybridisation and a square planar geometry?

(a)  $SF_6$  (b)  $BrF_5$  (c)  $PCl_5$  (d)  $[Ni(CN)_4]^{2-1}$ 

**11.** Which of the following statements is true about hybridisation?

(a) The hybridized orbitals have different energies for each orbital.

- (b) The number of hybrid orbitals is equal to the number of atomic orbitals that are hybridized.
- (c) Hybrid orbitals form multiple bonds.
- (d) The orbitals with different energies undergo hybridisation.

12. Hybridisation state of Xe in  $XeF_2$ ,  $XeF_4$  and  $XeF_6$  respectively are

(a) $sp^2$ , $sp^3$ , $d$ , $sp^3d^2$	(b) $sp^3d, sp^3d^2, sp^3d^3$
(c) $sp^3d^2$ , $sp^3d$ , $sp^3d^3$	(d) $sp^2$ , $sp^3$ , $sp^3d$

13. Match the column I with column II and mark the appropriate choice.

	Column I	Column II			
	(1) $C_2 H_2$	(i) $Sp^{3}d^{2}$ hybridisation			
	(2) SF <sub>6</sub>	(ii) Sp <sup>3</sup> d <sup>3</sup> hydridisation			
	(3) SO <sub>2</sub>	(iii) sp hybridisation			
	(4) IF <sub>7</sub>	(iv) Sp <sup>2</sup> hybridisation			
(8	$(a) (1) \rightarrow (i), (2) \rightarrow (iii), (3) \rightarrow (ii), (4) \rightarrow (iv)$				
(b) (1) $\rightarrow$ (iii), (2) $\rightarrow$ (i), (3) $\rightarrow$ (iv), (4) $\rightarrow$ (ii)					
(c) (1) $\rightarrow$ (ii), (2) $\rightarrow$ (iii), (3) $\rightarrow$ (i), (4) $\rightarrow$ (iv)					
(0	(d) (1) $\rightarrow$ (iv), (2) $\rightarrow$ (i), (3) $\rightarrow$ (iii), (4) $\rightarrow$ (ii)				

14. The hybridisation of the underline atom changes in :

(a) AlH <sub>3</sub> changes to AlH <sub>4</sub> <sup><math>-</math></sup>	(b) $H_2O$ changes to $H_3O^+$
(c) $NH_3$ changes to $NH_4^+$	(d) In all cases

- 15. In which of the following compounds B F bond length is shortest ? (a)  $BF_4^-$ (b)  $BF_3 \rightarrow NH_3$ (c) BF3 (d)  $BF_3 \leftarrow N(CH_3)_3$
- **16.** Number of bonds in  $SO_2$  are :

(a) two $\sigma$ and two $\pi$	(b) two $\sigma$ and one $\pi$
(c) two $\sigma$ , two $\pi$ and one lone pair	(d) none of these

17. Choose the molecules in which hybridisation occurs in the ground state ?

(a) $\mathbf{BCl}_{3}$		(b) $NH_3$	(c) PCl <sub>3</sub>	(d) $BeF_2$
The co	rrect answer is -	-		
(a) 1, 2	2, 4	(b) 1, 2, 3		
(c) 2, 3	;	(d) 3, 4		
<b>18.</b> sp <sup>2</sup> – hybrid	lisation is shown	by :		
(a) BeCl <sub>2</sub>	(b)) BF <sub>3</sub>			
(c) NH <sub>3</sub>	(d) XeF <sub>2</sub>			
<b>19.</b> The hybridis	sation of carbon i	n diamond, graphit	e and acetylene is (respective	ely) –
(a) sp <sup>3</sup> , sp <sup>2</sup> , sp	(b) sp <sup>3</sup> , sp, sp <sup>2</sup>			
(c) $sp^2$ , $sp^3$ , $sp$	(d) sp, sp <sup>3</sup> , sp <sup>2</sup>			
<b>20.</b> Each carbon	in carbon subox	ide $(C_3O_2)$ is :		
(a) sp <sup>2</sup>	- hybridized	(b) sp <sup>3</sup>	-hybridized	
(-) 1		(1) 2	1 1 1 1 1 1 1 1 1 1	1 1 /

(d) sp<sup>2</sup>-hybridized but linked with one co-ordinate bond (c) sp-hybridized

<sup>21.</sup> Among the following pairs in which the two species are not isostructural is :

www.neetjeenotes.com		NEET/JE	E MAIN PRACTICE P	APER 2024-2025
(a) $SiF_4$ and $SF_4$	(b) IO <sub>3</sub> - a	nd XeO <sub>3</sub>		
(c) $BH_4^-$ and $NH_4^+$	(d) $PF_6^-$ and $SF_6^-$	-		
<b>22.</b> Consider the following iodides :	PI <sub>3</sub>	AsI <sub>3</sub>	SbI <sub>3</sub>	
	102°	100.2°	99°	
The bond angle is maximum i	in Pl <sub>3</sub> , which is :			
(a) due to small size of phosp	horus (I	b) due to more b	p–bp repulsion in PI <sub>3</sub>	
(c) due to less electronegativit	ty of P (o	d) none of these		
<b>23.</b> $OF_2$ is gS %				
<ul><li>(a) Linear molecule and sp hy</li><li>(c) Bent molecule and sp<sup>3</sup> hyb</li></ul>	vbridised (b) Tetral pridised (d) None	nedral molecule a of these	and sp³ hybridised	
<b>24.</b> Select the correct statement for the su	alphuric acid.			
(I) It has high boiling point ar	nd viscosity.			
(II) There are two types of bo	nd lengths in its biva	lent anion.		
(III) $p\pi$ - $d\pi$ bonding between s	sulphur and oxygen i	s observed.		
(IV) Sulphur has the same hyl	bridisation that is of	boron in diboran	e.	
(a) II and III only (b) II	, III and IV only			
(c) I, III and IV only (d) II	II and IV only			
<b>25.</b> Identify the correct match.				
(i) XeF <sub>2</sub>	(a) Centra	al atom has sp <sup>3</sup> h	ybridisation and bent ge	cometry.
(ii) N <sub>3</sub> -	(b) Central atom l	nas sp <sup>3</sup> d <sup>2</sup> hybridi	sation and octahedral.	
(iii) $PCl_{s}(s)$ anion	(c) Central atom h	nas sp hybridisat	ion and linear geometry	
(iv) $I_2 Cl_6(\ell)$ cation	(d) Central atom l	nas sp <sup>3</sup> d hybridis	ation and linear geometr	ry.
(a) $(i - a)$ , $(ii - b)$ , $(iii - c)$ , $(iv - d)$	(1	b) $(i - d)$ , $(ii - b)$	, (iii – d), (iv – c)	
(c) $(i - b)$ , $(ii - c)$ , $(iii - a)$ , $(iv - d)$	((	d) $(i - d)$ , $(ii - c)$	, (iii − b), (iv − a)	
<b>26.</b> Correct statement regarding this react	tion :			
$BF_3 + NH_3 \longrightarrow [F_3B \leftarrow N]$	H.]			
(a) hybridisation of only N ch	anges (1	o) hybridisation	of only B changes	
(c) hybridisation of N and B b	both change	(d) none of	of these.	
(a) dsoy N dk ladj.k cnyrk g	S (1	o) dsoy B dk lad	lj.k cnyrk gSA	
(c) N rFkk B nksuksa ds ladj	j.k cnyrs gaSA	(d) buesa	a ls dksbZ ugha	
<b>27.</b> The type of hybrid orbitals used by cl	hlorine atom in CIO-, C	$CIO_2^-$ , $CIO_3^-$ and $C$	$2IO_4^-$ are :	
(a) sp. sp <sup>2</sup> . sp <sup>3</sup> and sp <sup>3</sup> d (b) sr	$o$ and $sp^3$			
(c) only $sp^3$ (d) or	nly sp			
<b>28.</b> The hybridisation of orbitals of N ato	om in $NO_3^-$ , $NO_2^+$ and N	${\rm IH}_{4}^{+}$ are respective	ely :	
(a) sp, sp <sup>2</sup> , sp <sup>3</sup> (b) sp <sup>2</sup> , sp, sp	3			
(c) sp, sp <sup>3</sup> , sp <sup>2</sup> (d) sp <sup>2</sup> , sp <sup>3</sup> , sp	p			

**29.** The species in which the N atom is in a state of sp hybridization is :

(a)  $NO_2^-$  (b)  $NO_3^-$ (c)  $NO_2$  (d)  $NO_2^+$ 

**30.** The type of hybridisation and number of lone pair(s) of electrons of Xe in XeOF<sub>4</sub>, respectively, are :

(a)  $sp^3d^2$  and 1(b)  $sp^3d^2$  and 2(c)  $sp^3d$  and 1(d)  $sp^3d$  and 2

#### NEET/JEE MAIN PRACTICE PAPER 2024-2025

1. (c)

The middle carbon atom is sp hybridised therefore the number of p-orbitals not involved in hybridisation is 2

# 2. (d)

Here  $\overset{1}{C}$ ,  $\overset{2}{C}$ ,  $\overset{5}{C}$ , &  $\overset{6}{C}$  are sp<sup>2</sup> hybridised So total no. of carbon atoms having sp<sup>2</sup> configuration is 4

3. (b)  

$$B(OH)_3 + OH^- \rightarrow B(OH)_4^-$$
  
 $sp^2$   $sp^3$ 

# 4. (c)

Hybridisation and structure of  $IF_2^- \rightarrow sp^3d$  & linear Hybridisation and structure of  $XeF_2 \rightarrow sp^3d$  & linear

So Both  $XeF_2$  and  $IF_2^-$  are isostructural and isoeletronic

Hybridisation and structure of  $IF_4^- \rightarrow sp^3d^2$  & square

Planar

Hybridisation and structure of  $XeF_4 \rightarrow sp^3d^2$  & square

Planar

 $\Rightarrow$  Both XeF<sub>4</sub> and IF<sub>4</sub><sup>-</sup> are isoelectronic and isostructural

# 5. (c)

The small size of both  $Li^+$  and  $F^-$  ion leads to a very high value of lattice energy and thus crystal of LiF is very difficult to break

#### 6. (b)

Rest of the molecules are sp hybridized

## 7. (b)

lp of e<sup>-</sup>s of nitrogen is donated to the vacant d-orbital of Si.

- **8.** (b) : The angle corresponds to  $sp^2$  hybridisation triangular planar is  $120^0$
- **9.** (a) :



Four  $sp^3$  hybrid orbitals are formed when one s and three p – orbitals hybridise.

**10.** (d) :



11. (b) : The number of orbitals which hybridise remains same after the hybridisation also. E.e.,



#### NEET/JEE MAIN PRACTICE PAPER 2024-2025



**12.** (b) : 
$$XeF_2 - sp^3d$$



Total no. of valence electrons = 22  $\frac{22}{8} = 2(Q) + 6(R), \frac{6}{2} = 3(Q)$  X = 2 + 3 = 5Hybridisation is sp<sup>3</sup>d. EeF<sub>4</sub> -



Hybridisation is  $sp^{3}d^{2}$ Total no. of electrons in outermost shells = 8 +28 =36  $\frac{36}{8} = 4(Q) + 4(R), \frac{4}{2} = 2(Q) + 0(R)$ X = 4 + 2 + 0 = 6Hybridisation is  $Sp^{3}d^{2}$ XeF<sub>6</sub>



Total no. of valence electrons = 8+42 = 50  $\frac{50}{8} = 6(Q) + 2(R), \frac{2}{2} = 1(Q)$ X= 6+1=7Hybridisation is Sp<sup>3</sup>d<sup>3</sup>.

**13.** (b) : In  $C_2H_2$ , C undergoes, sp hybridisation. Ground state

Excited state

$$\underbrace{\uparrow} \underbrace{\uparrow} \uparrow \uparrow \uparrow$$
 H - C = C - H

In  $SF_6$ , S undergoes  $sp^3$ ,  $d^2$  hybridisation. Ground state)

Excited state

Excited state

#### NEET/JEE MAIN PRACTICE PAPER 2024-2025



In  $So_2$ , S undergoes  $sp^2$  hybridization.

In  $\mathbf{IF}_7 \mathbf{I}$  undergoes  $\mathbf{sp}^3 \mathbf{d}^3 \mathbf{hybridization}$ . Ground state



14. (a)



## 15. (c)

Partial double bond character is developed in B-F bond of BF3 (due to formation of dative p bond)

In other case B is sp<sup>3</sup> hybridised (tetrahedral molecule or ion) and the possibility for  $\pi$  bonding no longer exists.

16. (c)

(c)  $\bigvee_{\sigma \sigma}^{\pi}$ 





(b) Electronic configuration of nitrogen in ground state is 1s<sup>2</sup>2s<sup>2</sup>2p<sup>3</sup>.



(c) Electronic configuration of phosphorus in ground state is 1s<sup>2</sup>2s<sup>2</sup>2p<sup>6</sup>3s<sup>2</sup>3p<sup>3</sup>.



(d) Electronic configuration of boron in ground state is  $1s^22s^2$ .



**18.** (b) Steric number = 0 + 3 = 3; so sp<sup>2</sup> hybridisation

**19.** (a) Diamond  $sp^3$ ; graphite =  $sp^2$ ; Acetylene = sp

$$O = C = C = C = O$$
$$| \qquad | \qquad |$$
$$sp \qquad sp \qquad sp$$

20. (c)

**21.** (a) (a) Tetrahedral and see-saw shaped.



(b) Both are sp<sup>3</sup> hybridised and trigonal pyramid.

- (c) Both are sp<sup>3</sup> hybridised and tetrahedral.
- (d) Both are sp<sup>3</sup>d<sup>2</sup> hybridised and octahedral.



Phosphorus is the most electronegative of the central atoms. Consequently, it exerts the strongest pull on shared electrons, concentrating these electrons near P and increasing bonding pair-bonding pair repulsions–hence, the largest angle in  $PI_3$ . Sb, the least electronegative central atoms, has the opposite effect : Shared electrons are attracted away from Sb, reducing repulsions between the Sb–I bonds. The consequence is that the effect of the lone pair is greatest in  $SbI_3$ , which has the smallest angle.

Atomic size arguments can also be used for these species. Larger outer atoms result in larger angles ; larger central atoms result in smallest angles.

# NEET/JEE MAIN PRACTICE PAPER 2024-2025



www.neetjeenotes.com

28.	(b)	$NO_2^+$	Number of electron pairs $= 2$
			Number of bond pairs = 2 Number of lone pair = 0 So, the species is linear with sp hybridisation.
			O = N = O
		NO3-	Number of electron pairs = 3 Number of bond pairs = 3 Number of lone pair = 0 So, the species is trigonal planar with sp <sup>2</sup> hybridisation. $\overline{O} - \overset{+}{N} \overbrace{O}^{\overline{O}} \rightarrow sp^{2}$
		$\mathrm{NH}_4^+$	Number of electron pairs = 4 Number of bond pairs = 4 Number of lone pair = 0 So, the species is tetrahedral with sp <sup>3</sup> hybridisation. $\begin{bmatrix} H \\ H \\ H \end{bmatrix}^{+} \rightarrow sp^{3}$
29.	(d)	$NO_2^- = sp^2$	
		$NO_3^- = sp^2$ $NO_2^- = sp^2$ $NO_2^+ = sp$	
30.	(a)	F F F	