		INDIAN SCHOOL AL WADI AL KABIR		
Class: XI	[Department: SCIENCE 2021 - 22 SUBJECT :CHEMISTRY	Date of submis 27.02.2022	sion:
Worksheet No: 13 WS WITH ANS.		Chapter: 9 – Coordination Compounds	Note: A4 FILE FORMAT	
NAME OF THE STUDENT		CLASS & SEC:	ROLL NO.	
Q. No.		Questions and Answers		Marks
1.				1 each

Q. NO.		Questions and A	Inswers	iviarks		
1.	Name the following complex compounds or ions.					
	1. [Al (H ₂ O) ₆] Br ₃					
	2. [Cr (NH ₃) ₆] Cl ₃	Hexaamminechromium (III) chlo	oride			
	3. K ₃ [FeF ₆]	Potassium hexafluoroferrate (III)			
	4. [Zn (OH) ₄] ⁻²	Tetrahydroxozincate (II) ion				
	5. [Co (H ₂ O) ₄ Cl ₂] Cl	Tetraaquadichlorocobalt (III) ch				
	6. [Cu (NH ₃) ₄] ⁺²	NH ₃) ₄] ⁺² Tetraamminecopper (II) ion				
	7. K ₂ [SnCl ₆] Potassium hexachlorostannate (IV)					
	8. [Pt (NH ₃) ₄ Cl ₂] [PtCl ₆] Tetraamminedichloroplatinum (IV) hexachloroplatinate (IV)					
2.	Write the formu	la for each of the following com	nlex compounds or ions	1 each		
	white the forma					
	1. Hexaamminecob	alt (III) chloride	[Co (NH ₃) ₆] Cl ₃			
	2. Diamminetetrabi	romoplatinum (VI) bromide	[Pt (NH ₃) ₂ Br ₄] Br ₂			
	3. Tetraaquacadmiu	um (II) nitrate	[Cd (H ₂ O) ₄] (NO ₃) ₂			
	4. Diamminesilver	(I) ion	[Ag (NH ₃) ₂] ⁺			
	5. Sodium tetracyan	nocuprate (I)	Na ₃ [Cu (CN) ₄]			
	6. Silver hexacyand	oferrate (II)	Ag ₄ [Fe (CN) ₆]			
	7. Tetraammineoxa	latonickel (II)	[Ni (NH ₃) ₄ C ₂ O ₄]			

3.	Identify the denticity of the ligands given below:					1 each	
	aqua ammin benzen	e e	$\begin{array}{c} H_2O\\ NH_3\\ C_6H_6 \end{array}$	mor mor mor	odentate odentate odentate		
	carbon nitrosy	yl I	CO NO	mor mor	odentate odentate		
	methyl dimeth trimeth	amine ylamine ylamine	CH ₃ NH ₂ (CH ₃) ₂ NH ₂ N(CH ₃) ₃	mor mor mor	odentate odentate odentate		
	ethylen diethyl triethyl	ediamine enediami enetetraa	or en ne or dien mine or trien	H2NCH NH(CH N(CH2	I2CH2NH2 I2CH2NH2)2 CH2NH2)3	bidentate tridentate tetradentate	
	Pyridin trimeth	e or py ylphosph	ine (PMe3)	C5H5N P(CH3)	3	monodentate monodentate	
4.	Identify the geometry of the given complexes:						1 each
	CN	Geometry	,		Hybridization	Example]
	2	Linear			sp	[Ag(NH ₃) ₂]*	
	4	Tetrahedra	1		sp^3	[Cd(NH ₃) ₄] ²⁺	
	4	square pla	nar		sp²d	[Cu(OH ₂) ₄] ²⁺	
	5	trigonal bi	pyramid		$sp^{3}d$	Fe(CO) ₅	
		Szmara nya	amidal		sp^2d^2	[MnCl 5]3-	
	<u> </u>	oquare py					
	6	Octahedra	1		sp ³ d ²	[Fe(CN) ₆] ⁴ ·	

5.	Show the: Formation of [CoCl ₄] ²⁻	2 each
	Co atom in the ground state $3d$ $4s$ $4p$	
	Co^{2+} ion	
	Formation of $[CoCl_4]^{2-}$ $\downarrow \downarrow \downarrow \downarrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \uparrow \downarrow \downarrow \downarrow \downarrow \downarrow \downarrow$	
6.	Show the : Formation of [Ni(CO) ₄]:	2 each
	Atomic orbitals of Ni in (Z=28) ground state3d4s4p	
	Hybridized sp ³ orbitals of Ni	
	Formation of [Ni(CO)4] (1)	
	FOUR ELECTRON PAIRS DONATED BY FOUR CO MOLECULES	
	sp ³ HYBRIDIZATION	



12.	What are the factors affecting the stability of the complexes?	1 each
	Factors affecting the stability of a complex ion(i) Nature of metal ion : Greater the charge and smaller the size of the ion, more is its charge density and greater will be stability of the complex.(ii) Nature of ligand : More the basicity of ligand, more is its tendency to donate electron pair and therefore, more is the stability of the complex.	
13.	 When a coordination compound CoCl₃.6NH₃ is mixed with AgNO₃, 3 moles of AgCl are precipitated per mole of the compound. Write (i) Structural formula of the complex (ii) IUPAC name of the complex . 	1 each
	(i) Complex so formed is: $CoCl_3.6NH_3 + AgNO_3 \rightarrow [Co(NH_3)_6]Cl_3$	
	(ii) IUPAC name of complex is: Hexaamminecobalt (III) chloride	
14.	Using IUPAC norms write the formulae for the following: (a) Potassium trioxalatoaluminate (III) (b) Dichloridobis(ethane-l, 2-diamine) cobalt (III) ion	1 each
	(a) $K_3[Al(C_2O_4)_3]$ (b) $[Co(Cl)_2(en)_2]^+$	
15.	For the complex [Fe(en) ₂ Cl ₂], Cl, (en = ethylene diamine), identify (i) the oxidation number of iron, (ii) the hybrid orbitals and the shape of the complex, (i) [Fe(en) ₂ Cl ₂] Cl or $x + 0 + 2$ (-1) + (-1) = 0 x + (-3) = 0 or $x = + 3\therefore Oxidation number of iron, x = + 3(ii) The complex has two bidentate ligands and two monodentate ligands. Therefore,the coordination number is 6 and hybridization will be d2sp3 and shape will beoctahedral.$	1 each
16	 Explain the following terms giving a suitable example in each case : (i) Ambidentate ligand (ii) Denticity of a ligand (iii) Crystal field splitting in an octahedral field (All India 2011) (i) Ambidentate ligand: The monodentate ligands with more than one coordinating 	
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	atoms is known as ambidentate ligand. Monodentate ligands have only one atom capable of binding to a central metal atom or ion. For example, the nitrate ion NO_2^- can bind to the central metal atom/ion at either the nitrogen atom or one of the oxygen atoms.					
	Example : — SCN thiocyanato - S, — NCS thiocyanato-N					
	(ii) Denticity of a ligand: The number of donor atoms in a ligand which forms coordinate bond with the central metal atom are called denticity of a ligand. Example : If donor atom is one then it is called Monodentate ligand, if it is two, then it is called Bidentate and so on.					
	(iii) Crystal field splitting: It is the splitting of the degenerate energy levels due to the presence of ligands. When ligand approaches a transition metal ion, the degenerate d-orbitals split into two sets, one with lower energy and the other with higher energy. This is known as crystal field splitting and the difference between the lower energy set and higher energy set is known as crystal field splitting energy (CFSE)					
17	 State a reason for each of the following situations : (i) Co²⁺ is easily oxidized to Co³⁺ in presence of a strong ligand. (ii) CO is a stronger complexing reagent than NH₃. (iii) The molecular shape of [Ni(CO)₄] is not the same as that of [Ni(CN)₄]²⁻ (Delhi 2011) 	1 each				
	(i) Because in the presence of strong ligands, the crystal field splitting energy is more than the energy required to oxidise Co^{2+} .					
	(ii) This is due to the formation of π – bond by back donation of electrons from metal to carbon of CO or due to synergic bonding.					
	(iii) CO is a stronger field ligand than CN. Ni is in zero oxidation state in Ni(CO) ₄ and has tetrahedral geometry. But, Ni is in +2 oxidation state in $[Ni(CN)_4]^{2-}$ and has dsp ² hybridization (different geometry than tetrahedral sp ³).					
18.	Draw the geometry of the various complexes.	1 each				
	$[Fe(CN)_6]^4$ [Ni(CN)_4] ²⁻ [Cr(C ₂ O ₄) ₃] ³⁻					
	$\begin{bmatrix} CN & CN \\ CN & Fe & CN \\ CN & CN \\ CN & CN \end{bmatrix}^{2} $					
PREPAR	RED BY : Ms. Jenifer Robinson					