# **JEE Main/Boards**

# **Exercise 1**

**Q.1** Copper can be extracted by hydrometallurgy but not zinc. Explain.

**Q.2** Out of C and CO, which is a better reducing agent at 673 K?

**Q.3** How is leaching carried out in case of low grade copper ores?

**Q.4** Why is zinc not extracted from zinc oxide, through reduction using CO?

**Q.5** Name the common elements present in the anode mud in electrolytic refining of copper. Why are they so present?

**Q.6** State the role of silica in the metallurgy of copper.

Q.7 What is meant by the term "chromatography"?

Q.8 How is 'cast iron' different from 'pig iron"?

Q.9 Differentiate between "minerals" and "ores".

**Q.10** Out of C and CO. which is a better reducing agent for ZnO?

**Q.11** What is the role of graphite rod in the electrometallurgy of aluminium?

Q.12 Describe a method for refining nickel.

**Q.13** Giving examples, differentiate between 'roasting' and 'calcination".

Q.14 Why copper matte is put in silica lined converter?

**Q.15** What is the role of cryolite in the metallurgy of aluminium?

Q.16 What types of ores are roasted?

Q.17 Name one each of (a) acidic flux (b) basic flux.

Q.18 What is gangue?

**Q.19** What name is given to carbon reduction process for extracting the metal?

**Q.20** Outline the principles of refining of metals by the following methods:

(i) Zone refining (ii) Electrolytic refining

(iii) Vapour phase refining

**Q.21** What is the role of depressant in froth floatation process?

**Q.22** Why is the extraction of copper from pyrites more difficult than that from its oxide ore through reduction?

Q.23 Explain:

(i) Zone refining (ii) Column chromatography

**Q.24** Write down the reactions taking place in different zones in the blast furnace during the extraction of iron.

**Q.25** Write chemical reactions taking place in the extraction of zinc from zinc blende.

**Q.26** What criterion is followed for the selection of the stationary phase in chromatography'?

**Q.27** How can you separate alumina from silica in a bauxite ore associated with silica? Give equations, if any?

**Q.28** The value of  $\Delta_{12}G^{0}$  for formation of Cr<sub>2</sub> O<sub>3</sub> is -540 kJ mol<sup>-1</sup> and that of Al<sub>2</sub>O<sub>3</sub> is - 827 kJ mol<sup>-1</sup>. Is the reduction of Cr<sub>2</sub>O<sub>3</sub> possible with Al?

**Q.29** The choice of a reducing agent in a particular case depends on thermodynamic factor. How far do you agree with this statement? Support your opinion with two examples.

**Q.30** Name the processes from which chlorine is obtained as a by-product. What will happen if an aqueous solution of NaCl is subjected to electrolysis?

**Q.31** Predict conditions under which Al might be expected to reduce MgO.

Exercise 2			<b>Q.11</b> Purification of silicon element used in semi- conductors is done by							
Single Correct Choice Type			(A) Zone ref	ining	(B) Hea	ting				
<b>Q.1</b> In presence of oxygen, removal of sulphur from an ore is included in		(C) Froth flo	atation	(D) Hea	iting in v	vacuum				
(A) Calcina	ation	(B) Roasting		Q.12 Alumir	nothermic p	orocess i	is used f	for meta	llurg	jy of
(C) Smelti	ng	(D) Fluxing		(A) Pb		(B) Ag				
	5			(C) Al		(D) Nor	ne of the	ese		
<b>Q.2</b> In the metal is fo	extraction o ormed by re	of copper from induction of Cu <sub>2</sub> C	ts sulphide ore, the ) with	Q.13 In elect	trorefining	of coppe	er some	gold is d	lepo	sited
(A) FeS	(B) CO	(C) Cu <sub>2</sub> S	(D) SO <sub>2</sub>	as						
				(A) Anode m	nud	(B) Cath	node mu	bu		
Q.3 Pyrolu	isite is a/an			(C) Cathode		(D) Elec	trolyte			
(A) Oxide	ore	(B) Sulphide d	bre	<b>011</b> Poling	process is	ucod				
(C) Carbid	e ore	(D) Not an or	e	(A) For the r		useu	m Cu			
<b>0.4</b> The m	netal always	found in the fre	e state is	(A) For the r	emoval of					
(A) Au	(B) 7n	(C) Cu	(D) Na	(B) For the removal of $Al_2O_3$ from Al						
(/)//0		(0) 60		(C) For the removal $Fe_2O_3$ from Fe						
<b>Q.5</b> In the	extraction	of iron, the slag	produced is?	(D) In all the	e above					
(A) CO	(B) FeSiC	O <sub>3</sub> (C) MgSiO <sub>3</sub>	(D) CaSiO <sub>3</sub>	Q.15 Which	of the follo	owing is	not a ba	asic flux?	?	
Q.6 Which	alloy of alu	ıminium is used	in aircraft industry	(A) CaCO <sub>3</sub>	(B) CaO	(C)	SiO <sub>2</sub>	(D) M	gO	
(A) Duralu	min	(B) Magnaliur	n	<b>0 16</b> Zone r	efining is a	methor	to obt:	ain		
(C) Nickelo	су	(D) alunico		(A) Very high temperature						
				(B) Ultra pure Al						
<b>Q.7</b> The co	ommon imj	purities present	in the bauxite ore	(C) Ultra pure metal						
	and CuO	(B) Eq. $(O)$ and	PhO	(D) Ultra pure oxides						
(A) $Fe_2O_3$ (C) $Fe_2O_3$ (C)		(D) $\operatorname{Fe}_2 \operatorname{O}_3$ and (D) $\operatorname{SiO}_2$ and (	FDO							
(C) $\operatorname{Fe}_2 \operatorname{O}_3 \operatorname{d}$		(D) SIO <sub>2</sub> and $(D)$	200	<b>Q.17</b> Mac A	rthur proce	ess is use	e for			
<b>Q.8</b> The m	naterial used	d in solar cell co	ntains	(A) Ag	(B) Fe	(C)	Cl	(D) O <sub>2</sub>		
(A) Si	(B) Sn	(C) Ti	(D) Cs	<b>Q. 18</b> Use of	f electrolys	is is				
<b>Q.9</b> Cyanide process is used for the extraction of:		(A) Electropl	lating	(B) Ele	ectrorefi	ining				
(A) Ag	(B) Hg	(C) Cu	(D) Zn	(C) Both (A)	and (B)	(D) N	one of t	hese		
<b>Q.10</b> Durii	ng smelting	, an additional s	substance is added	<b>Q.19</b> Calcina	ation is use	ed in met	tallurgy	for remo	oval	of
which con	mbines wit	th impurities t	o form a fusible	(A) Water an	nd sulphide	(B) Wa	ater and	I CO <sub>2</sub>		
$(\Lambda)$ Slog		(C) Canqua		(C) CO <sub>2</sub> and	H <sub>2</sub> S	(D) H	$_{2}$ O and I	H <sub>2</sub> S		

(A) Slag (B) Mud (C) Gangue (D) Flux

<b>Q.20</b> In bla	st furnace,	maximun	n tempera	ature is in	Pre
(A) Zone of	fusion				01
(B) Zone of	combustio	n			Q. 1
(C) Zone of	slag comb	ustion			(A) (
(D) Zone of	reduction				(C) F
<b>Q.21</b> Of the	e following	g which	cannot b	be obtained by	Q.2
					(A) E
(A) Ag	(B) Mg and		(C) Cu	(D) Cr	(C) Z
<b>Q.22</b> Van Arconverting	rkel methoo the metal i	d of purifi nto a	cation of	metals involves	Q.3
(A) Volatile	stable com	pound			(A) k
(B) Volatile	unstable co	ompound			(C) C
(C) Non-vo	latile stable	e compou	Ind		
(D) None o	f the above				<b>Q.4</b> . is
					(A) C
Q.23 In the	froth float	ation pro	cess for	the purification	(B) A
(A) They are	ore partici	es noat p	ecause		(C) Z
(A) They are light			(D) N		
by wate	r	αιορπου	ic i.e., no	t easily wetted	0.5
(C) They be	ar electrost	tatic char	ge		<b>Q.5</b>
(D) They are	e insoluble				(A) (
<b>Q.24</b> Whicl	n method	of purific	ation is	represented by	(C) C
the followir	ng equatior	ns?		. ,	(A) I the o
Ti + 2I <sub>2</sub> —	<sup>773K</sup> →TiI₂	1675K	→Ti + 2	I <sub>2</sub>	(B) I
(A) C					not
(A) Cupella	tion	(B) Polir	ng		(C) I1
(C) Van Ark	el	(D) Zon	e refining	l	(D) l
Q.25 Calcin	ation is the	e process	of heatin	g the ore	06/
(A) In a blast furnace (B) In absence of air			auro		
(C) In prese	nce of air	(D) Nor	e of thes	e	Reas
<b>Q.26</b> Whic extracted b	h one of · y carbon re	the follo	wing me process?	tals cannot be	<b>Q.7</b> the r
(A) Pb	(B) Al	(C)	Hg	(D) Zn	Reas
<b>Q.27</b> Cyani	de process	is used fo	or obtain	ing	Q.8

(A) Cr	(B) Ag	(C) Cu	(D) Zn
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# **Previous Year's Questions**

	<b>Q.1</b> Which of the following mineral does not contain Al (1992)		
	(A) Cryolite	(B) Mica	(,
	(C) Feldspar	(D) Fluorspar	
by	Q.2 Which of the foll	lowing is not an ore	(1982)
	(A) Bauxite	(B) Malachite	
	(C) Zinc blende	(D) Pig iron	
lves	Q.3 Copper can be e	extracted from	(1978)
	(A) Kupfernickel	(B) Dolomite	
	(C) Galena	(D) Malachite	
	<b>Q.4</b> Among the follo is	wing statements, the inco	rrect one <b>(1997)</b>
	(A) Calamine and sid	erite are carbonates	
tion	(B) Argentite and cup	orite are oxides	
	(C) Zinc blende and	pyrites are sulphides	
ted	(D) Malachite and az	urite are ores of copper	
	<b>Q.5</b> Which ore conta	ins both iron and copper	(2005)
	(A) Cuprite	(B) Chalcocite	
	(C) Chalcopyrite	(D) Malachite	
by	(A) If both assertion the correct explanati	and reason are true and on of assertion, then mark	reason is : (A)
	(B) If both assertion not the correct expla	and reason are true but mation of assertion, then r	reason is nark (B)
	(C) If assertion is true	e but reason is false, then i	mark (C)
	(D) If both assertion	and reason are false, then	mark (D)
	<b>Q.6 Assertion:</b> Gold is aurocyanide complex	recovered from its solution coby adding zinc dust.	ontaining <b>(1982)</b>
	Reason: Zinc is more	e electropositive than gold	•
be	<b>Q.7 Assertion:</b> Zinc the recovery of Ag from the second s	is used and copper is no om the complex [Ag(CN),]	t used in <b>(1995)</b>
	Reason: Zinc is a pov	werful reducing agent that	n copper.
	Q.8 Assertion: Coke a	and flux are used in smelting	g. <b>(1998)</b>

**Reason:** The phenomenon in which ore is mixed with suitable flux and coke is heated to fusion is known as smelting.

Q.9 Assertion: Leaching is a process of reduction. (1999)

**Reason:** Leaching involves treatment of the ore with a suitable reagent so as to make it soluble while impurities remains insoluble.

Q.10 Assertion: Ethyl xanthate is used as a collector in froth floatation process. (1999)

**Reason:** Collectors depress the floatation property of one of the components of the ore and thus help in the separation of different minerals present in the same ore.

**Q.11** Which of the following factors is of no significance for roasting sulphide ores to the oxides and not subjecting the sulphide ores to carbon reduction directly? (2008)

(A) Metal sulphides are thermodynamically more stable than  $\mathsf{CS}_{\scriptscriptstyle 2}$ 

(B) CO<sub>2</sub> is thermodynamically more stable than CS<sub>2</sub>

- (C) Metal sulphides are less stable than the corresponding oxides
- (D)  $CO_2$  is more volatile than  $CS_2$

**Q.12** Which method of purification is represented by the following equation:  $Ti+2I_2 \rightarrow TiI_4$  (2012)

(A) Zone refining	(B) Cupellation
(C) Doling	$(\mathbf{D})$ $(\mathbf{D})$

(C) Poling (D) Van Arkel

**Q.13** In the context of the Hall-Heroult process for the extraction of Al, which of the following statements is false? (2015)

(A) CO and CO<sub>2</sub> are produced in this process

(B)  $Al_2O_3$  is mixed with  $CaF_2$  which lowers the melting point of the mixture and brings conductivity

(C)  $AI^{3+}$  is reduced at the cathode to form AI

(D) Na<sub>3</sub>AlF<sub>6</sub> serves as the electrolyte

**Q.14** Which one of the following ores is best concentrated by froth floatation method? (2016)

(A) Siderite (B) Galena

(C) Malachite (D) Magnetite

Q.15 Galvanization is applying a coating of(2016)(A) Cr(B) Cu(C) Zn(D) Rb

# **JEE Advanced/Boards**

## **Exercise 1**

Q.1 What is Goldschmidt Thermite Process?

 $\mathbf{Q.2}~\mathrm{Al_2O_3}$  can't be reduced by carbon to get Al metal. Explain.

**Q.3** Cu can reduce  $Ag^+$  to metallic Ag but Ag can't reduce  $Cu^{2+}$  to metallic Cu. Explain.

**Q.4** A metal is in combined state as sulphide. Identify the steps (A), (B), (C)



**Q.5** Write chemical equations for metallurgical processes to represent:

(a) Roasting of galena (PbS).

(b) Reduction of  $Cu_2O$  using charcoal as a reducing agent.

(c) deposition of pure silver from an aqueous solution of  $Ag^{+}$ .

Q.6 Why zinc and not copper is used for the

recovery of metallic silver from the complex [Ag(CN),]?

**Q.7** Sodium is prepared by the electrolysis of molten NaCl but not by the electrolysis of its aqueous solution.

**Q.8** Elements of alkali metal group are strong reducing agent. Explain.

**Q.9** In the metallurgy of iron, why limestone is added to the ore.

**Q.10** What is the actual reducing agent of haematite in blast furnace?

**Q.11** Write chemical equations to represent the most probable outcome in each of the following. If no reaction is likely to occur, state so:

(i)  $CdCO_3 \xrightarrow{\Delta}$ (ii) MgO  $\xrightarrow{\Delta}$ (iii) SnO<sub>2</sub> + CO  $\xrightarrow{\Delta}$ (iv) CdSO<sub>4</sub> (aq) <u>electrolysis</u> (v) 2HgO  $\xrightarrow{\Delta}$ (vi) MgO + Zn  $\xrightarrow{\Delta}$ 

Q.12 Why graphite is used as anode but not diamond?

**Q.13** The following reactions take place during the extraction of copper from copper ore:

(i)  $2Cu_2S(\ell) + 3O_2(g) \rightarrow 2Cu_2O(\ell) + +2SO_2(g)$ 

(ii)  $2Cu_2O(\ell) + Cu_2S(\ell) \rightarrow 6Cu(\ell) + SO_2(g)$ 

Identify the oxidizing and reducing agents.

**Q.14** Dolomite (MgCO<sub>3</sub>.CaCO<sub>3</sub>) can also be treated to get MgCl<sub>2</sub> which in turn is electrolyzed to get Mg. (Dow natural brine process). Give reactions of this process.

**Q.15**  $[CaCO_3.MgCO_3] \xrightarrow{\Delta} CaO.MgO$ CaO. MgO  $\xrightarrow{Fe/Si}$  Mg + Fe + Ca<sub>2</sub>SiO<sub>4</sub> Name this process.

Q.16 Convert Mg into MgO.

**Q.17** (i) A black mineral (A) on treatment with dilute sodium cyanide solution in presence of air gives a clear solution of (B) and (C).

(ii) The solution of (B) on reaction with zinc gives precipitate of a metal (D).

(iii) (D) is dissolved in dilute  $HNO_3$  and the resulting solution gives a white precipitate (E) with dilute HCl.

(iv) (E) on fusion with sodium carbonate gives (D).

(v) (E) dissolves in aqueous solution of ammonia giving a colourless solution of (F). Identify compounds (A) to (F) and give chemical equations for reactions in steps (i) to (iv). **Q.18** Name two metals which are used for reduction in metallurgical process. Give one chemical equation for each

**Q.19** Aluminium metal is frequently used as a reducing agent for the extraction of metals such as chromium or manganese from their respective oxides. Why?

**Q.20** What is the function of basic furnace linings in steels manufacture?

**Q.21** Write down reactions involved in the extraction of Pb. What is the oxidation number of lead in litharge?

**Q.22**  $A_1$  and  $A_2$  are two ores of metal M.  $A_1$  on calcination gives black precipitate, CO<sub>2</sub> and water?



**Q.23** Gold is also extracted by cyanide process as in case of silver. Outline the reactions.

Q.24 How is Ag extracted from silver coin?

**Q.25** In the purification of bauxite ore as preliminary step in the production of Al,  $[Al(OH)_4]^-$  can be converted to  $Al(OH)_3$  by passing CO<sub>2</sub> through it. Write an equation for the reaction that occurs.

**Q.26** When the ore hematite is burnt in air with coke around 2000°C along with lime, the process not only produces steel, but also produces a silicate slag that is useful in making building material such as cement. Discuss the same and show through balanced chemical equation.

#### Q.27 Explain the following

(i) In the metallurgy of iron, lime-stone is added to the ore.

(ii) Although Au is soluble in aqua-regia, Ag is not.

## **Exercise 2**

## Single Correct Choice Type

<b>Q</b> .1	Sulphide	ores a	re ger	nerally	concentrated	by
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(A) Froth floatation (E	B) Roasting
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(C) Gravity (D) Reduction by carbon

**Q.2** Which method of purification is represented by the following equations?

$Ti + 2I_2 \xrightarrow{523K} TiI_4$	$\xrightarrow{1700K} Ti + 2I_2$
(A) Cupellation	(B) Poling
(C) Van Arkel	(D) Zone refining

## Q.3 Cryolite is

(A)  $Na_3AIF_6$  and is used in the electrolysis of alumina for decreasing electrical conductivity

(B)  $Na_3AIF_6$  and is used in the electrolysis of alumina for lowering the melting point of alumina

(C)  $Na_3AIF_6$  and is used in the electrolytic purification of alumina

(D) Na<sub>3</sub>AlF<sub>6</sub> and is used in the electrolysis of alumina

Q.4 Cassiterite is an ore of

(A) Mn (B) Ni (C) Sb (D) Sn

**Q.5** When an aqueous solution of sodium chloride is electrolysed using platinum electrodes, the ions discharged at the electrodes are

(A) Sodium and hydrogen

(B) Sodium and chloride

(C) Hydrogen and chloride

(D) Hydroxyl and chloride

Q.6 The luster of a metal is due to

(A) Its high density

(B) Its high polishing

(C) Its chemical inertness

(D) Presence of free electrons

Q.7 Which metal can't be obtained from electrolysis?

(A) Ca (B) Mg

(C) Cr (D) All of the above

- Q.8 The function of flux during smelting of ores is to:
- (A) Make the ore porous
- (B) Remove gangues
- (C) Facilitate reduction
- (D) Precipitate slag

**Q.9** The chemical composition of slag formed during the smelting process in the extraction of copper is:

(A) Cu <sub>2</sub> O + FeS	(B) FeSiO <sub>3</sub>
(C) CuFeS <sub>2</sub>	(D) Cu <sub>2</sub> S + FeO

**Q.10** The chemical process in the production of steel from haemotite ore involves:

- (A) Reduction
- (B) Oxidation

(C) Reduction followed by oxidation

(D) Oxidation followed by reduction

**Q.11** In the commercial electrochemical process for alumniuim extraction, the electrolyte used is

(A) Al  $(OH)_3$  in NaOH solution

(B) An aqueous solution of  $Al_2(SO_4)_3$ 

- (C) A molten mixture of Al<sub>2</sub>O<sub>3</sub> and Na<sub>3</sub>AlF<sub>6</sub>
- (D) A molten mixture of AlO(OH) and Al(OH)<sub>3</sub>

Q.12 Calcination is used in metallurgy to remove

(A) $H_2O$ and $H_2S$	(B) $H_2O$ and $CO_2$
(C) $CO_2$ and $H_2S$	(D) $H_2O$ and $H_2S$ .

Q.13 The metallic luster exhibited by sodium is

explained by the presence of

(A) Na<sup>+</sup> ions

- (B) Conducting electrons
- (C) Free protons
- (D) A body-centred cubiclattice.

Q.14 Stainless steel contains

(A) Fe, Cr, Cu	(B) Fe, Cr, Ni
(C) Fe, Ni, C	(D) Fe, Ni, Cu

**Q.15** Which one of the following beneficiation processes is used for the mineral,  $AI_2O_3.2H_2O$ ?

(A) Froth floatation	(B) Leaching
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(C) Liquation (D) Magnetic separation

**Q.16** The extraction of which of the following metals involves bessemerisation ?

(A) Fe (B) Ag (C) Al (D) Cu

**Q.17** One of the characteristic properties of non-metals is that they

(A) Are reducing agents

(B) Form basic oxides

(C) Form cations by electron gain

(D) Are electronegative

**Q.18** In the froth floatation process for the purification of ores, the ore particles float because

(A) They are light

(B) Their surface is hydrophobic i.e., not easily wetted by water

(C) They bear electrostatic charge

(D) They are insoluble

Q.19 Which is not a basic flux?

(A) CaCO<sub>3</sub> (B) Lime (C) SiO<sub>2</sub> (D) CaO

**Q.20** The substance which is added to remove impurities is known as

(A) Slag (B) Flux (C) Gangue (D) Catalyst

**Q.21** Electrolytic reduction method is used in the extraction of

(A) Highly electronegative elements

(B) Highly electropositive elements

(C) Transition metals

(D) Noble metals

Q.22 The role of calcination in metallurgical operations is

(A) To remove moisture

(B) To decompose carbonate

(C) To drive off organic matter

(D) To achieve all the above

#### **Comprehension Type**

A black coloured compound (A) on reaction with dil  $H_2SO_4$  form a gas 'B' and a solution of compound (C). When gas B is passed through solution of compound (C), a black coloured compound A is obtained which is soluble in 50% HNO<sub>3</sub> and forms blue coloured complex 'D' with excess of  $NH_4OH$  and chocolate brown ppt. 'E' with  $K_4$ [Fe(CN)<sub>6</sub>]

<b>Q.23</b> 'A' is				
(A) CuS	(B) FeS	(C) PbS	(D) HgS	
<b>Q.24</b> 'B' is				
(A) H <sub>2</sub> S	(B) SO <sub>2</sub>	(C) NH <sub>3</sub>	(D) SO <sub>3</sub>	
<b>Q.25</b> 'C' is				
(A) CuS	(B) CuSO <sub>4</sub>	(C) Cu(NO <sub>3</sub> ) <sub>2</sub>	(D) HgSO <sub>4</sub>	
<b>Q.26</b> 'D' is				
(A) Cu(OH) <sub>2</sub>		(B) [Cu(NH <sub>3</sub> ) <sub>2</sub> ]SO	4	
(C) [Cu(NH <sub>3</sub> ) <sub>4</sub> ] (NO <sub>3</sub> ) <sub>2</sub>		(D) [Cu(NH <sub>3</sub> ) <sub>6</sub> ]SO	4	
<b>Q.27</b> 'E' is				
(A) Cu <sub>2</sub> [Fe(CN) <sub>6</sub> ]		(B) Cu <sub>4</sub> [Fe(CN) <sub>6</sub> ]		
(C) $Cu_3[Fe(CN)_6]_2$		(D) None of thes	e	
		-		

#### **Assertion Reasoning Type**

(A) If both assertion and reason are true and reason is the correct explanation of assertion, then mark (a)

(B) If both assertion and reason are true but reason is not the correct explanation of assertion, then mark (b)

(C) If assertion is true but reason is false, then mark (c)

(D) If both assertion and reason are false, then mark (d)

**Q.28 Assertion:** Calamine is a ore of boron. **Reason:** Boric acid is a tribasic acid.

**Q.29 Assertion:** FeS<sub>2</sub> is also known as fools' gold. **Reason:** FeS<sub>2</sub> has yellow metallic appearance.

**Q.30 Assertion:** Pb<sup>4+</sup> can be reduced easily to Pb<sup>2+</sup>. **Reason:** Pb<sup>2+</sup> is paramagnetic.

**Q.31 Assertion:** Diamond is harder than graphite.

Reason: Graphite is more stable than diamond.

**Q.32 Assertion:** Carbon dioxide and nickel react to form tetracaronyl nickel (0).

**Reason:**  $Ni(CO)_4$  decomposes on heating to form Ni and CO.

## **Previous Years Questions**

Q.1 Which ore contains both iron and copper (2005)

(A) Cuprite	(B) Chalcocite

(C) Chalcopyrite (D) Malachite

Q.2 Extraction of zinc from zinc blende is achieved by

(A) Electrolytic reduction

(B) Roasting followed by reduction with carbon

(C) Roasting followed by reduction with another metal

(D) Roasting followed by self-reduction

**Q.3** Native silver metal forms a water soluble complex with a dilute aqueous solution of NaCN in the presence of (2008)

(A) Nitrogen	(B) Oxygen
(C) Carbon dioxide	(D) Argon

**Q.4** Oxidation states of the metal in the minerals haematite and magnetite, respectively, are (2011)

(A) II, III in haematite and III in magnetite

(B) II, III in haematite and II in magnetite

(C) II in haematite and II, III in magnetite

(D) III in haematite and II, III in magnetite

Q.5 Addition of high proportions of manganese makes steel useful in making rails (1998)

(A) Gives hardness to steel

(B) Helps the formation of oxides of iron

(C) Can remove oxygen and sulphur

(D) Can show highest oxidation state of +7

Q.6 Extraction of metal from the ore cassiterite involves (2011)

(A) Carbon reduction of an oxide ore

(B) Self-reduction of a sulphide ore

(C) Removal of copper impurity

(D) Removal or iron impurity

### Paragraph

(2007)

Copper is the most noble of the first row transition metals and occurs in small deposits in several countries. Ores of copper include chalcanthite ( $CuSO_4$  .5H<sub>2</sub>O), atacamite ( $Cu_2Cl(OH)_3$ , cuprite ( $Cu_2O$ ), copper glance ( $Cu_2S$ ) and malachite ( $Cu_2(OH)_2CO_3$ ). However, 80% of the world copper production comes from the ore chalcopyrite ( $CuFeS_2$ ). The extraction of copper from chalcopyrite involves partial roasting, removal of iron and self-reduction. (2010)

**Q.7** Partial roasting of chalcopyrite produces

(A) Cu <sub>2</sub> S and FeO	(B) Cu <sub>2</sub> O and FeO
(C) CuS and $Fe_2O_3$	(D) $Cu_2O$ and $Fe_2O_3$

Q.8 Iron is removed from chalcopyrite as

(A) FeO (B) FeS (C)  $Fe_2O_3$  (D)  $FeSiO_3$ 

**Q.9** In self-reduction, the reducing species is

(A) S (B)  $O^{2-}$  (C)  $S^{2-}$  (D)  $SO_{2}$ 

Q.10 Match the following Metals listed in column I with extraction processes listed in column II. (1979)

Column I	Column II
(A) Silver	(p) Fused salt Electrolysis
(B) Calcium	(q) Carbon Reduction
(C) Zinc	(r) Carbon monoxide reduction
(D) Iron	(s) Amalgamation
(E) Copper	(t) Self reduction

**Q.11** Write the chemical reactions involved in the extraction of silver from argentite. (2000)

**Q.12** Write the balanced chemical equation for developing photographic films.

**Q.13** Write balanced chemical equation for developing a black and white photographic film. Also give reason, why the solution of sodium thiosulphate on acidification turns milky white and give balance equation of this reaction. (2005)

**Q.14** Given the number of water molecule (s) directly bonded to the metal center in  $CuSO_4$ .  $5H_2O$ . **(2009)** 

**Q.15** Given the coordination number of Al in the crystalline state of  $AlCl_3$ . **(2009)** 

Q.16 Extraction of zinc from zinc blende is achieved by,

(A) Electrolytic reduction

(B) Roasting followed reduction by reduction with carbon

(C) Roasting followed by reduction with another metal

(D) Roasting following by self-reduction

Q.17 Match the conversions in column I with the type (s) of reaction (S) give in column II. (2008)

Column I	Column II
(A) $PbS \rightarrow PbO$	(p) Roasting
(B) $CaCO_3 \rightarrow CaO$	(q) Calcination
(C) $ZnS \rightarrow Zn$	(r) Carbon reduction
(D) $Cu_2 S \rightarrow Cu$	(s) Self reduction

Q.18 Sulfide ores are common for the metals (2013)

- (A) Ag, Cu and Pb (B) Ag, Cu and Su
- (C) Ag, Mg and Pb (D) Al, Cu and Pb

Q.19 The carbon-based reduction method in NOT used for the extraction of (2013)

(A) Tin from  $SnO_2$ 

(2007)

- (B) Iron from  $Fe_2O_3$
- (C) Aluminium from  $Al_2O_3$
- (D) Magnesium from MgCO<sub>3</sub>.CaCO<sub>3</sub>

**Q.20** Match the anionic species given in column I that are present in the ore (s) given in column II (2015)

Column I	Column II
(A) Carbonate	(p) Siderite
(B) Sulphide	(q) Malachite
(C) Hydroxide	(r) Bauxite
(D) Oxide	(s) Calamine
	(s) Argentite

Q.21 Extraction of copper from copper pyrite  $(CuFeS_2)$ involves(2016)

(A) Crushing following by concentration of the ore by froth-flotation

(B) Removal of iron as slag

(C) Self-reduction step to produce 'blister copper' following evolution of  $SO_2$ 

(D) Refining of 'blister copper' by carbon reduction

# **MASTERJEE Essential Questions**

# JEE Main/Boards

## **JEE Advanced/Boards**

Exersice 1			Exersice	1			
Q.2	Q.6	Q.15	Q.4	Q.12	Q.17		
Q.21	Q.27	Q.30	Q.20	Q.27			
Exersice 2			Exersice	Exersice 2			
Q.8	Q.24		Q.2	Q.13	Q.32		
Previous Years' Questions		Previous	Previous Years' Questions				
Q.9			Q.3	Q.10	Q.15		

	Answer Key						
JEE Mair	s/Boards						
Exercise 2							
Single Correct	Choice Type						
<b>Q.1</b> B	<b>Q.2</b> C	<b>Q.3</b> A	<b>Q.4</b> A	<b>Q.5</b> D	<b>Q.6</b> A	<b>Q.7</b> C	
<b>Q.8</b> A	<b>Q.9</b> A	<b>Q.10</b> D	<b>Q.11</b> A	<b>Q.12</b> D	<b>Q.13</b> A	<b>Q.14</b> A	
<b>Q.15</b> C	<b>Q.16</b> C	<b>Q.17</b> A	<b>Q.18</b> C	<b>Q.19</b> B	<b>Q.20</b> B	<b>Q.21</b> B	
<b>Q.22</b> A	<b>Q.23</b> A	<b>Q.24</b> C	<b>Q.25</b> B	<b>Q.26</b> B	<b>Q.27</b> A		
Previous Years' Questions							
<b>Q.1</b> D	<b>Q.2</b> D	<b>Q.3</b> D	<b>Q.4</b> B	<b>Q.5</b> C	<b>Q.6</b> A	<b>Q.7</b> B	
<b>Q.8</b> B	<b>Q.9</b> D	<b>Q.10</b> C	<b>Q.11</b> A	<b>Q.12</b> D	<b>Q.13</b> D	<b>Q.14</b> B	
<b>Q.15</b> C							

**JEE Advanced/Boards** 

Exercise 2

Single Correct	Choice Type					
<b>Q.1</b> A	<b>Q.2</b> C	<b>Q.3</b> B	<b>Q.4</b> D	<b>Q.5</b> C	<b>Q.6</b> D	<b>Q.7</b> C
<b>Q.8</b> D	<b>Q.9</b> B	<b>Q.10</b> D	<b>Q.11</b> C	<b>Q.12</b> C	<b>Q.13</b> C	<b>Q.14</b> B
<b>Q.15</b> B	<b>Q.16</b> D	<b>Q.17</b> D	<b>Q.18</b> B	<b>Q.19</b> C	<b>Q.20</b> B	<b>Q.21</b> B
<b>Q.22</b> D						
Comprehensio	n Type					
<b>Q.23</b> A	<b>Q.24</b> A	<b>Q.25</b> B	<b>Q.26</b> C	<b>Q.27</b> A		
Assertion Reas	oning Type					
<b>Q.28</b> D	<b>Q.29</b> C	<b>Q.30</b> C	<b>Q.31</b> C	<b>Q.32</b> D		
Previous Years' Question						
<b>Q.1</b> C	<b>Q.2</b> B	<b>Q.3</b> B	<b>Q.4</b> D	<b>Q.5</b> A, C	<b>Q.6</b> A, D	<b>Q.7</b> B
<b>Q.8</b> D	<b>Q.9</b> C	<b>Q.10</b> A $\rightarrow$ s; B –	$\rightarrow$ p; C $\rightarrow$ q; D $\rightarrow$	q, r; E $\rightarrow$ t	<b>Q.16</b> B	
<b>Q.17</b> A → p; B	$\rightarrow$ q; C $\rightarrow$ p, r; D	$\rightarrow$ p, s	<b>Q.18</b> A	<b>Q.19</b> C, D		
<b>Q.20:</b> $A \rightarrow p$ , q,	s; $B \rightarrow t$ ; $C \rightarrow q$ ,	r; D → r	<b>Q.21</b> A, B, C			

# **Solutions**

# JEE Main/Boards

## **Exercise 1**

**Sol 1:** The reduction potential of zinc and iron is lower than that of copper. In hydrometallurgy, Zn/Fe can be used to displace Cu from the solution

$$Fe_{(s)} + Cu^{2+}_{(aq.)} \longrightarrow Fe^{2+}_{(aq.)} + Cu_{(s)} \downarrow$$

But to displace Zn, more reactive metals i.e. Na, K are required but they violently react with water and gives hydrogen gas.

$$K + H_2O \longrightarrow KOH + \frac{1}{2}H_2 \uparrow$$

 $\therefore$  By hydrometallurgy, Cu can be extracted but not Zn.

#### Sol 2: Ellingham diagrams.

In this range of temperature, diagrams show that carbon monoxide act as stronger reduction agent than carbon.



 $2CO + O_2 \rightarrow 2CO_2$ 

has more negative free energy than

 $2C + O_2 \rightarrow 2CO$ 

 $\therefore$  In the upper part of furnace, haematite is reduced by CO even in the presence of C.

**Sol 3:** In case of low grade copper ores, such as chalcopyrites, Ag is used to reduced the remaining copper.

$$Cu^{2+} + 2Ag \rightarrow Cu + 2Ag^{+}$$

and by formation of shiny  $Ag_2S$  layer, it is confirmed that Cu got reduced.

**Sol 4:** Ellingham diagram clearly indicates that the CO graph cut at much higher temperature to Zinc oxide's graph.



 $T_0 > T_{B,P \text{ of } Zn}$  So, Zn will be in vapour form and its very tough to maintain such high temperature in furnace for long time and at this temperature, carbon forms carbides.

**Sol 5:** Element present in the anode mud:- Se, Te, Ag, Au, Pt and Sb

These elements are very less reactive and are not affected during the purification process. Hence, they settle down below anode as anode mud.

**Sol 6:** In the process of Bessemerisation, air is blown into molten matte with silica. So, by self reduction CuS gets reduced to Cu but at same time, FeS also gets converted to FeO but to avoid further reaction, silica is added.

FeO + SiO<sub>2</sub> 
$$\rightarrow$$
 FeSiO<sub>3</sub> (Slag)  
Thomas slag

By this, molten copper gets separated out.

Sol 7: Refer Theory - Chromatography

**Sol 8:** The composition difference between cast and Pig iron is, Pig iron has 4% carbon content whereas cast iron has at least 2% carbon content.



After extracting from furnace, we get pig iron and we put them in a case from which impurities can be cut out so there is difference of amount in impurity. **Sol 9:** Minerals are the substances in which metals are present either in native or combined forms but ores are that minerals from which metal can be extracted economically and conveniently.

**Sol 10:** From Ellingham diagram, graph of C cuts the Zn graph at lower temperature than CO.



 $\therefore$  ZnO + C  $\rightarrow$  Zn + CO,  $\Delta G_1$ 

 $ZnO + CO \rightarrow Zn + CO_{2'} \Delta G_{2}$ 

 $|\Delta G_1| > |\Delta G_2|$  so, first reaction is favourable

 $\therefore$  C is better reducing agent than CO.

**Sol 11:** Graphite rod is used as an anode and graphite line iron as cathode during the electrolysis of alumina.

Sol 12: Refer theory

Sol 13: Refer theory

**Sol 14:** Copper matte contains sulfides of copper and iron. While reducing, it is Bessemer converter by self-reduction, CuS gets converted to Cu but to avoid Fe in the final product, silica is used which forms slag with FeO

 $FeO + SiO_2 \rightarrow FeSiO_3$ (slag)

**Sol 15:** Generally, aluminium is extracted by using electrometallurgy, but for that purpose Al should be in molten state. But, melting point of aluminium is very high. So, to increase the solubility and to decrease the melting point, cryolite is added.

**Sol 16:** Roasting is a process in which ore is heated in presence of oxygen to convert it into corresponding oxides. So the ores whose oxides can be easily reduced by C/CO or by self reduction are roasted. E.g. CuS, FeS, but ores of Al, Na, K cannot be roasted.

**Sol 17:** (a) Acidic flux:- Used to remove basic gangues

E.g. SiO<sub>2</sub>, Na<sub>2</sub>B<sub>4</sub>O<sub>7</sub>.10H<sub>2</sub>O

(b) Basic flux:- Used to remove acidic gangues

E.g. CaO, MgCO<sub>3</sub>, CaCO<sub>3</sub>.

Sol 18: Gangue:- (Matrix)

These are the non-metallic impurities present in the ore. In fact, impurities associated with an ore is known as gangue.

Sol 19: Refer theory

**Sol 20:** (1) Zone Refining: Fractional crystallisation :-Based on difference in solublities of impurities in molten and solid state.

E.g. Ge, Ga, Si etc (To extrapurify the metal)

(2) Electrolytic Refining: Faraday's law of electrolysis

Metals which are not reduced by C/CO, are reduced by this method. E.g. Na, Ca, K, Al etc.

(3) Vapour phase refining:

Ni+4CO $\rightarrow$  [Ni(CO)<sub>4</sub>]  $\xrightarrow{\Lambda}$  Ni $\downarrow$  +4CO Purification by forming volatile carbonyl complex.

**Sol 21:** Froath floatation:- Based on the different wetting property of the ore and gangue with oil and water. Depressants are used to depress the frothing tendency of the frothing agent Eg. NaCN, KCN

Example: In the selective froth floatation of PbS, ZnS can be depressed using depressant.

$$Zn + 4CN^{-} \rightarrow [Zn(CN)_4]^2$$

**Sol 22:** Copper pyrites  $\rightarrow$  CuFeS<sub>2</sub> composes sulfides of CuS and FeS both.

While extracting copper from pyrites, when we oxidise the ore 2FeS +  $3O_2 \rightarrow 2FeO+2SO_2$ , ferrous oxide also gets formed which has to be removed by slag formation

$$\begin{array}{c} CuS + O_2 \rightarrow CuO + SO_2 \\ \hline \\ 2CuO + CuS \rightarrow 3Cu + SO_2 \end{array} \qquad Self \\ reduction \end{array}$$

But in simple oxide 'CuO', it can be easily reduced by carbon reduction

$$CuO + C \rightarrow Cu + CO_{2}$$

**Sol 23:** (i) Zone Refining: Method to obtain metals of very high purity metals like Ge, Ga, Si etc. This method is based on difference in solubilities of impurities in molten and solid state. This refining is as shown below:



Heater melts that part of metal and impurities flows in the molten part and by moving this, eventually get at the end and cut out, this process is repeated several times.

(ii) Column Chromatography:- This technique is based on different adsorbing power of different metal ions on a surface. This technique is used for that metals whose separation is difficult. Eg. Lanthanoids, actinoids.

**Sol 24:** At 600-900°C:  $Fe_2O_3+3CO \rightarrow 2Fe+3CO_2$ At 900-1000°C ;  $CaCO_3 \rightarrow CaO + CO_2$  ;  $CO_2 + C \rightarrow 2CO$ 

**Sol 25:** Extraction of any ore is done in 3 parts.

(i) Concentration:- froth floatation process.

(ii) Roasting:-

$$2ZnS + 3O_2 \longrightarrow 2ZnO + 2SO_2$$

Carbon Reduction

(iii) Purification:- fractional distillation & electrolysis.

#### Sol 26: Refer theory

**Sol 27:** Silica in alumina is a natural impurity and before electrolysis, this impurity should be removed. So any of the following chemical process can be used to purify alumina from bauxite.

(i) Bayer's process:- Finely powdered

Bauxite 
$$\xrightarrow{\text{Roasted}}_{\text{FeO}\rightarrow\text{Fe}_2O_3}$$
 Roasted ore



Filtrate (Sod. aluminate + Sod. Silicate)  $\xrightarrow{CO_2}$ Al(OH)<sub>3</sub>  $\xrightarrow{\Delta}$  Pure Al<sub>2</sub>O<sub>3</sub>

(ii) Hall's process:

Finely Bauxite:

Finely Bauxite:-

 $\begin{tabular}{|c|c|c|c|} Na_2CO_3 & Solution+ \\ Fused, extracted & (Residue SiO_2 + Fe_2O_3) \\ with water & \\ \end{tabular}$ 

$$\begin{array}{c} \underline{CO_2} \\ 50^{\circ} - 60^{\circ} \& \text{ filtered} \\ \hline \\ \text{Filtrate (Na_2CO_3)} \end{array} \rightarrow \text{ppt. Al(OH)_3} \\ \downarrow \\ \Delta \\ Pure Al_2O_3 \end{array}$$

(iii) Serpeck's Process:-

Finely powdered bauxite ore  $\xrightarrow{\text{Coke}+\text{N}_2}{\Delta \text{ to } 1800^{\circ}\text{C}}$ silica reduced + AIN to Si which volatises  $H_2\text{O}$ Pure Al<sub>2</sub>O<sub>3</sub>  $\xleftarrow{}^{\Delta}$  Al(OH)<sub>3</sub>

Sol 28: Refer Elingham diagram

**Sol 29:** Yes, Reducing agent's selection widely depends on thermodynamic factor because free energy is the factor which decides the reaction feasibility.

E.g. (1) In the upper part of blast furnace, carbon monoxide is better reducing agent than carbon because in that temperature range,  $\Delta G$  of CO's reaction is less than for C.

(2) Reduction of ZnO is not feasible by carbon because it can be only possible at high temperature at which both compounds are in vapour phase.

**Sol 30:** In the electrolysis of NaCl,  $Cl_2$  is obtained as a by-product

Sol 31: Ellingham Diagram:-



Reduction of MgO by Al is feasible at very high temperature around 1750°C

 $2AI+3MgO \rightarrow 3Mg\downarrow + Al_2O_3$ 

Because  $\Delta G$  is negative after this temperature. So, aluminothermite is not useful to extract Mg from MgO. Thus, electrolytic reduction is used.

## **Exercise 2**

**Sol 1: (B)** During roasting, air is blown to convert metal sulphides into metal oxides.

$$FeS + \frac{3}{2}O_2 \rightarrow FeO + SO_2$$

Sol 2: (C) Cu can be reduced by self-reduction. So,

 $Cu_2S + \frac{3}{2}O_2 \rightarrow Cu_2O + SO_2$ ;  $2Cu_2O + Cu_2S \rightarrow 6Cu + SO_2$ 

**Sol 3: (A)** Pyrolusite  $\rightarrow$  MnO<sub>2</sub>

Sol 4: (A) Ag, Au, Hg, Pt are chemically inert metals.

**Sol 5: (D)**  $C_{ao} + SiO_2 \longrightarrow CaSiO_3$ 

**Sol 6: (A)** Duralumin an alloy of aluminium is used in aircraft industry

**Sol 7: (C)** Bauxite ore impurities:-  $SiO_2 + Fe_2O_3$ 

Sol 8: (A) Si is a semi-conductor.

Sol 9: (A) Ag + 2CN<sup>-</sup> 
$$\longrightarrow$$
 Ag(CN)<sub>2</sub><sup>-</sup>  
 $\downarrow$   
Zn, 2CN<sup>-</sup>  
Ag + [Zn(CN)<sub>4</sub>]<sup>2-</sup>

**Sol 10: (D)** Flux + Gangue  $\rightarrow$  slag.

**Sol 11: (A)** Zone refining:- Purification is done at microscopic level because we need ultra-purified element to use in semi-conductors. Zone refining is based on fractional crystallisation.

**Sol 12: (D)** None of this elements can be reduced by Al but other example are Cr, Mn, Ti, etc.

**Sol 13: (A)** Anode mud are the elements which get deposited in the bottom of anode Eg. Ag, Au, Pt.

**Sol 14: (A)** Poling:- Molten metal is stirred with green anode poles which releases hydrocarbon (gas) which reduces the oxide e.g. Cu

$$Cu_2O + CO \rightarrow 2Cu + CO_2$$
;  $Cu_2O + H_2 \rightarrow 2Cu + H_2O$ 

**Sol 15: (C)** SiO<sub>2</sub> is an acidic flux  $SiO_2 + CaO \longrightarrow CaSiO_3$ 

**Sol 16: (C)** Zone refining is a method to obtain ultra pure metal.

**Sol 17: (A)** Mac Arthur process is basically cyanide process.

**Sol 18: (C)** Electroplating: To deposit a specific metal on some other to protect the inner ore is done by electrolysis.

Electrorefining: Electrochemical series decide which metal to be deposited first on anode and using it, we can remove the impurities by electrolysis.

**Sol 19: (B)** Calcination: Ore is heated at high temperature in the absence of air.

E.g. Applied for carbonate and hydrated ores.

 $CuCO_3.Cu(OH)_2 \rightarrow CuO + CO_2 + H_2O$ 



**Sol 21: (B)** Mg and Al's aqueous solution on electrolysing gives hydrogen at anode.

.:. Molten mixture is used.

**Sol 22: (A)** Van-Arkel method:- Metal is converted into a volatile stable compound (e.g., lodine) and impurities are not affected during compound formation

E.g. Zr, Ti  
Ti + 2I<sub>2</sub> 
$$\rightarrow$$
 TiI<sub>4</sub> (g)  
Ti I<sub>4</sub>(g)  $\xrightarrow{1700K}$  Ti<sub>(S)</sub> + 2I<sub>2(g)</sub>

**Sol 23: (A)** In the froth floatation process for the purification of ores, the ore particles float because they are light.

Sol 24: (C) Van arkle method is used.

**Sol 25: (B)** Calcination is the process of heating the ore in absence of air.

**Sol 26: (B)** Highly electropositive metal can not be reduced because it requires high temperature at which carbide formation occurs.

Sol 27: (A) Cyanide process is used for obtaining Cr.

## **Previous Years' Questions**

**Sol 1: (D)** Fluorspar (CaF<sub>2</sub>), Cryolite (Na<sub>3</sub>AlF<sub>6</sub>), Feldspar (KAlSi<sub>3</sub>O<sub>8</sub>), Mica (K<sub>2</sub>O.3Al<sub>2</sub>O<sub>3</sub>.6SiO<sub>2</sub>.2H<sub>2</sub>O).

**Sol 2: (D)** Pig iron  $\rightarrow$  It is the most impure form of iron and contains highest proportion of carbon (2.5 – 4%)

Malachite  $\rightarrow$  Cu(OH)<sub>2</sub>.CuCO<sub>3</sub>

Zinc blende  $\rightarrow$  ZnS

 $Bauxite \rightarrow Al_2O_3.2H_2O$ 

**Sol 3: (D)** Malachite (Cu(OH)<sub>2</sub>.CuCO<sub>3</sub>).

**Sol 4: (B)** Cuprite (Cu<sub>2</sub>O) and Argentite (Ag<sub>2</sub>S).

**Sol 5: (C)** Among cuprite [Cu<sub>2</sub>O], Chalcocite [Cu<sub>2</sub>S], Chalcopyrite [CuFeS<sub>2</sub>] & Malachite [Cu(OH)<sub>2</sub>.CuCO<sub>3</sub>].

**Sol 6: (A)** Au is recovered from the solution by the addition of electropositive metal.

 $2NaAu(Cu)_2 + Zn \rightarrow Na_2Zn(CN)_4 + 2Au^{1}$ Soluble complex

**Sol 7: (B)** Both assertion and reason are true and reason is the correct explanation of assertion.

**Sol 8: (B)** Both assertion and reason are true but reason is not the correct explanation of assertion. Non-fusible mass present in ore in mixing with suitable flux are fused which are then reduced by coke to give free metal.

Sol 9: (D) Leaching is a process of concentration.

**Sol 10: (C)** Assertion is true but reason is false. Collectors absorb themselves on polar groups to grains of ores and thus derive them on the surface to pass on into the froth.

**Sol 11: (A)** Metal sulphides are thermodynamically more stable than CS<sub>2</sub>

Sol 12: (D) Van Arkel method

$$\begin{aligned} &\mathsf{Ti}(s) + 2I_{2}(g) \xrightarrow{523\mathsf{K}} \mathsf{Til}_{4}(g) \\ &\mathsf{Til}_{4}(g) \xrightarrow{1700\mathsf{K}} \mathsf{Ti}(s) + 2I_{2}(g) \end{aligned}$$

**Sol 13: (D)** 13 In the context of the Hall-Heroult process for the extraction of Al,  $Na_3AlF_6$  serves as the electrolyte

Sol 14: (B) Galena = Pbs

For sulphur ores froth floatation is carried out.

Sol 15: (C) Galvanization is applying a coating of Zn.

## **JEE Advanced/Boards**

## **Exercise 1**

**Sol 1:** Goldschmidt thermite process:- Reduction by powdered aluminium. This process is employed in cases where metals have very high m.pt and are to be extracted from their oxides.

Process:-

 $Mg + BaO_2 \rightarrow BaO + MgO + Heat \rightarrow$ (in large amount) Ignition mixture

$$Cr_2O_3 + AI \xrightarrow{\Delta} 2Cr(\ell) + AI_2O_3 + Heat$$

E.g. Reduction of Cr, Ti, W, Mn etc.



**Sol 2:**  $Al_2O_3 + C \rightarrow 2Al + CO_2$ 

 $\Delta G$  of this reaction is negative at very high temperature, so there are 2 reasons that it is not useful:-

(i) Such high temperature is difficult to achieve in furnace for very long time and at such temperature,  $Al_2O_3$  will also be in vapour phase.

(ii) At high temperature, C instead of reducing forms carbide  $Al_4C_3$  with aluminium.

**Sol 3:** Redox reaction depends upon standard reduction potential of that metals.

 $E^o_{Ag^+/Ag}-E^o_{Cu^{2+}/Cu}>0\,$  and to fease the reaction,  $E^o$  should be positive of that whole reaction.

$$\therefore$$
 Cu + 2Ag<sup>+</sup>  $\rightarrow$  2Ag + Cu<sup>2+</sup>

Oxidation half:- Cu  $\rightarrow$  Cu<sup>2+</sup> + 2e<sup>-</sup>; E<sub>1</sub> = - E<sup>o</sup><sub>Cu<sup>2+</sup>/Cu</sub> Reduction half:- Ag<sup>+</sup> + e<sup>-</sup>  $\rightarrow$  Ag ;E<sub>2</sub> = E<sup>o</sup><sub>Aq<sup>+</sup>/Aq</sub>

$$E^o_{Reaction} = 2E^o_{Ag^+/Ag} - E^o_{Cu^{2+}/Cu} > 0$$



E.g. FeS 
$$\xrightarrow{O_2} FeO \xrightarrow{C+CO} Fe \xrightarrow{Fe} (inpure) \xrightarrow{Fe} (pure)$$

**Sol 5:** (a) PbS + 
$$\frac{3}{2}O_2 \longrightarrow$$
 PbO + SO<sub>2</sub>

$$\rightarrow Pb_3O_4 + SO_2$$

- (b)  $Cu_2O + C \rightarrow 2Cu + CO$
- (c)  $Ag^+_{(aq.)} + e^- \longrightarrow Ag_{(s)}$

**Sol 6:**  $E_{(OP)}^{o}$  of Zn is more positive than  $E_{OP}^{o}$  of Cu and this Zn is more powerful reductant than Cu and thus, Zn can easily replace Ag from  $[Ag(CN)_2^-]$ 

$$Zn + [Ag(CN)_{2}^{-}] \longrightarrow Ag + [Zn(CN)_{4}]^{2-} E_{1}, K_{1}$$

$$Cu + [Ag(CN)_{2}]^{-} \longrightarrow Ag + [Cu(CN)_{4}]^{2-} E_{2}, K_{2}$$

$$K_{1} >> K_{2} \text{ because } |E_{1}| > |E_{2}|$$

Furthermore, Zn being cheaper in comparison to Cu.

**Sol 7:** Instead of molten NaCl, if we use aqueous solution of NaCl, then, at cathode, Na<sup>+</sup> ions are not discharged at cathode on the contrary, H<sup>+</sup> ions get discharged to form  $H_2$ .

$$\mathsf{H}^{+} + \mathsf{e}^{-} \rightarrow \frac{1}{2}\mathsf{H}_{2}^{+} \uparrow$$

At anode,  $Cl_2^{\uparrow}$  will form irrespective of solvent.

In molten NaCl, there is no competition for Na $^{+}$  to get discharged at cathode.

 $Na^{+}_{(aq.)} + e^{-} \longrightarrow Na_{(s)}$ 

**Sol 8:** The members of group 1 i.e. alkali metals have low ionization potential value and thus possess high electropositive character as well as high oxidation potential which enables then to get easily oxidised so that's why they are strong reducing agent but they are not widely used because they react violently with water Na + H<sub>2</sub>O  $\rightarrow$  NaOH + H<sub>2</sub> $\uparrow$ 

**Sol 9:** In blast furnace, lime stone,  $CaCO_3$  is useful in fusion zone, where at 800°C – 1200°C,  $CaCO_3$  gets dissociated in CaO + CO<sub>2</sub> and it acts as flux, which reacts with acidic gangue SiO<sub>2</sub> to form slag. CaCO<sub>3</sub>  $\rightarrow$  CaO + CO<sub>2</sub>

 $CaO + SiO_2 \rightarrow CaSiO_3$  (slag)

**Sol 10:** In blast furnace, temperature differs widely from top to bottom.

So, in different temperature range different compound behaves as a reducing agent.

CO is better reducing agent than C

$$3Fe_2O_3 + C \rightarrow 2Fe_3O_4 + CO$$
  
 $Fe_3O_4 + CO \rightarrow 3FeO + CO_2$   
 $FeO + CO \rightarrow Fe + CO_2$   
For this in zone of combust

For this, in zone of combustion.

 $C + O_2 \rightarrow CO_2$  $CO_2 + C \rightarrow 2CO$ 

**Sol 11:** (i)  $CdCO_3 \xrightarrow{\Delta} CdO + CO_2$ Thermal decomposition (ii) MgO  $\xrightarrow{\Delta}$  No reaction (iii) SnO<sub>2</sub> + CO  $\xrightarrow{\Delta}_{\text{Reduction}}$  Sn + 2CO<sub>2</sub> (iv) CdSO<sub>4</sub>(aq)  $\xrightarrow{\text{electrolysis}} \underbrace{H_2SO_4 + O_2}_{\text{at anode}} + \underbrace{Cd}_{\text{at cathode}}$  (v) 2HgO  $\xrightarrow{\Lambda}$  Hg + O<sub>2</sub>

Thermal decomposition

(vi) MgO + Zn  $\xrightarrow{\Delta}$  NO reaction

**Sol 12:** Graphite and diamond, both are isomers but their physical properties differ widely.



Graphite conducts electricity because there exist free electrons between two sheets of graphite. On the other hand, diamond is a bad conductor because it does not have free electrons and during electrolysis, it is important for anode to conduct electricity and complete the circuit.



Oxidising Reducing agent agent

Sol 14: Calcination:-

$$\begin{split} \mathsf{MgCO}_3.\mathsf{CaCO}_3 & \xrightarrow{\mathsf{Calcination}} \mathsf{MgO} + \mathsf{CaO} + 2\mathsf{CO}_2^{\uparrow} \\ \mathsf{Dolomite} \\ \mathsf{MgCl}_2 + \mathsf{CaCO}_3 & \downarrow & \xrightarrow{\mathsf{CO}_2} \mathsf{MgCl}_2 + \mathsf{CaCl}_2 \\ \mathsf{Now}, \text{ electrolysis of } \mathsf{MgCl}_2 \text{ gives } \mathsf{Mg} \\ \mathsf{Mg}^{2+} + 2e^- & \rightarrow \mathsf{Mg} \text{ Anode reaction} \\ 2\mathsf{Cl}^- & \rightarrow \mathsf{Cl}_2^{\uparrow} + 2e^- \mathsf{Cathode reaction} \end{split}$$

Sol 15: Calicnation

Sol 16: Mg 
$$\xrightarrow{HNO_3}$$
 Mg(NO<sub>3</sub>)<sub>2</sub>  $\xrightarrow{\Delta}$  MgO  
 $\downarrow$  Cl<sub>3</sub>  
MgCl<sub>2</sub>  $\xrightarrow{KHO}$  Mg(OH)<sub>2</sub>  $\xrightarrow{}$  H<sub>2</sub>O



(A)  $\rightarrow$  (D) leaching process

 $\therefore$  these are the compounds of Ag

 $\therefore$  (A) Ag<sub>2</sub>S

(B) [Ag(CN)<sub>2</sub>]<sup>-</sup> Na<sup>+</sup>

(C)  $Na_2S$ 

(D) Ag

- (E) AgCl
- (F)  $[Ag(NH_3)_2]CI$

Sol 18: (i) Aluminium:- Gold Schimdt process:-

Al + Mn<sub>3</sub>O<sub>4</sub>  $\rightarrow$  Mn + Al<sub>2</sub>O<sub>3</sub> (ii) Fe is also used

 $Fe + HgS \rightarrow Hg + FeS$ 

**Sol 19:** Aluminium metal is frequently used as a reducing agent for the extraction of metal such as Cr and Mn from their respective oxides because aluminium is more electropositive than Cr/Mn. The process of reduction is known as Aluminothermite process (Gold shmidt process)

 $Cr_2O_3 + 2AI \rightarrow Al_2O_3 + 2Cr; \Delta H = -ve$  $3Mn_2O_4 + 8AI \rightarrow 4Al_2O_2 + 9Mn; \Delta H = -ve$  **Sol 20:** During the extraction of iron (Fe) from haematite  $(Fe_2O_3)$  it contains some impurities such as  $P_4O_{10}$  or  $SO_2$ , which are acidic in nature. So, to remove this gangue as slag basic lining of lime (CaO) or magnesia (MgO) is used

 $6CaO + P_4O_{10} \rightarrow 2Ca_3(PO_4)_2$ MgO + SO\_2  $\rightarrow$  MgSO\_3

**Sol 21:** Lead is extracted from its mineral galena (PbS). (i) It is firstly concentrated by froth floatation process.

(ii) Roasting in reverbatory furnace

 $2PbS + 3O_2 \rightarrow 2PbO + 2SO_2 \downarrow$ 

 $PbS + 2O_2 \rightarrow PbSO_4$ 

The idea of Cu ores come from calcined black solid and ppt. after reaction with KI.

**Sol 22:** A<sub>1</sub> will be malachite i.e., CuCO<sub>3</sub>.Cu(OH)<sub>2</sub>

 $A_2$  will be copper glance i.e.,  $Cu_2S$  The confirmatory reaction are

(a) For malachite

(i) 
$$CuCO_3.Cu(OH)_2 \xrightarrow{Calcinations} 2CuO + CO_2 \uparrow H_2O$$
  
Black solid

(ii) 
$$CuCO_3.Cu(OH)_2 \xrightarrow{HCI} CuCI_2 + CO_2^+ + H_2O$$

$$\mathsf{CuCl}_{2} \xrightarrow{\mathsf{KI}} \mathsf{Cu}_{2} \operatorname{I}_{2} \downarrow + \mathsf{KCl} + \operatorname{I}_{2}$$

(b) For copper glance

$$Cu_2S \xrightarrow{Roasting} Cu_2O + SO_2$$

 $Cu_2S + Cu_2O \rightarrow Cu + SO_2$ 

 $SO_2$  is the gas which gives green colour with acidified  $K_2Cr_2O_7$  as

$$3SO_2 + K_2Cr_2O_7 + H_2SO_4 \rightarrow K_2SO_4 + Cr_2(SO_4)_3 + 4H_2O$$

Now reduction of AgCl,

(i) 
$$2\text{AgCl} + \text{Na}_2\text{CO}_3 \rightarrow \text{Ag}_2\text{CO}_3 + 2\text{NaCl}$$

$$Ag_2CO_3 \xrightarrow{\Lambda} 2Ag \downarrow + CO_2\uparrow + \frac{1}{2}O_2\uparrow$$

(ii) Cyanide process with AgCl.

**Sol 23:** Cyanide process is a type of leaching process which a chemical reaction is used to extract metal from ore. In this process, crushed rock containing traces of gold is treated with 0.1 - 0.2% solution of NaCN and aerated  $O_2$  (air) oxidized free metal to Au<sup>+</sup> which complexes with CN<sup>-</sup>

 $4Au + 8CN^{-} + 2H_2O + O_2 \rightarrow 4[Au(CN)_2]^{-} + 4OH^{-}$ 

The pure metal is displaced from solution by an active metal

 $2[Au(CN)_2]^+ Zn \rightarrow [Zn(CN)_4]^2 + 2Au \downarrow$ 

**Sol 24:** Silver coin contains Ag, Au and Cu also. So, to extract silver, it should undergo various processes.

So, individually every metal impurities can be removed.

Silver coin

$$(Cu+Ag+Au) \xrightarrow{Conc.} (Cu^{2+} + Ag^{+} + Au^{3+})NO_{3}^{-}$$

$$\downarrow NaCl$$

$$AgCl \downarrow (white ppt.)$$

$$AuCl_{3} \downarrow$$

Now, since Ag, Au, both are noble metals, they possess similar chemical properties.

But due to difference in physical properties, Au can be removed by fractional distillation.

After oxidising minerals,

The roasted mineral is smelted into lead

$$2PbO + PbS \rightarrow 3Pb + SO_2 \uparrow$$
  
PbSO<sub>4</sub> + PbS → 2Pb + 2SO<sub>2</sub> ↑ Self reduction

PbO (litharge) oxidation state  $\rightarrow +2$ 

Purification is done by cupellation, polling and most importantly, electrolysis.

**Sol 25:** In the purification of bauxite, due to impurities such as silica  $(SiO_2)$  and other silicates, Bauxite is converted into alumina by Bayer's process. During this process,  $CO_2$  is passed through it. Aqueous solution of  $CO_2$  is acidic. Hence,  $CO_2$  when passed into  $[Al(OH)_4]^-$  solution makes it acidic. Hence,  $Al(OH)_3$  is precipitated.

 $CO_{2} + H_{2}O \rightarrow H_{3}O^{+} + HCO_{3}^{-}$  $[AI(OH)_{4}]^{-} + H_{3}O^{+} \rightarrow AI(OH)_{3}\downarrow + 2H_{2}O$ 

**Sol 26:** Haematite ore majorily contain  $[Fe_2O_3/FeO]$  but also contains silica  $(SiO_2)$ . So, during roasting in blast furnance, carbon reduction is done along with lime. So, silica can be removed as slag.

$$CaO + SiO_2 \rightarrow CaSiO_3$$

Simultaneously, Fe also gets reduced

$$C + O_2 \rightarrow CO_2; CO_2 + C \rightarrow 2CC$$
$$3CO + Fe_2O_3 \rightarrow 2Fe_{(Steel)} + 3CO_2$$

So difference in densities make them easily separable at the bottom of furnace and this calcium silicate because of its hard nature is widely used in cement industry.

**Sol 27:** (ii) Au and Ag both are generally chemically inert but they differ during their reaction with HCl +  $HNO_3$ 

Au + 4HCl + 3HNO<sub>3</sub> 
$$\rightarrow$$
 HAuCl<sub>4</sub> + 3NO<sub>2</sub> + 3H<sub>2</sub>O  
Ag + HCl + HNO<sub>3</sub>  $\rightarrow$  AgCl + NO<sub>2</sub> + H<sub>2</sub>O

So, in Aquaregia, Au forms a soluble complex compound,  $HAuCI_4$  but Ag forms a precipitate which do not dissolve.

# Exercise 2

#### Single Correct Choice Type

**Sol 1: (A)** Sulphides have fundamental property based on the different wetting property of the ore and gangue with oil and water.

## Sol 2: (C) Van Arkel

This is vapour phase refining process by forming volatile complex compound.

**Sol 3: (B)** Cryolite-Na<sub>3</sub>AlF<sub>6</sub> is used during the electrolysis of alumina to increase the electrical conductivity and decrease the melting point of alumina (Al<sub>2</sub>O<sub>3</sub>).

Sol 4: (D) Cassiterite is an ore of Sn.

**Sol 5: (C)** Discharging of element at cathode and anode depends upon various factors mainly on

(i) standard reduction potential

(ii) Nature of electrode

Using platinum and graphite electrode, we get different products so it is entirely experimental.

**Sol 6: (D)** Metal makes lattices in which nucleus is arranged in specific array of position but valence electrons are free to move into whole lattice this property gives lustre to a metal.

**Sol 7: (C)** For Cr, instead of electrolysis, we can use more cheaper process, aluminothermite, because electrolysis gives high yield but it's a very costly process.

**Sol 8: (D)** Flux's basic use is to combine with gangues/ matrix or other impurities and then remove them by slag formation.

E.g. Formation of CaSiO<sub>3</sub> in extraction of iron (Fe).

**Sol 9: (B)** In extraction of Cu, FeO is a gangue to remove it,  $SiO_2$  (silica) is added and it forms

Thomas slag

$$\underset{\text{Gangue}}{\text{FeO}} + \underset{\text{Flux}}{\text{SiO}_2} \longrightarrow \underset{\text{Thomas slag}}{\text{FeSiO}_3}$$

Sol 10 (D) In blast furnace

$$\begin{array}{c} \mathsf{FeO}/\mathsf{Fe}_2\mathsf{O}_3 \twoheadrightarrow \mathsf{Fe}_3\mathsf{O}_4 \longrightarrow \mathsf{Fe} \\ & & & \\ & & & \\ & & & \\ \mathsf{Roasting} \\ & & & \\ & & & \\ & & & \\ \mathsf{Fe}_3\mathsf{O}_4 \end{array}$$

**Sol 11: (C)**  $AI_2O_3$  is used in molten state because aqueous solution will give  $H_2$  at cathode and  $Na_3AIF_6$  is used to increase the conductivity of  $AI_2O_3$  and to decrease the melting point of mixture.

**Sol 12: (C)** Calcination is used in metallurgy to remove CO<sub>2</sub> and H<sub>2</sub>S

**Sol 13: (C)** The metallic luster exhibited by sodium is explained by the presence of Free protons

Sol 14: (B) Stainless steel:- Fe + Cr + Ni

Sol 15: (B) Leaching:-

 $\begin{aligned} AI_2O_3 + 2NaOH &\rightarrow 2NaAIO_2 + H_2O \\ 2NaAIO_2 + 4H_2O &\rightarrow 2AI(OH)_3 + 2NaOH \\ 2AI(OH)_3 &\xrightarrow{A}_{1200} & AI_2O_3 + 3H_2O \end{aligned}$ 

Sol 16: (D) Bessemerisation of Cu



Bessemer converter

 $2FeS + 3O_2 \rightarrow 2FeO + 2SO_2$   $2CuS + SiO_2 \rightarrow FeSiO_3 \text{ (slag)}$   $2CuS + 3O_2 \rightarrow 2Cu_2O + 2SO_2$  $2Cu_2O + Cu_2S \rightarrow 6Cu + SO_2$ 

**Sol 17: (D)** Generally they are more electronegative than metals.

**Sol 18: (B)** In the froth floatation process for the purification of ores, the ore particles float because their surface is hydrophobic i.e. not easily wetted by water

Sol 19: (C) SiO<sub>2</sub> Acidic flux.

**Sol 20: (B)** Flux + Gangue  $\rightarrow$  slag

Sol 21: (B) E.g. Na, K, Ca, Al, etc.

Group I, II metals.

**Sol 22: (D)** The role of calcination in metallurgical operations is to remove moisture, to decompose carbonate,to drive off organic matter.

#### **Comprehension type**

Sol 23: (A) CuS

**Sol 24: (A)** H<sub>2</sub>S

**Sol 25: (B)** CuSO<sub>4</sub>

**Sol 26: (C)** [Cu(NH<sub>3</sub>)<sub>4</sub>](NO<sub>3</sub>)<sub>2</sub>

#### Sol 27: (A)

$$(A) = Black \xrightarrow{Dil. H_2SO_4} (B) + (C)$$

$$(XS) NH_4OH$$

$$(D) = Chocolate brown ppt.$$

$$(E) = Chocolate brown ppt. form with Cu metal$$

$$(E) is Cu_2 [Fe(CN)_6]$$

#### **Assertion Reasoning Type**

**Sol 28: (D)** Calamine  $(ZnCO_3)$ (R) Boric acid is monobasic acid  $B(OH)_3 + H_2O \rightarrow [B(OH)_4]^- + H^+$ 

Sol 29: (D) Based on experiments.

**Sol 30: (C)**  $Pb^{4+}$  is easily reduced to  $Pb^{2+}$  because of inert pair effect.

(R) Pb<sup>2+</sup> is diamagnetic.

**Sol 31: (C)** Diamond is hardest natural occurring substance because of its strong 4 valence lattice.

(R) Graphite is more stable because of electrical conductivity and availability of free electrons.

Sol 32: (C) Mond's Process:-

 $\begin{array}{ccc} \mathsf{Ni} + 4\mathsf{CO} & \xrightarrow{\Delta} & \mathsf{Ni}(\mathsf{CO})_{4} & \xrightarrow{\Delta} & \mathsf{Ni} \downarrow + 4\mathsf{CO} \\ & & & & \\ & & &$ 

## **Previous Years' Questions**

Sol 1: (C) Chalcopyrite contain both iron and copper.

**Sol 2: (B)** Zinc blende contain ZnS which is first roasted partially and then subjected to reduction with carbon

 $ZnS + O_2 \rightarrow ZnO + SO_2$  Roasting

 $ZnO + C \xrightarrow{\Delta} Zn + CO^{\uparrow}$  Carbon reduction

**Sol 3: (B)** A water soluble complex with silver and dilute aqueous solution of NaCN is Na[Ag(CN)<sub>2</sub>]. In the cyanide process, the native silver is crushed and treated with aqueous NaCN solution and aerated.

$$4Ag + 8NaCN + 2H_2O + O_2 \rightarrow 4Na[Ag(CN)_2] + 4NaOH$$

**Sol 4: (D)** Haematite is  $Fe_2O_3$ , in which oxidation number of iron is III. Magnetite is  $Fe_3O_4$  which is infact a mixed oxide (FeO.Fe\_2O\_3), hence, iron is present in both II and III oxidation state.

**Sol 5: (A, C)** Addition of manganese to iron improve hardness of steel as well as remove oxygen and sulphur.

**Sol 6: (A, D)** The important ore of tin is cassiterite  $(SnO_2)$ . Tin is extracted from cassiterite ore by carbon reduction method in a blast furnace.

 $SnO_2 + 2C \rightarrow Sn + 2CO$ 

The product often contain traces of iron which is remov-ed by blowing air through the melt to oxidize to FeO which then floats to the surface.

 $2\text{Fe} + \text{O}_2 \rightarrow 2\text{FeO}$ 

**Sol 7: (B)** 
$$2CuFeS_2 + O_2 \rightarrow Cu_2S + 2FeS + SO_2^{\uparrow}$$
  
 $2Cu_2S + 3O_2 \rightarrow 2Cu_2O + 2SO_2^{\uparrow}$   
 $2FeS + 3O_2 \rightarrow 2FeO + 2SO_2^{\uparrow}$ 

**Sol 8: (D)** FeO + SiO<sub>2</sub>  $\rightarrow$  FeSiO<sub>3</sub> (slag)

**Sol 9: (C)**  $Cu_2S + 2Cu_2O \rightarrow 6Cu + SO_2$ 

In Cu<sub>2</sub>S, sulphur is S<sup>2-</sup> and in SO<sub>2</sub>, sulphur is in +4 state. Hence, S<sup>2-</sup> is acting as a reducing agent.

Sol 10: (A)  $A \rightarrow s$ ;  $B \rightarrow p$ ;  $C \rightarrow q$ ;  $D \rightarrow q$ , r;  $E \rightarrow t$ 

Silver is extracted by amalgamation process

 $Ag + Hg \rightarrow Ag(Hg) \xrightarrow{\text{Distillation}} Ag(s) + Hg(g)^{\uparrow}$ 

(B) Calcium is extracted by electrolysis of fused CaCl<sub>2</sub>.

(C) Zinc is extracted by carbon reduction method

 $ZnO + C \rightarrow Zn + CO$ 

(D) Iron is extracted by both, carbon reduction method and CO reduction methods

 $Fe_2O_3 + 3C \rightarrow 2Fe + 3CO$  $Fe_2O_3 + 3CO \rightarrow 2Fe + 3CO_2$  (E) Copper is extracted by self-reduction methods

$$Cu_2S + O_2 \rightarrow Cu_2O + SO_2$$
$$Cu_2O + Cu_2S \rightarrow Cu + SO_2$$

**Sol 11:**  $4NaCN + Ag_2S \rightarrow 2Na[Ag(CN)_2] + Na_2S$  $2Na[Ag(CN)_2] + Zn \rightarrow Na_2[Zn(CN)_4] + 2Ag$ 

**Sol 12:** The common photographic film is coated with AgBr and during developing of photographic film, the unreacted AgBr is removed by Na<sub>2</sub>S<sub>2</sub>O<sub>2</sub> as

$$AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaBr$$

Sol 13: 
$$2AgBr + C_6H_4(OH)_2(Hydroquinone) \rightarrow$$
  
 $2Ag + 2HBr + C_6H_4O_2$   
(developer) quinone  
 $AgBr + 2Na_2S_2O_3 \rightarrow Na_3[Ag(S_2O_3)_2] + NaBr$   
(b)  $Na_2S_2O_3 + 2H^+ \rightarrow 2Na^+ + H_2SO_3 + S\downarrow$   
Colloidal  
Sulphur

**Sol 14:** Four, the complex has formula  $[Cu(H_2O)_4] SO_4.H_2O$ 

**Sol 15:** In crystalline state,  $AICI_3$  has rock-salt like structure with coordination number of AI = 6.

**Sol 16: (B)** Extraction of zinc from zinc blende is achieved by

**Sol 17:**  $A \rightarrow p$ ;  $B \rightarrow q$ ;  $C \rightarrow p$ , r;  $D \rightarrow p$ , s PbS  $\rightarrow$  PbO - Roasting CaCO<sub>3</sub>  $\rightarrow$  CaO - Calcination ZnS  $\rightarrow$  Zn - Roasting, Carbon reduction Cu<sub>2</sub>S  $\rightarrow$  Cu - Roasting, Self reduction

#### Sol 18: (A)

Sulfinde ore of Ag  $\rightarrow$  Argentite (Ag<sub>2</sub>S) Pb  $\rightarrow$  Galena (PbS) Cu  $\rightarrow$  Copper glance (Cu<sub>2</sub>S) Hence (A) is correct.

**Sol 19 : (C, D)**  $Fe_2O_3$  and  $SnO_2$  undergoes C reduction. Hence (C) and (D) are correct. **Sol 21: (A, B, C)** Extraction of copper from copper pyrite  $(CuFeS_2)$  involve crushing following by concentration of the ore by froth-flotation, removal of iron as slag, self-reduction step to produce 'blister copper' following evolution of SO<sub>2</sub>