### The D and F Block Elements

### Question1

Given below are two statements:

Statement (I): The 4f and 5f - series of elements are placed separately in the Periodic table to preserve the principle of classification. Statement (II): S-block elements can be found in pure form in nature. In the light of the above statements, choose the most appropriate answer from the options given below:

[27-Jan-2024 Shift 1]

### **Options:**

- A. Statement I is false but Statement II is true
- B. Both Statement I and Statement II are true
- C. Statement I is true but Statement II is false
- D. Both Statement I and Statement II are false

**Answer: C** 

#### **Solution:**

Solution:

s-block elements are highly reactive and found in combined state.

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### **Question2**

NaCl reacts with conc.  $H_2SO_4$  and  $K_2Cr_2O_7$  to give reddish fumes (B), which react with NaOH to give yellow solution (C). (B) and (C) respectively are;

[27-Jan-2024 Shift 1]

#### **Options:**

- A.  $CrO_2Cl_2$ ,  $Na_2CrO_4$
- B.  $Na_2CrO_4$ ,  $CrO_2Cl_2$
- C.  $\mathrm{CrO_2Cl_2}$ ,  $\mathrm{KHSO_4}$
- D.  $CrO_2Cl_2$ ,  $Na_2Cr_2O_7$

**Answer: A** 

$$\begin{aligned} \text{NaCl+} & \text{conc.} & \text{H}_2\text{SO}_4 + \text{K}_2\text{Cr}_2\text{O}_7 &\longrightarrow \text{CrO}_2\text{Cl}_2 &+ \text{KHSO}_4 + \text{NaHSO}_4 + \text{H}_2\text{O} \\ \text{CrO}_2\text{Cl}_2 + \text{NaOH} &\longrightarrow \text{Na}_2\text{CrO}_4 &+ \text{NaCl} + \text{H}_2\text{O} \\ & \text{(C) Yellow colour} \end{aligned}$$

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### **Question3**

Given below are two statements:

Statement (I): In the Lanthanoids, the formation of  $\mathrm{Ce}^{+4}$  is favoured by its noble gas configuration.

Statement (II) :  $Ce^{+4}$  is a strong oxidant reverting to the common +3 state.

In the light of the above statements, choose the most appropriate answer from the options given below:

[27-Jan-2024 Shift 2]

### **Options:**

- A. Statement I is false but Statement II is true
- B. Both Statement I and Statement II are true
- C. Statement I is true but Statement II is false
- D. Both Statement I and Statement II are false

**Answer: B** 

#### **Solution:**

#### Solution:

Statement (1) is true. Ce<sup>+4</sup> has noble gas electronic configuration.

Statement (2) is also true due to high reduction potential for  $\mathrm{Ce}^{4+}$  /  $\mathrm{Ce}^{3+}$  (+1.74V), and stability of  $\mathrm{Ce}^{3+}$ ,  $\mathrm{Ce}^{4+}$  acts as strong oxidizing agent.

### **Question4**

Choose the correct option having all the elements with d<sup>10</sup> electronic configuration from the following: [27-Jan-2024 Shift 2]

### **Options:**

A. 
$$^{27}$$
 Co,  $^{28}$  Ni,  $^{26}$  Fe,  $^{24}$  Cr

D. 
$$^{28}$$
 Ni,  $^{24}$  Cr,  $^{26}$  Fe,  $^{29}$  Cu



**Answer: B** 

### **Solution:**

$$[Cr] = [Ar]4 s^1 3d^5$$

$$[Cd] = [Kr]5 s^2 4d^{10}$$

$$[Cu] = [Ar]4 s^1 3d^{10}$$

$$[Ag] = [Kr]5 s^1 4d^{10}$$

$$[Zn] = [Ar]4 s^2 3d^{10}$$

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## **Question5**

In alkaline medium.  $MnO_4$  oxidises I to [29-Jan-2024 Shift 1]

**Options:** 

A.  $IO_4^-$ 

B. IO

C. I<sub>2</sub>

D. IO<sub>3</sub>

**Answer: D** 

### **Solution:**

**Solution:** 

$$2MnO_4^- + H_2O + I^-$$
 alkaline medium  $= 2MnO_2 + 2OH^- + IO_3^-$ 

**Question6** 

In chromyl chloride test for confirmation of  $Cl^-$  ion, a yellow solution is obtained. Acidification of the solution and addition of amyl alcohol and  $10\%~H_2O_2$  turns organic layer blue indicating formation of chromium pentoxide. The oxidation state of chromium in that is [29-Jan-2024 Shift 1]

**Options:** 

A. +6

B. +5

C. +10

D. +3

**Answer: A** 

### **Solution:**

$$\begin{array}{c} \text{C1}^- + \text{K}_2 \text{Cr}_2 \text{O}_7 + \text{H}_2 \text{SO}_4 \longrightarrow \text{CrO}_2 \text{C1}_2 \xrightarrow{\text{Basic medium}} & \text{CrO}_4^{2^-} + \text{C1}^-\\ \text{yellow solution} & \\ \begin{array}{c} \text{CrO}_4^{2^-} & \xrightarrow{\text{1.Acidification}} \\ \text{2. Amyl alcohol} & \\ \text{3.10\% H}_2 \text{O}_2 & \\ \text{blue compound} \end{array}$$

### **Question7**

Which of the following statements are correct about Zn, Cd and Hg? A. They exhibit high enthalpy of atomization as the d-subshell is full. B. Zn and Cd do not show variable oxidation state while Hg shows +I and + II.

C. Compounds of Zn, Cd and Hg are paramagnetic in nature.

D. Zn, Cd and Hg are called soft metals.

Choose the most appropriate from the options given below: [29-Jan-2024 Shift 2]

### **Options:**

A. B, D only

B. B, C only

C. A, D only

D. C, D only

**Answer: A** 

#### **Solution:**

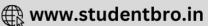
#### **Solution:**

(A) Zn, Cd, Hg exhibit lowest enthalpy of atomization in respective transition series.

(C) Compounds of Zn, Cd and Hg are diamagnetic in nature.

### **Question8**

Which of the following acts as a strong reducing agent? (Atomic number : Ce = 58, Eu = 63, Gd = 64, Lu = 71)



### [29-Jan-2024 Shift 2]

### **Options:**

A. Lu<sup>3+</sup>

B. Gd<sup>3+</sup>

C. Eu<sup>2+</sup>

D. Ce<sup>4+</sup>

**Answer: C** 

### **Solution:**

#### Solution:

Eu<sup>+2</sup> 
$$\blacktriangleright$$
 Eu<sup>+3</sup> + 1e<sup>-</sup>
[Xe]4f<sup>7</sup>6 s<sup>0</sup> [Xe]4f<sup>6</sup>6 s<sup>0</sup>

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### **Question9**

### **Match List-II with List-II**

List-I	List-II	
Species	Electronic distribution	
(A) Cr <sup>-2</sup>	(I) 3d <sup>8</sup>	
(B) Mn <sup>+</sup>	(II) 3d <sup>3</sup> 4 s <sup>1</sup>	
(C) Ni <sup>-2</sup>	(III) 3d <sup>4</sup>	
(D) V	(IV) 3d <sup>5</sup> 4 s <sup>1</sup>	

# Choose the correct answer from the options given below: [30-Jan-2024 Shift 1]

### **Options:**

A. (A)-I, (B)-II, (C)-III, (D)-IV

B. (A)-III, (B) - IV, (C) - I, (D)-II

C. (A)-IV, (B)-III, (C)-I, (D)-II

D. (A)-II, (B)-I, (C)-IV, (D)-III

**Answer: B** 



$$_{24}$$
Cr → [Ar]3d<sup>5</sup>4 s<sup>1</sup>; Cr<sup>2+</sup> → [Ar]3d<sup>4</sup>  
 $_{25}$ Mn → [Ar]3d<sup>5</sup>4 s<sup>2</sup>; Mn<sup>+</sup> → [Ar]3d<sup>5</sup>4 s<sup>1</sup>  
 $_{28}$ Ni → [Ar]3d<sup>8</sup>4 s<sup>2</sup>; Ni<sup>2+</sup> → [Ar]3d<sup>8</sup>  
 $_{33}$ V → [Ar]3d<sup>3</sup>4 s<sup>2</sup>; V<sup>+</sup> → [Ar]3d<sup>3</sup>4 s<sup>1</sup>

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### Question 10

# Diamagnetic Lanthanoid ions are: [30-Jan-2024 Shift 1]

### **Options:**

A.  $Nd^{3+}$  and  $Eu^{3+}$ 

B. La<sup>3+</sup> and Ce<sup>4+</sup>

C. Nd<sup>3+</sup> and Ce<sup>4+</sup>

D.  $Lu^{3+}$  and  $Eu^{3+}$ 

Answer: B

### **Solution:**

 $Ce : [Xe]4f^{1}5d^{1}6 s^{2}; Ce^{4+} diamagnetic$ 

 $La: [Xe]4f^05d^16 s^2; La^{3+}$  diamagnetic

\_\_\_\_\_

### Question11

# The orange colour of $\rm K_2Cr_2O_7$ and purple colour of $\rm KMnO_4$ is due to [30-Jan-2024 Shift 2]

### **Options:**

A. Charge transfer transition in both.

B. d  $\rightarrow$  d transition in KMnO<sub>4</sub> and charge transfer transitions in K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>.

C. d  $\rightarrow$  d transition in  $\mathrm{K_2Cr_2O_7}$  and charge transfer transitions in  $\mathrm{KMnO_4}$ .

D.  $d \rightarrow d$  transition in both.

**Answer: A** 



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### **Question12**

Alkaline oxidative fusion of  $MnO_2$  gives " A " which on electrolytic oxidation in alkaline solution produces B. A and B respectively are: [30-Jan-2024 Shift 2]

**Options:** 

A. Mn<sub>2</sub>O<sub>7</sub> and MnO<sub>4</sub>

B.  $MnO_4^{2-}$  and  $MnO_4^{-}$ 

C.  $Mn_2O_3$  and  $MnO_4^{\ 2-}$ 

D.  $\mathrm{MnO_4}^{2-}$  and  $\mathrm{Mn_2O_7}$ 

**Answer: B** 

**Solution:** 

Alkaline oxidative fusion of MnO,:

 $2MnO_2 + 4OH^- + O_2 \rightarrow 2MnO_4^{2-} + 2H_2O$ 

Electrolytic oxidation of  $MnO_4^{\ 2-}$  in alkaline medium.

 $MnO_4^{2-} \longrightarrow MnO_4^{-} + e^{-}$ 

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## Question 13

A and B formed in the following reactions are:

 $CrO_2Cl_2 + 4 NaOH \rightarrow A + 2 NaCl + 2H_2O$ 

 $A + 2HCl + 2H_2O_2 \rightarrow B + 3H_2O$ 

[30-Jan-2024 Shift 2]

**Options:** 

$$A. A = Na_2CrO_4, B = CrO_5$$

$$B. A = Na_2Cr_2O_4, B = CrO_4$$

$$C. A = Na_2Cr_2O_7$$
,  $B = CrO_3$ 

D. 
$$A = Na_2Cr_2O_7$$
,  $B = CrO_5$ 

**Answer: A** 

#### **Solution:**

$$\begin{aligned} &\operatorname{CrO_2Cl_2} + 4\operatorname{NaOH} \longrightarrow \operatorname{Na_2CrO_4} + 2\operatorname{NaC1} + 2\operatorname{H_2O} \\ &\overset{\text{(A)}}{\underset{\text{(B)}}{\operatorname{CrO_4}}} + 2\operatorname{H_2O_2} + 2\operatorname{HC1} \longrightarrow & \operatorname{CrO_5} + \\ && \underset{\text{Missing from balanced eqaution}}{\operatorname{CrO_5}} + 3\operatorname{H_2O} \end{aligned}$$

### Question14

**Identify correct statements from below:** 

- A. The chromate ion is square planar.
- B. Dichromates are generally prepared from chromates.
- C. The green manganate ion is diamagnetic.
- D. Dark green coloured  $K_2MnO_4$  disproportionate in a neutral or acidic medium to give permanganate.
- E. With increasing oxidation number of transition metal, ionic character of the oxides decreases.

Choose the correct answer from the options given below:

[31-Jan-2024 Shift 1]

### **Options:**

A. B, C, D only

B. A, D, E only

C. A, B, C only

D. B, D, E only

**Answer: D** 

#### **Solution:**

B. 
$$2Na_2CrO_4 + 2H^+ \rightarrow Na_2Cr_2O_7 + 2Na^+ + H_2O$$

- C. As per NCERT, green manganate is paramagnetic with 1 unpaired electron.
- D. Statement is correct
- E. Statement is correct

### **Question15**

The 'Spin only' Magnetic moment for  $[Ni(NH_3)_6]^{2+}$  is  $\_\_\times 10^{-1}$  BM.

( given = Atomic number of Ni : 28)

[31-Jan-2024 Shift 1]



**Answer: 28** 

### **Solution:**

NH3 act as WFL with Ni2+

$$Ni^{2+} = 3d^8$$

No. of unpaired electron = 2

$$\mu = \sqrt{n(n+2)} = \sqrt{8} = 2.82 \,\text{BM}$$

$$= 28.2 \times 10^{-1} \, \text{BM}$$

$$x = 28$$

### **Question16**

Choose the correct statements from the following

- A. Mn<sub>2</sub>O<sub>7</sub> is an oil at room temperature
- B.  $V_2O_4$  reacts with acid to give  $VO_2^{\ 2+}$
- C. CrO is a basic oxide
- D. V<sub>2</sub>O<sub>5</sub> does not react with acid

Choose the correct answer from the options given below:

[31-Jan-2024 Shift 2]

### **Options:**

- A. A, B and D only
- B. A and C only
- C. A, B and C only
- D. B and C only

**Answer: B** 

### **Solution:**

- (A) Mn,O, is green oil at room temperature.
- (B) V<sub>2</sub>O<sub>4</sub> dissolve in acids to give VO<sup>2+</sup> salts.
- (C) CrO is basic oxide
- (D) V<sub>2</sub>O<sub>5</sub> is amphoteric it reacts with acid as well as base.





Number of moles of  $H^+$ ions required by 1 mole of  $MnO_4^-$ to oxidise oxalate ion to  $CO_2$  is\_\_\_\_\_ [31-Jan-2024 Shift 2]

**Answer: 8** 

### **Solution:**

$$2MnO_4^- + 5C_2O_4^{-2-} + 16H^+ \rightarrow 2Mn^{2+} + 10CO_2 + 8H_2O$$

∴ Number of moles of H<sup>+</sup>ions required by 1 mole of MnO<sub>4</sub><sup>-</sup>to oxidise oxalate ion to CO<sub>2</sub> is 8

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### **Question18**

In the reaction of potassium dichromate, potassium chloride and sulfuric acid (conc.), the oxidation state of the chromium in the product is (+)\_\_\_ [31-Jan-2024 Shift 2]

**Answer: 6** 

#### **Solution:**

#### **Solution:**

 $\mathrm{K_2Cr_2O_7}(s) + 4\,\mathrm{KCl}(s) + 6\mathrm{H_2SO_4} \ \, (\mathrm{conc.}) \\ \longrightarrow 2\mathrm{CrO_2Cl_2}(g) + 6\mathrm{KHSO_4} + 3\mathrm{H_2O}$ 

This reaction is called chromyl chloride test.

Here oxidation state of Cr is +6.

### **Question19**

Which of the following compounds show colour due to d-d transition? [1-Feb-2024 Shift 2]

**Options:** 

A.  $CuSO_4 \cdot 5H_2O$ 



B.  $K_2Cr_2O_7$ 

C. K<sub>2</sub>CrO<sub>4</sub>

D. KMnO<sub>4</sub>

**Answer: A** 

### **Solution:**

 $CuSO_4 \cdot 5H_2O$ 

 $Cu^{2+}: 3d^94 s^0$ 

unpaired electron present so it show colour due to d-d transition.

# Question20

# The transition metal having highest $3^{rd}$ ionisation enthalpy is : [1-Feb-2024 Shift 2]

### **Options:**

A. Cr

B. Mn

C. V

D. Fe

**Answer: B** 

#### **Solution:**

Mn :  $3d^54s^2$ 

Fe :  $3d^64s^2$ 

Cr : 3d 54s1

 $V:3d^34s^2$ 

So Mn has highest 3rd IE among all the given elements due to  ${
m d}^5$  configuration.

### Question21

Given below are two statements: one is labelled as Assertion (A) and the other is labelled as Reason (R).

Assertion (A) : In aqueous solutions  ${\rm Cr}^{2+}$  is reducing while  ${\rm Mn}^{3+}$  is oxidising in nature.



Reason (R): Extra stability to half filled electronic configuration is observed than incompletely filled electronic configuration. In the light of the above statement, choose the most appropriate answer from the options given below: [1-Feb-2024 Shift 2]

#### **Options:**

- A. Both (A) and (R) are true and (R) is the correct explanation of (A)
- B. Both (A) and (R) are true but (R) is not the correct explanation of (A)
- C. (A) is false but (R) is true
- D. (A) is true but (R) is false

**Answer: A** 

#### **Solution:**

#### Solution:

 $Cr^{2+}$  is reducing as it configuration changes from  $d^4$  to  $d^3$  due to formation of  $Cr^{3+}$ , which has half filled  $t_{2g}$  level, on other hand, the change  $Mn^{3+}$  to  $Mn^{2+}$  result half filled  $d^5$  configuration which has extra stability.

### **Question22**

The magnetic moment of a transition metal compound has been calculated to be 3.87 B.M. The metal ion is [24-Jan-2023 Shift 1]

#### **Options:**

- A. Cr<sup>2+</sup>
- B. Mn<sup>2+</sup>
- $C. V^{2+}$
- D. Ti<sup>2+</sup>

**Answer: C** 

#### **Solution:**

$$\begin{array}{l} Cr^{+2}: [Ar], \ 3d^4, \ 4\ s^0n = 4, \ \mu = \sqrt{4(4+2)} = \sqrt{24} \\ = 4.89 \ BM \\ Mn^{+2}: [Ar], \ 3d^5, \ 4\ s^0n = 5, \ \mu = \sqrt{5(5+2)} = \sqrt{35} \\ = 5.91 \ BM \\ V^{+2}: [Ar], \ 3d^3, \ 4\ s^0n = 3, \ \mu = \sqrt{3(3+2)} = \sqrt{15} \\ = 3.87 \ BM \\ Ti^{+2}: [Ar], \ 3d^2, \ 4\ s^0n = 2, \ \mu = \sqrt{2(2+2)} = \sqrt{8} \\ = 2.82 \ BM \end{array}$$

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An ammoniacal metal salt solution gives a brilliant red precipitate on addition of dimethylglyoxime. The metal ion is: [24-Jan-2023 Shift 1]

A. Cu<sup>2+</sup>

B. Co<sup>2+</sup>

C. Fe<sup>2+</sup>

D. Ni<sup>2+</sup>

**Answer: D** 

#### **Solution:**

#### **Solution:**

 $Ni^{+2} + 2DMG \xrightarrow{NH_3(aq)} [Ni(DMG)_2]$ 

### **Question24**

Which one amongst the following are good oxidizing agents?

A. Sm<sup>2+</sup>

**B.** Ce<sup>2+</sup>

C. Ce<sup>4+</sup>

**D. Tb**<sup>4+</sup>

Choose the most appropriate answer from the options given below: [24-Jan-2023 Shift 2]

### **Options:**

A. Conly

B. D only

C. A and B only

D. C and D only

**Answer: D** 

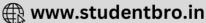
### **Solution:**

#### Solution:

Ce<sup>+4</sup> and Tb<sup>+4</sup> act as oxidising agent.

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K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> paper acidified with dilute H<sub>2</sub>SO<sub>4</sub> turns green when exposed to [24-Jan-2023 Shift 2]

**Options:** 

A. Carbon dioxide

B. Sulphur trioxide

C. Hydrogen sulphide

D. Sulphur dioxide

**Answer: D** 

**Solution:** 

$$3SO_2 + Cr_2O_7^{2-} + 2H^+ \rightarrow 3SO_4^{2-} + 2Cr^{+3} + H_2O_4^{2-}$$

### **Question26**

Potassium dichromate acts as a strong oxidizing agent in acidic solution. During this process, the oxidation state changes from [25-Jan-2023 Shift 2]

**Options:** 

A. +3 to +1

B. +6 to +3

C. +2 to +1

D. +6 to +2

**Answer: B** 

**Solution:** 

**Solution:** 

$$14H^{+} + 6e^{-} + Cr_{2}O_{7}^{-2} \longrightarrow 2Cr^{+3} + 7H_{2}O$$

### Question27

During the borax bead test with CuSO<sub>4</sub>, a blue green colour of the bead



# was observed in oxidising flame due to the formation of [29-Jan-2023 Shift 1]

### **Options:**

A.  $Cu_3B_2$ 

B. Cu

C.  $Cu(BO_2)_2$ 

D. CuO

**Answer: C** 

### **Solution:**

#### **Solution:**

Blue green colour is due to formation of Cu(BO<sub>2</sub>)<sub>2</sub>

$$\begin{aligned} &\operatorname{CuSO}_4 \overset{\Delta}{\longrightarrow} \operatorname{CuO} + \operatorname{SO}_3 \\ &\operatorname{CuO} + \operatorname{B}_2\operatorname{O}_3 \to \operatorname{Cu(BO}_2)_2 \end{aligned}$$

### **Question28**

The set of correct statements is:

- (i) Manganese exhibits +7 oxidation state in its oxide.
- (ii) Ruthenium and Osmium exhibit +8 oxidation in their oxides.
- (iii) Sc shows +4 oxidation state which is oxidizing in nature.
- (iv) Cr shows oxidising nature in +6 oxidation state.
- [29-Jan-2023 Shift 2]

### **Options:**

- A. (ii) and (iii)
- B. (i), (ii) and (iv)
- C. (i) and (iii)
- D. (ii), (iii) and (iv)

**Answer: B** 

#### **Solution:**

#### **Solution:**

(i), (ii) and (iv) correct. Manganese exhibits +7 oxidation state in its oxide. ( $\mathrm{Mn_2O_7}$ ) Ru&Os from  $\mathrm{RuO_4}$  &  $\mathrm{OsO_4}$  oxide in +8 oxidation state Cr in +6 oxidation act is oxidizing.

Sc does not show +4 oxidation state.

.....



A solution of CrO<sub>5</sub> in amyl alcohol has a....colour [29-Jan-2023 Shift 2]

**Options:** 

A. Green

B. Orange-Red

C. Yellow

D. Blue

**Answer: D** 

**Solution:** 

A solution of  ${\rm CrO_5}$  in amyl alcohol has a blue colour. So, option (4) is correct.

Question30

During the qualitative analysis of  $SO_3^{2-}$  using dilute  $H_2SO_4$ ,  $SO_2$  gas is evolved which turns  $K_2Cr_2O_7$  solution (acidified with dilute  $H_2SO_4$  ): [30-Jan-2023 Shift 1]

**Options:** 

A. Black

B. Red

C. Green

D. Blue

**Answer: C** 

**Solution:** 

$$Cr_2O_7^{2-} + SO_3^{2-} \xrightarrow{H^+} Cr^{3+} + SO_4^{2-}$$

### Question31

When Cu<sup>2+</sup> ion is treated with KI, a white precipitate, X appears in solution. The solution is titrated with sodium thiosulphate, the compound Y is formed. X and Y respectively are



### [31-Jan-2023 Shift 1]

### **Options:**

$$A. X = Cu_2I_2$$
$$Y = Na_2S_4O_5$$

B. 
$$X = Cu_2I_2 Y = Na_2S_4O_6$$

$$\mathrm{C.~X} = \mathrm{CuI}_2~\mathrm{Y} = \mathrm{Na}_2\mathrm{S}_4\mathrm{O}_3$$

D. 
$$X = CuI_2 Y = Na_2S_4O_6$$

**Answer: B** 

#### **Solution:**

#### **Solution:**

$$\begin{array}{l} \operatorname{Cu}^{2^+} + 2\operatorname{KI} \to & \operatorname{CuI}_2 & \downarrow + 2\operatorname{K}^+ \\ \Gamma^- \mathrm{is} \ \mathrm{strong} \ \mathrm{R.A} \ \mathrm{it} \ \mathrm{reduces} \ \mathrm{Cu}^{2^+} \ \mathrm{to} \ \mathrm{Cu}^+ \\ 2\operatorname{CuI}_2 \to & \operatorname{Cu}_2 \mathrm{I}_2 \downarrow + \mathrm{I}_2 \\ & \text{(White)} \text{X'} \\ \mathrm{KI} + \mathrm{I}_2 \to & \mathrm{K}^+ \mathrm{I}_3^- \ (\mathrm{Brown} \ \mathrm{solution}) \\ \mathrm{I}_3^- & \rightleftharpoons \mathrm{I}_2 + \mathrm{I}^- \\ \mathrm{KI}_3 + \operatorname{Na}_2 \mathrm{S}_2 \mathrm{O}_3 \to \mathrm{KI} + \operatorname{Na}_2 \mathrm{S}_4 \mathrm{O}_6 \\ & \text{(Y)} \end{array}$$

\_\_\_\_\_\_

### Question32

# The correct order of basicity of oxides of vanadium is [31-Jan-2023 Shift 1]

### **Options:**

A. 
$$V_2O_3 > V_2O_4 > V_2O_5$$

B. 
$$V_2O_3 > V_2O_5 > V_2O_4$$

$${\rm C.~V_2O_5} > {\rm V_2O_4} > {\rm V_2O_3}$$

$${\rm D.\ V_2O_4 > V_2O_3 > V_2O_5}$$

Answer: A

### **Solution:**

#### **Solution:**

With increase in % of oxygen acidic nature of oxide of an element increase and basic nature decreases

\_\_\_\_\_

## Question33





Highest oxidation state of Mn is exhibited in  $Mn_2O_7$ . The correct statements about  $Mn_2O_7$  are

- (A) Mn is tetrahedrally surrounded by oxygen atoms
- (B) Mn is octahedrally surrounded by oxygen atoms
- (C) Contains Mn-O-Mn bridge
- (D) Contains Mn-Mn bond.

Choose the correct answer from the options given below [1-Feb-2023 Shift 1]

#### **Options:**

- A. A and C only
- B. A and D only
- C. B and D only
- D. B and C only

**Answer: A** 

#### **Solution:**

### Question34

A solution of  ${\rm FeCl}_3$  when treated with  ${\rm K}_4[{\rm Fe(CN)}_6]$  gives a prussiun blue precipitate due to the formation of [1-Feb-2023 Shift 1]

#### **Options:**

- A. K[Fe<sub>2</sub>(CN)<sub>6</sub>]
- B. Fe[Fe(CN)<sub>6</sub>]
- C.  $Fe_3[Fe(CN)_6]_2$
- D.  $Fe_4[Fe(CN)_6]_3$

**Answer: D** 

#### **Solution:**

### Solution:

Formation of Prussian blue complex takes place.

Which of the following are the example of double salt?

- (A)  $FeSO_4 \cdot (NH_4)_2 SO_4 \cdot 6H_2O$
- (B)  $CuSO_4 \cdot 4NH_3 \cdot H_2O$
- (C)  $K_2SO_4 \cdot Al_2(SO_4)_3 \cdot 24H_2O$
- (D)  $Fe(CN)_2 \cdot 4 KCN$

Choose the correct answer.

[1-Feb-2023 Shift 1]

#### **Options:**

- A. A and C only
- B. A and B only
- C. A, B and D only
- D. B and D only

**Answer: A** 

#### **Solution:**

#### Solution:

Double salt contain's two or more types of salts.  $CuSO_4.4NH_3 \cdot H_2O$  and  $Fe(CN)_2.4$  KCN are complex compounds.

\_\_\_\_\_

### Question36

Given below are two statements: one is labelled as Assertion (A) and the other is labelled as Reason (R).

Assertion (A):  $Cu^{2+}$  in water is more stable than  $Cu^{+}$ .

Reason (R): Enthalpy of hydration for  $Cu^{2+}$  is much less than that of  $Cu^{+}$ .

In the light of the above statements, choose the correct answer from the options given below:

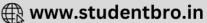
[1-Feb-2023 Shift 2]

#### **Options:**

- A. Both (A) and (R) are correct and (R) is the correct explanation of (A).
- B. (A) is correct but (R) is not correct.
- C. (1) is not correct but (R) is correct.
- D. Both (A) and (R) are correct but (R) is not the correct explanation of (A).

**Answer: A** 





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

The stability of  $Cu^{2+}$  (aq) rather than  $Cu^{+}(aq)$ , is due to the much more negative  $\Delta_{hyd}H$  of  $Cu^{2+}$  (aq) than  $Cu^{+}(aq)$ , which more than compensates for the second ionisation enthalpy of Cu.

\_\_\_\_\_\_

### **Question37**

Which element is not present in Nessler's reagent? [1-Feb-2023 Shift 2]

**Options:** 

- A. Mercury
- B. Potassium
- C. Iodine
- D. Oxygen

**Answer: D** 

**Solution:** 

Solution:

(Nessler's Reagent  $\rightarrow K_2[HgI_4]$ 

Question38

In ammonium-phosphomolybdate, the oxidation state of Mo is + \_\_\_ [6-Apr-2023 shift 1]

Answer: 6

**Solution:** 

 $(NH_4)_3PO_4 \cdot 12MoO_3$ Let X = oxidation state of Mo in  $MoO_3$ 

 $X + (-2) \times 3 = 0$ 

X = +6 Ans: 6

-----

### **Question39**

In chromyl chloride, the number of d-electrons present on chromium is same as in (Given at no. of Ti : 22, V: 23, Cr : 24, Mn : 25, Fe : 26)



### [8-Apr-2023 shift 1]

#### **Options:**

A. Fe (III)

B. V (IV)

C. Ti (III)

D. Mn (VII)

**Answer: D** 

#### **Solution:**

**Solution:** 

$$CrO_2Cl_2 \rightarrow Chromyl chloride$$
 $\downarrow \downarrow$ 
 $Cr^{+6} \rightarrow 4s^0 3d^0$ 
 $Mn(vii) \rightarrow Mn^{+7}$ 
 $\downarrow \downarrow$ 
 $4s^0 3d^0$ 

Same

### Question 40

Which halogen is known to cause the reaction given below:

 $2Cu^{2+} + 4X^{-} \rightarrow Cu_2X_2(s) + X_2$ 

[8-Apr-2023 shift 1]

### **Options:**

A. All halogens

B. Only chlorine

C. Only Bromine

D. Only Iodine

**Answer: D** 

### **Solution:**

(Only iodine) 
$$2Cu^{2+} + 4I^{-} \rightarrow Cu_2I_2 + I_2$$

### Question41

Given below are two statements:



Statement I : Aqueous solution of  $K_2Cr_2O_7$  is preferred as a primary standard in volumetric analysis over  $Na_2Cr_2O_7$  aqueous solution. Statement II :  $K_2Cr_2O_7$  has a higher solubility in water than  $Na_2Cr_2O_7$  In the light of the above statements, choose the correct answer from the options given below: [10-Apr-2023 shift 1]

#### **Options:**

- A. Statement I is false but Statement I is true
- B. Statement I is true but Statement II is false
- C. Both Statement I and Statement II are true
- D. Both Statement I and Statement II are false

**Answer: B** 

#### **Solution:**

#### **Solution:**

(1)  $K_2Cr_2O_7$  is used as primary standard. The concentration  $Na_2Cr_2O_7$  changes in aq. solution.

(2) It is less soluble than  $Na_2Cr_2O_7$ 

### Question42

Prolonged heating is avoided during the preparation of ferrous ammonium sulphate to [10-Apr-2023 shift 1]

#### **Options:**

- A. prevent hydrolysis
- B. prevent reduction
- C. prevent breaking
- D. prevent oxidation

**Answer: C** 

#### **Solution:**

#### **Solution:**

It may oxidise ferrous ion to ferric ions.

### **Question43**

Which of the following statements are correct?

- (A) The  $M^{3+}$  /  $M^{2+}$  reduction potential for iron is greater than manganese
- (B) The higher oxidation states of first row d-block elements get stabilized by oxide ion.
- (C) Aqueous solution of Cr<sup>2+</sup> can liberate hydrogen from dilute acid.
- (D) Magnetic moment of  $V^{2+}$  is observed between 4.4-5.2 BM. Choose the correct answer from the options given below:

[10-Apr-2023 shift 1]

### **Options:**

- A. (C), (D) only
- B. (B), (C) only
- C. (A), (B), (D) only
- D. (A), (B) only

**Answer: B** 

#### **Solution:**

#### **Solution:**

(A) The  ${
m M}^{3+}$  /  ${
m M}^{2+}$  reduction potential for manganese is greater than iron

(B)  $E_{Fe^{+3}/Fe^{+2}}^{0} = +0.77$ 

$$E_{M_{p+3}/M_{p+2}}^{0} = +1.57$$

$$E_{Mn^{+3}/Mn^{+2}}^{Pe^{+3}/Pe^{-2}} = +1.57$$
  
(C)  $E_{Cr^{+3}/Cr^{+2}}^{0} = -0.26$ 

$$\therefore \operatorname{Cr}^{2\oplus} + \operatorname{H}^{\oplus} \to \operatorname{Cr}^{3\oplus} + \frac{1}{2}\operatorname{H}_{2}$$

(D)  $V^{2\oplus} = 3$  unpaired electron Magnetic Moment = 3.87 B.M

### **Question44**

When a solution of mixture having two inorganic salts was treated with freshly prepared ferrous sulphate in acidic medium, a dark brown ring was formed whereas on treatment with neutral FeCl<sub>3</sub>. it gave deep red colour which disappeared on boiling and a brown red ppt was formed. The mixture contains

[11-Apr-2023 shift 1]

#### **Options:**

A. 
$$C_2O_4^{\ 2}$$
-& $NO_3$ 

B. 
$$SO_3^{2}$$
-& $C_2O_4^{2}$ -

**Answer: C** 

### **Solution:**

```
Solution:
```

```
CH<sub>3</sub>COO<sup>-</sup> + FeCl<sub>3</sub> → Fe(CH<sub>3</sub>COO)<sub>3</sub> or [Fe<sub>3</sub> (OH)<sub>2</sub> (CH<sub>3</sub>COO)<sub>6</sub>]<sup>+</sup>

Blood red colour

\downarrow \Delta

Fe(OH)<sub>2</sub> (CH<sub>3</sub>COO) \downarrow

Red-brown precipitate

2NO_3^- + 4H_2SO_4 + 6Fe^{2+} \rightarrow 6Fe^{3+} + 2NO \uparrow + 4SO_4^{2-} + 4H_2O

[Fe(H<sub>2</sub>O)<sub>6</sub>]<sup>2+</sup> + NO → [Fe(H<sub>2</sub>O)<sub>5</sub> (NO)]<sup>2+</sup> + H<sub>2</sub>O

Brown
```

-----

### Question45

The pair of lanthanides in which both elements have high third - ionization energy is: [13-Apr-2023 shift 1]

### **Options:**

A. Dy, Gd

B. Eu, Gd

C. Lu, Yb

D. Eu, Yb

**Answer: D** 

### **Solution:**

#### Solution:

 $\mathrm{Eu^{+2}:[Xe]4f^7}$  $\mathrm{Yb^{+2}:[Xe]4f^{14}}$  High IE due to half filled & fully filled configurations

------

### **Question46**

In Chromyl chloride, the oxidation state of chromium is (+) \_\_\_\_\_. [15-Apr-2023 shift 1]

**Answer: 6** 

$$CrO_2 CO_2$$
 (Chromylchloride)  
 $\downarrow$   
 $x-4-2=0$   
 $\boxed{x=+6}$ 

\_\_\_\_\_

### **Question47**

The difference in oxidation state of chromium in chromate and dichromate salts is \_\_\_\_ [24-Jun-2022-Shift-1]

Answer: 0

**Solution:** 

#### **Solution:**

Chromate ion  $\rightarrow$  CrO<sub>4</sub><sup>2-</sup>, oxidation state of Cr = +6 Dichromate ion  $\rightarrow$  CrO<sub>7</sub><sup>2-</sup>, oxidation state of Cr = +6  $\rightarrow$  Difference in oxidation state = zero

\_\_\_\_\_

### **Question48**

Manganese (VI) has ability to disproportionate in acidic solution. The difference in oxidation states of two ions it forms in acidic solution is\_\_\_\_

[24-Jun-2022-Shift-2]

**Answer: 3** 

**Solution:** 

#### **Solution:**

Manganese (VI) disproportionates in acidic medium as  $3MnO_4^{-2} + 4H^+ \longrightarrow 2MnO_4^{-} + MnO_2 + 2H_2O$ The difference in oxidation states of Mn in the products formed = 7 – 4 = 3

### **Question49**

Cerium (IV) has a noble gas configuration. Which of the following is

### correct statement about it? [25-Jun-2022-Shift-1]

### **Options:**

- A. It will not prefer to undergo redox reactions.
- B. It will prefer to gain electron and act as an oxidizing agent.
- C. It will prefer to give away an electron and behave as reducing agent.
- D. It acts as both, oxidizing and reducing agent.

**Answer: B** 

#### **Solution:**

#### Solution:

Cerium exists in two different oxidation state +3, +4 $Ce^{+4} + e^{-} \rightarrow Ce^{3+} E^{0} = +1.61V$   $Ce^{+3} + 3e^{-} \rightarrow Ce E^{0} = -2.336V$ 

It shows Ce<sup>+4</sup> acts as a strong oxidising agent \& accepts electron.

### Question 50

### Among the following, which is the strongest oxidizing agent? [25-Jun-2022-Shift-1]

### **Options:**

- A. Mn<sup>3+</sup>
- B.  $Fe^{3+}$
- $C Ti^{3+}$
- D. Cr<sup>3+</sup>

**Answer: A** 

### **Solution:**

Strongest oxidising agent have highest reduction potential value

$$E^{\circ}_{\ Fe^{3+} \ | \ Fe^{2+}} = +0.77 V \ E^{\circ}_{\ Ti^{3+} \ | \ Ti^{2+}} = -0.37 V$$

 $E^{\circ}_{Mn^{+3} \mid Mn^{+2}} = +1.57V \ E^{\circ}_{Cr^{3+} \mid Cr^{2+}} = -0.41V$ 

 $\mathrm{Mn}^{+3}$  is the best oxidising agent among the given series.

### Question51

The metal ion (in gaseous state) with lowest spin-only magnetic moment value is

### [25-Jun-2022-Shift-2]

**Options:** 

A.  $V^{2+}$ 

B. Ni<sup>2+</sup>

C. Cr<sup>2+</sup>

D. Fe<sup>2+</sup>

Answer: B

### **Solution:**

#### Solution:

	Valence shell configuration	Unpaired electrons
V <sup>2+</sup>	3d <sup>3</sup> 4s <sup>0</sup>	n = 3
Ni <sup>2+</sup>	3d <sup>8</sup> 4s <sup>0</sup>	n = 2
Cr <sup>2+</sup>	3d <sup>4</sup> 4s <sup>0</sup>	n = 4
Fe <sup>2+</sup>	3d <sup>6</sup> 4s <sup>0</sup>	n = 4

\_\_\_\_\_

### **Question52**

The spin-only magnetic moment value of the most basic oxide of vanadium among  $V_2O_3$ ,  $V_2O_4$  and  $V_2O_5$  is\_\_\_\_\_ B.M. (Nearest integer) [26-Jun-2022-Shift-1]

**Answer: 3** 

#### **Solution:**

#### Solution:

The most basic oxide among  $V_2O_3$ ,  $V_2O_4$  and  $V_2O_5$  is  $V_2O_3$   $V_2O_3=V^{+3}(d^2)$  Magnetic moment  $=\sqrt{2(2+2)}=\sqrt{8}$ 

Magnetic moment =  $\sqrt{2(2+2)} = \sqrt{8}$ = 2.83  $\approx$  3

### Question53

The most common oxidation state of Lanthanoid elements is +3. Which of the following is likely to deviate easily from +3 oxidation state?



[26-Jun-2022-Shift-2]
Options:
A. Ce (At. No. 58)
3. La (At. No. 57)
C. Lu (At. No. 71)
D. Gd (At. No. 64)
Answer: A
Solution:
Solution: $Ce \rightarrow [Xe]4f^{1}5d^{1}6s^{2}$ $Ce^{+4} \rightarrow [xe]4f^{\circ}5d^{\circ}6s^{\circ}$ Cerium in +4 oxidation state acquires inert gas configuration.
Question54
(A) Cu(II) complexes are always paramagnetic. (B) Cu(I) complexes are generally colourless (C) Cu(I) is easily oxidized (D) In Fehling solution, the active reagent has Cu(I) [27-Jun-2022-Shift-1]
Answer: 3
Solution:
<b>Solution:</b> A) $Cu$ (II) complexes are always paramagnetic as they have one unpaired electron due to $d^9$ configuration of $Cu(II)$ B) $Cu(I)$ complexes are generally colourless due to $d^{10}$ configuration. C) $Cu(I)$ is easily oxidised to $Cu^{+2}$ in aqueous solution $2Cu^+ \rightarrow Cu^{+2} + Cu$ $Cu^{+1}$ disproportionates to $Cu^{+2}$ and $Cu$ $E_{cell}^{\ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ $
Question55
Acidified potassium permanganate solution oxidises oxalic acid. The spin-only magnetic moment of the manganese product formed from the above reaction isB.M. (Nearest integer)

### [27-Jun-2022-Shift-1]

**Answer: 6** 

#### **Solution:**

```
KMnO<sub>4</sub> (acidic medium) + H_2C_2O_4 → CO_2 + Mn^{+2} Mn<sup>+2</sup> has 5 unpaired electrons

∴ Spin only magnetic moment = \sqrt{5(5+2)} = \sqrt{5} \times 7 = \sqrt{35} ≈ 5.92 B.M.

≈ 6 B.M.
```

### Question 56

The 'f' orbitals are half and completely filled, respectively in lanthanide ions

[Given : Atomic no. Eu, 63; Sm, 62; Tm, 69; Tb, 65; Yb, 70; Dy, 66] [27-Jun-2022-Shift-2]

### **Options:**

- A. Eu<sup>2+</sup> and Tm<sup>2+</sup>
- B.  $Sm^{2+}$  and  $Tm^{3+}$
- C.  $Tb^{4+}$  and  $Yb^{2+}$
- D.  $Dy^{3+}$  and  $Yb^{3+}$

**Answer: C** 

### **Solution:**

```
Solution:
```

```
T b \rightarrow 4f^{9}6s^{2}
Tb^{+4} \rightarrow 4f^{7}
Yb \rightarrow 4f^{14}6s^{2}
Yb^{+2} \rightarrow 4f^{14}
```

Hence, the pair  ${\rm Tb}^{+4}$  and  ${\rm Yb}^{+2}$  have half filled and completely filled f subshells respectively.

-----

### Question57

Dihydrogen reacts with CuO to give [28-Jun-2022-Shift-1]

**Options:** 



A. CuH<sub>2</sub>

B. Cu

C. Cu<sub>2</sub>O

D. Cu(OH)<sub>2</sub>

**Answer: B** 

### **Solution:**

#### **Solution:**

 $\mathrm{CuO} + \mathrm{H_2} \rightarrow \mathrm{Cu} + \mathrm{H_2O}$  (under hot conditions)

### **Question58**

Which one of the lanthanoids given below is the most stable in divalent form?

[28-Jun-2022-Shift-1]

### **Options:**

A. Ce (Atomic Number 58)

B. Sm (Atomic number 62)

C. Eu (Atomic Number 63)

D. Yb (Atomic Number 70)

**Answer: C** 

#### **Solution:**

#### Solution:

Electronic configuration of Europium (Eu) is  $[Kr]4f^76 s^2$ . It can loose two electron to gain the half filled stable state. Hence, +2 oxidation state is most stable.

### Question59

The number of terminal oxygen atoms present in the product B obtained from the following reaction is

$$FeCr_2O_4 + Na_2CO_3 + O_2 \rightarrow A + \overline{Fe_2O_3} + CO_2$$

$$A + H^+ \rightarrow B + H_2O + Na^+$$

[29-Jun-2022-Shift-1]



**Answer: 6** 

#### **Solution:**

$$\begin{aligned} & \operatorname{FeCr_2O_4} + \operatorname{Na_2CO_3} + \operatorname{O_2} \rightarrow \operatorname{Fe_2O_3} + \operatorname{CO_2} + \operatorname{Na_2CrO_4} \\ & \operatorname{Na_2CrO_4} + \operatorname{H}^+ \rightarrow \operatorname{Cr_2O_7}^{-2} + \operatorname{H_2O} + \operatorname{Na}^+ \end{aligned}$$

$$_{2\text{Na}^{+}}\begin{bmatrix}0\\O\\O\end{bmatrix}\text{Cr}\begin{bmatrix}O\\O\\O\end{bmatrix}^{2-}$$

-----

### Question60

An acidified manganate solution undergoes disproportionation reaction. The spin-only magnetic moment value of the product having manganese in higher oxidation state is \_\_\_\_\_B.M. (Nearest integer) [29-Jun-2022-Shift-1]

Answer: 0

**Solution:** 

**Solution:** 

 $3MnO_4^{\ 2^-} + 4H^+ \longrightarrow 2MnO_4^{\ -} + MnO_2^{\ +4} + 2H_2O$   $Mn^{+7} = \text{ no. of unpaired electrons is ' 0 '}$  $\mu = 0 \ B.M.$ 

.....

### Question61

The reaction of  $H_2O_2$  with potassium permanganate in acidic medium leads to the formation of mainly : [25-Jul-2022-Shift-1]

**Options:** 

A.  $M n^{2+}$ 

B. M n<sup>4+</sup>

C.  $M n^{3+}$ 

D. M n<sup>6+</sup>

**Answer: A** 



\_\_\_\_\_

### **Question62**

Among  $\mathrm{Co}^{3+}$ ,  $\mathrm{Ti}^{2+}$ ,  $\mathrm{V}^{2+}$  and  $\mathrm{Cr}^{2+}$  ions, one if used as a reagent cannot liberate H  $_2$  from dilute mineral acid solution, its spin-only magnetic moment in gaseous state is B.M. (Nearest integer) [25-Jul-2022-Shift-1]

Answer: 5

**Solution:** 

**Solution:** 

 ${\rm Co}^{3+}$  will not liberate  ${\rm H_2}$  gas an reaction with dilute acid  ${\rm F}$  0 = +1.07

 $E_{\text{Co}^{3+}/\text{Co}^{2+}}^{O} = +1.97$ 

And  $Co^{3+}$  has electronic configuration = [Ar]3d  $^{6}$ 

 $\div 4 \text{ unpaired } e^- \text{are present in it}$ 

∴ Spin-only magnetic moment  $= \sqrt{4(4+2)} = 4.92 \approx 5$ 

-----

### Question63

The spin-only magnetic moment value of the compound with strongest oxidizing ability among  $MnF_4$ ,  $MnF_3$  and  $MnF_2$  is \_\_\_\_\_ B.M. [nearest integer]

[26-Jul-2022-Shift-2]

**Answer: 5** 

**Solution:** 

**Solution:** 

$$MnF_4$$
  $MnF_3$   $MnF_2$   
 $+4$   $+3$   $+2$   
 $E.C = [Ar] 3d^3$   $[Ar] 3d^4$   $[Ar] 3d^5$ 

Hence  $MnF_3 \Rightarrow$  strongest O.A

 $\mu = \sqrt{4(4+2)} = \sqrt{24} = 4.89 = 5$ 

-----





Given below are two statements.

Statement I: Iron (III) catalyst, acidified K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> and neutral KMnO<sub>4</sub>

have the ability to oxidise  $I^-$  to  $I_2$  independently.

Statement II: Manganate ion is paramagnetic in nature and involves  $p\pi - p\pi$  bonding.

In the light of the above statements, choose the correct answer from the options given below.

[27-Jul-2022-Shift-1]

### **Options:**

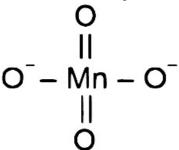
- A. Both Statement I and Statement II are true.
- B. Both Statement I and Statement II are false.
- C. Statement I is true but Statement II is false.
- D. Statement I is false but Statement II is true.

**Answer: B** 

#### **Solution:**

#### **Solution:**

Manganate ion  $MnO_4^{2-}$  has tetrahedral structure



has only  $d\pi - p\pi\pi$ -bonds.

 ${\rm Fe}^{3+}$  is not used as a catalyst in the conversion of I' to  ${\rm I_2}$  by  ${\rm K_2Cr_2O_7\cdot K_2Cr_2O_7}$  oxidise I $^-$ in acidic medium easily

### Question65

The total number of Mn = O bonds in  $Mn_2O_7$  is \_\_\_\_\_. [27-Jul-2022-Shift-1]

#### **Options:**

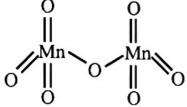
- A. 4
- B. 5
- C. 6
- D. 3

**Answer: C** 

### **Solution:**

**Solution:** 

Structure of mathrm  $M n_2$  mathrm  $O_7$  is as :



 $\therefore$  There are total 6M = O bonds are present in  $Mn_2O_7$  compound.

### **Question66**

In the titration of  $KMnO_4$  and oxalic acid in acidic medium, the change in oxidation number of carbon at the end point is \_\_\_\_\_. [27-Jul-2022-Shift-1]

**Answer: 1** 

**Solution:** 

**Solution:** 

 $16 \text{H}^+ + 2 \text{MnO}_4^- + 5 \text{C}_2 \text{O}_4^{\ 2-} \rightarrow 10 \text{CO}_2 + 2 \text{Mn}^{2+} + 8 \text{H}_2 \text{O}$ 

During titration of oxalic acid by KMnO<sub>4</sub>, oxalic acid converts into CO<sub>2</sub>.

 $\therefore$  Change in oxidation state of carbon = 1

\_\_\_\_\_\_

### Question67

In neutral or alkaline solution,  ${\rm MnO_4}^{-}{\rm oxidises}$  thiosulphate to : [27-Jul-2022-Shift-2]

**Options:** 

A. 
$$S_2O_7^{2-}$$

B. 
$$S_2O_8^{2}$$

**Answer: D** 

------

## **Question68**

The oxidation state of manganese in the product obtained in a reaction of potassium permanganate and hydrogen peroxide in basic medium is

\_\_\_\_. [27-Jul-2022-Shift-2]

**Answer: 4** 

**Solution:** 

**Solution:** 

$$2 \text{KMnO}_4 + 3 \text{H}_2 \text{O}_2 \overset{\text{basic medium}}{\longrightarrow} 2 \text{MnO}_2 + 3 \text{O}_2 + 2 \text{H}_2 \text{O} + 2 \text{ KOH}$$

\_\_\_\_\_

### **Question69**

Which of the following has least tendency to liberate H<sub>2</sub> from mineral acids?

[28-Jul-2022-Shift-1]

**Options:** 

A. Cu

B. Mn

C. Ni

D. Zn

**Answer: A** 

**Solution:** 

The metal atom whose oxidation potential is less than that of hydrogen can release  $H_2$  from mineral acids.

$$E_{Zn/Zn^{+2}} = 0.76$$

$$E_{Ni/Ni^{\circ 2}} = 0.25$$

$$E_{Mn/Mn^{+2}} = 1.18$$

$$E_{Cu^{\circ}/Cu^{+2}} = -0.34$$

-----





The disproportionation of  $MnO_4^{\ 2^-}$  in acidic medium resulted in the formation of two manganese compounds A and B. If the oxidation state of M n in B is smaller than that of A, then the spin-only magnetic moment ( $\mu$ ) value of B in BM is \_\_\_\_\_.(Nearest integer) [28-Jul-2022-Shift-1]

**Answer: 4** 

### **Solution:**

```
Solution:

3\text{MnO}_4^{-2} + 4\text{H}^+ \rightarrow \text{MnO}_2 + \text{MnO}_4^-

\text{Mn} \rightarrow 4\text{s}^2 3\text{d}^5

\text{Mn}^{+4} \rightarrow 3\text{d}^3

\text{n} = 3

\mu = \sqrt{\text{n(n+2)}}

= \sqrt{3}(5)

= \sqrt{15}

= 3.87 \approx 4 \text{ B.M.}
```

### Question71

Which of the following pair is not isoelectronic species? (At. no. Sm, 62; Er, 68; Yb, 70; Lu, 71; Eu, 63; Tb, 65; Tm, 69) [28-Jul-2022-Shift-2]

#### **Options:**

B. 
$$Yb^{2+}$$
 and  $Lu^{3+}$ 

D. 
$$Tb^{2+}$$
 and  $Tm^{4+}$ 

**Answer: 0** 

#### **Solution:**

#### **Solution:**

Species having same number of electrons are isoelectronic

Tb 
$$\rightarrow 65$$
Tb<sup>+2</sup>  $\rightarrow 63$  electrons
Tm  $\rightarrow 69$ Tm<sup>+4</sup>  $\rightarrow 65$  electrons
Sm<sup>+2</sup>  $\rightarrow 60$  electrons
Er<sup>+3</sup>  $\rightarrow 65$  electrons
$$\begin{array}{c} \text{not isoelectronic} \\ \text{not isoelectronic} \end{array}$$

The reaction of zinc with excess of aqueous alkali, evolves hydrogen gas and gives:

[29-Jul-2022-Shift-1]

### **Options:**

- A. Zn(OH)<sub>2</sub>
- B. ZnO
- C.  $[Zn(OH)_4]^{2-}$
- D.  $[ZnO_2]^{2-}$

**Answer: C** 

#### **Solution:**

#### **Solution:**

Zinc dissolves in excess of aqueous alkali.  $Zn + 2OH^- + 2H_2O \longrightarrow [Zn(OH)_4]^{2^-} + H_2 \uparrow \text{ (Tetrahydroxozincate(II) ion)} \\ \text{However, this reaction in NCERT is given as} \\ Zn + 2\,NaOH \longrightarrow Na_2ZnO_2 + H_2 \uparrow \\ ZnO_2^{2^-} \text{ is anhydrous form of } [Zn(OH)_4]^{2^-}. \\ ZnO_2^{2^-} + 2H_2O \rightleftharpoons [Zn(OH)_4]^{2^-} \\ \text{So in aqueous medium best answer of this question is } [Zn(OH)_4]^{2^-}.$ 

\_\_\_\_\_

# Question73

In following pairs, the one in which both transition metal ions are colourless is : [29-Jul-2022-Shift-1]

## **Options:**

- A.  $Sc^{3+}$ ,  $Zn^{2+}$
- B. Ti<sup>4+</sup>, Cu<sup>2+</sup>
- C.  $V^{2+}$ ,  $Ti^{3+}$
- D.  $Zn^{2+}$ ,  $Mn^{2+}$

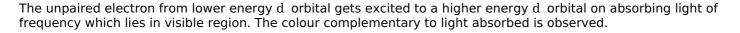
**Answer: A** 

#### **Solution:**

 $Sc^{+3}$  and  $Zn^{+2}$  are colourless as they contain no unpaired electron. Whereas the transition metal ions  $Cu^{+2}$ ,  $Ti^{+3}$ ,  $V^{+2}$  and  $Mn^{+2}$  are coloured as they contain unpaired electrons.







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# **Question74**

In neutral or faintly alkaline medium,  $KMnO_4$  being a powerful oxidant can oxidize, thiosulphate almost quantitatively, to sulphate. In this reaction overall change in oxidation state of manganese will be : [29-Jul-2022-Shift-1]

#### **Options:**

A. 5

B. 1

C. 0

D. 3

Answer: D

#### **Solution:**

#### **Solution:**

In neutral or Faintly alkaline medium, thiosulphate is oxidised almost quantitatively to sulphate ion according to reaction given below,

 $8MnO_4^- + 3S_2O_3^{2-} + H_2O \rightarrow 8MnO_2 + 6SO_4^{2-} + 2OH^-$ 

Here the Mn changes from  $Mn^{+7}$  to  $Mn^{+4}$ 

Thus overall change in its oxidation number would be of 3.

\_\_\_\_\_

## Question 75

Which of the following 3d-metal ion will give the lowest enthalpy of hydration ( $\Delta_{hyd}$  H) when dissolved in water? [29-Jul-2022-Shift-2]

#### **Options:**

A. Cr<sup>2+</sup>

B. Mn<sup>2+</sup>

C. Fe<sup>2+</sup>

D.  $Co^{2+}$ 

Answer: B



Ion	$\Delta H_{Hyd}^{O}(kJ/mole)$
Cr <sup>2+</sup>	-1925
Mn <sup>2+</sup>	-1862
Fe <sup>2+</sup>	-1998
Co <sup>2+</sup>	-2079

\_\_\_\_\_

# **Question76**

# The major components of German silver are [25 Feb 2021 Shift 2]

## **Options:**

A. Cu, Zn and Ag

B. Cu, Zn and Ni

C. Ge, Cu and Ag

D. Zn, Ni and Ag

**Answer: B** 

#### **Solution:**

#### **Solution:**

The major components of German silver are Cu: 25-30%, Zn: 25-30% and Ni: 40-30%.

\_\_\_\_\_\_

# **Question77**

In which of the following pairs, the outer most electronic configuration will be the same?
[25 Feb 2021 Shift 1]

### **Options:**

A.  $V^{2+}$  and  $Cr^{+}$ 

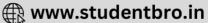
B. Cr<sup>+</sup>and M n<sup>2+</sup>

C. N i<sup>2+</sup> and Cu<sup>+</sup>

D.  $Fe^{2+}$  and  $Co^{+}$ 

**Answer: B** 





(a)  $V^{2+} - [Ar]3d^3$ ;  $Cr^+ - [Ar]3d^5$ (b)  $Cr^+ - [Ar]3d^5$ ;  $M n^{2+} - [Ar]3d^5$ (c)  $N i^{2+} - [Ar]3d^8$ ;  $Cu^+ - [Ar]3d^{10}$ (d)  $F e^{2+} - [Ar]3d^6$ ;  $Co^+ - [Ar]3d^7 4s^1$ Thus, in option (b), both ions have same outer most electronic configuration.

\_\_\_\_\_

# **Question78**

What is the correct order of the following elements with respect to their density?

[24 Feb 2021 Shift 2]

### **Options:**

A. Cr < Zn < Co < Cu < Fe

B. Zn < Cu < Co < Fe < Cr

C. Zn < Cr < Fe < Co < Cu

D. Cr < Fe < Co < Cu < Zn

**Answer: C** 

#### **Solution:**

#### **Solution:**

Generally, due to decrease in metallic radius and increase in atomic mass density increase across the period from left to right.

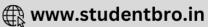
Metal	Density (g/cm³)
Zn	7.13
Cr	7.19
Fe	7.8
Со	8.7
Cu	8.9

Correct order is Cu > Co > Fe > Cr > Zn.

-----

# **Question79**

**Answer: 6** 



#### **Solution:**

#### Solution:

Solution:
$$S_{2}O_{3}^{2-} + MnO_{4}^{-} \xrightarrow{Mild} SO_{4}^{2-} + MnO_{4}^{2-}$$
Thiosulphate ion [Reducing agent] Permanganate ion Alkaline sulphate ion(A) Manganate ion Manganate ion (Oxidation number of sulphur in  $SO_{4}^{2-}(A)$  is  $+6$ 

$$[ :x + 4(-2) = -2 ]$$

$$\Rightarrow x = +6$$

# Question80

On treating a compound with warm dil. H <sub>2</sub>SO<sub>4</sub>, gas X is evolved, which turns K<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> paper acidified with dil. H<sub>2</sub>SO<sub>4</sub> to a green compound Y . X and Y respectively are [26 Feb 2021 Shift 1]

### **Options:**

$$A. X = SO_2, Y = Cr_2O_3$$

B. 
$$X = SO_3$$
,  $Y = Cr_2O_3$ 

$$C. X = SO_2, Y = Cr_2(SO_4)_3$$

D. 
$$X = SO_3$$
,  $Y = Cr_2(SO_4)_3$ 

**Answer: C** 

#### **Solution:**

Gas, X tums acidified dichromate ( $K_2Cr_2O_7$ ) green that means it is a reducing agent.

Cr (VI) from compound  $K_{22}Cr_2O_7$  on reduction changes its colour from orange to green which is the colour of Cr (III)

 $SO_2$  can show both reducing and oxidising properties whereas  $SO_3$  cannot show reducing property because of highest group number 16, oxidation state (+6) of sulphur in  $SO_3$ .

As, X reacts with K 2Cr2O7 in dil. H 2SO4 medium, the green coloured Cr (III) compound, Y must be Cr (III) sulphate or  $Cr_2(SO_4)_3$ .

So, Y is  $Cr_2(SO_4)_3$ .

The overall reactions can be shown as : 
$$(Compound) \xrightarrow{Dil.} SO_2(g) \xrightarrow{K_2Cr_2O_7} Cr_2(SO_4)_3$$
 water soluble  $SO_3^-$  salt  $H_2SO_4$   $(X)$   $dil.H_2SO_4$   $(Y)$   $Green$   $(i) SO_3^{-2} + H_2SO_4 \rightarrow SO_2 \uparrow + H_2O + SO_4^{-2}$   $(X)$   $(Y)$   $(Green)$ 





On treating a compound with warm dil. H  $_2\mathrm{SO}_4$ , gas X is evolved, which turns K  $_2\mathrm{Cr}_2\mathrm{O}_7$  paper acidified with dil. H  $_2\mathrm{SO}_4$  to a green compound Y . X and Y respectively are [26 Feb 2021 Shift 1]

### **Options:**

A. 
$$X = SO_2$$
,  $Y = Cr_2O_3$ 

B. 
$$X = SO_3$$
,  $Y = Cr_2O_3$ 

$$C. X = SO_2, Y = Cr_2(SO_4)_3$$

D. 
$$X = SO_3, Y = Cr_2(SO_4)_3$$

**Answer: C** 

#### **Solution:**

Gas, X tums acidified dichromate ( $K_2Cr_2O_7$ ) green that means it is a reducing agent.

Cr (VI) from compound  $K_{22}Cr_2O_7$  on reduction changes its colour from orange to green which is the colour of Cr (III) compound Y.

 $SO_2$  can show both reducing and oxidising properties whereas  $SO_3$  cannot show reducing property because of highest group number 16, oxidation state (+6) of sulphur in  $SO_3$ .

So, X is  $SO_2$ 

As, X reacts with K  $_2$ Cr $_2$ O $_7$  in dil. H  $_2$ SO $_4$  medium, the green coloured Cr (III) compound, Y must be Cr (III) sulphate or Cr $_2$ (SO $_4$ ) $_3$ .

So, Y is  $Cr_2(SO_4)_3$ .

The overall reactions can be shown as

(Compound) Dil. 
$$SO_{2}(g)$$
  $Cr_{2}(SO_{4})_{3}$   $Cr_{2}(SO_{4})_{4}$   $Cr_{2}(SO_{4})_{4}$   $Cr_{2}(SO_{4})_{4}$   $Cr_{2}(SO_{4})_{4}$   $Cr_{2}(SO_{4})_{4}$ 

\_\_\_\_\_

## **Question82**

Which one of the following lanthanoids does not form  $MO_2$ ? [ M is lanthanoid metal] [26 Feb 2021 Shift 1]

#### **Options:**

- A. Pr
- B. Dy
- C. Nd
- D. Yb

**Answer: D** 

#### **Solution:**

#### **Solution:**

In oxides, M  $\rm O_2$  ( M is lanthanoid metal) only four lanthanoids exhibit +4 oxidation state. These lanthanoids are praseodymium (Pr, Z = 59), neodymium (N d , Z = 60) terbium (T b, Z = 65) dysprosium (Dy, Z = 66) So, Y b (ytterbium) option (d) does not form M  $\rm O_2$  type of oxide.

**Note** The common and predominant oxidation state of lanthanoids is +3. Consequently M  $^{4+}$  compounds are strong oxidising agents which changes to the common +3 state. Similarly, lanthanoid compounds of +2 state have a tendency to show reducing property as they get changed to +3 state easily.

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# **Question83**

#### Given below are two statements:

Statement I CeO<sub>2</sub> can be used for oxidation of aldehydes and ketones.

Statement II Aqueous solution of EuSO  $_{4}$  is a strong reducing agent.

In the light of the above statements, choose the correct answer from the options given below.

[25 Feb 2021 Shift 1]

#### **Options:**

- A. Both statement I and statement II are true.
- B. Both statement I and statement II are false.
- C. Statement I is true but statement II is false.
- D. Statement I is false but statement II is true.

**Answer: A** 

#### **Solution:**

#### Solution:

Both statement I and statement II are true. The +3 oxidation state of lanthanide is most stable and therefore lanthanide in +4 oxidation state has strong tendency to gain electrons and converted into +3 and therefore act as strong oxidising agent, e.g.  $Ce^{4+}$ .

 $\therefore$ CeO<sub>2</sub> is used to oxidised alcohol, aldehyde and ketones. Lanthanides in +2 oxidation state has strong tendency to loss electron and converted into +3 oxidation state e.g. E u<sup>+2</sup>.

 $\therefore$  EuSO  $_4$  acts as strong reducing agent.

# Question84

# The major components in "Gun Metal" are: [24 Feb 2021 Shift 1]

#### **Options:**

- A. Cu, Zn and Ni
- B. Cu, Sn and Zn



C. Al , Cu, Mg and Mn
D. Cu, Ni and Fe
Answer: B
Solution:
Solution: The major components in "Gun Metal" are Cu : 87%; Z n : 3%; Sn : 10%.
Question85
The electrode potential of M $^{2+}$ / M of 3d -series elements shows positive value of [24 Feb 2021 Shift 1]
Options:
A. Z n
B. Fe
C. Co
D. Cu
Answer: D
Solution:
<b>Solution:</b> Only copper shows positive value for electrode potential of M <sup>2+</sup> / M of 3d -series elements.
Question86
The incorrect statement among the following is [24 Feb 2021 Shift 2]
Options:
A. VOSO $_4$ is a reducing agent.
B. $Cr_2O_3$ is an amphoteric oxid
${\rm C.~RuO_4}$ is an oxidising agent.
D. Red colour of ruby is due to the presence of Co <sup>3+</sup> .

**Answer: B** 

Red colour of ruby is due to presence of  $Cr^{3+}$  ions in Al  $_2O_3$ .

Chromium is the trace element that causes ruby's red colour, which ranges from an orange red to a publish red. The strength of ruby's red depends on how much chromium is present.

-----

## **Question87**

# The common positive oxidation states for an element with atomic number 24 , are [17 Mar 2021 Shift 2]

#### **Options:**

A. +2 to +6

B. +1 and +3 to +6

C. +1 and +3

D. +1 to +6

**Answer: A** 

#### **Solution:**

#### **Solution:**

Common positive oxidation states for an element with atomic number 24 , are +2 to +6 Chromium (  $Z\,$  =  $\,24$  )

Electronic configuration: [Ar]  $4s^13d^5$  It has five electrons in 3d-subshell and one electron in 4s-subshell. Thus, chromium metal has six valence electrons.

Chromium can lose some or all of its valence electrons to form ions with different oxidation states. Thus, chromium shows the oxidation states of +1, +2, +3, +4, +5 and +6 respectively.

The most common oxidation states of chromium are +2, +3 and +6.

Question88

# The oxide that shows magnetic property is [18 Mar 2021 Shift 2]

#### **Options:**

A. SiO<sub>2</sub>

B.  $M n_3 O_4$ 

C. N  $a_2$ O

D. MgO

**Answer: B** 



Generally, d-block elements shows magnetic property due to presence of unpaired electrons in its d-block. Mn i.e. manganese shows +2 and +3 oxidation state.

Both having unpaired electrons as shown below

$$_{25}$$
M n = [Ar]3d  $^5$ 4s<sup>2</sup>

 ${\rm M}\,{\rm n}^{2+}({\rm 3d}\,{}^5{\rm 4s}^0)$  having 5 unpaired electrons in 3d orbital.

 $\mathrm{M}\,\mathrm{n}^{+3}$ ,  $(\mathrm{3d}\,^4\mathrm{4s}^0)$  having 4 unpaired electrons in 3d orbital.

 $\mathrm{SiO}_{2}$ , p-block element compound is commonly believed to be a typical diamagnetic as no unpaired electrons are present.

 $M\ gO$  and  $N\ a_2O$  also have unpaired electrons and hence are diamagnetic.

Hence,  $M n_3 O_4$  show magnetic property.

### \_\_\_\_\_

# **Question89**

# Fex $_2$ and $_{Lt}$ are known when x and y are [16 Mar 2021 Shift 2]

#### **Options:**

A. x = F, Cl, Br, I and y = F, Cl and Br

B. 
$$x = F$$
, Cl, Br and  $y = F$ , Cl, Br and I

C. x = Cl, Br, I and y = F, Cl, Br and I

D. 
$$x = F$$
, Cl, Br, I and  $y = F$ , Cl, Br and I

**Answer: A** 

#### **Solution:**

 $F\ el\ _3$  does not exist because of  $I\ ^-$ . Being a very good reducing agent,  $I\ ^-$ reduces  $F\ e^{3+}$  to  $F\ e^{2+}$ .

$$2Fe^{3+} + 2I^{-} \longrightarrow 2Fe^{2+} + I_{2}$$

Reducing nature order I > Br > Cl > F

$$F e^{3+} + e^{-} \rightarrow F e^{2+} E^{\circ} = 0.77V$$

$$I_2 + 2e^- \rightarrow 2I^- E^\circ = -0.54V$$

As standard reduction potential value of F  $e^{3+}$  is more than I  $_2$ , so it is reduced to F  $e^{2+}$ . Hence, except F  $e^{1}$   $_3$ , all other halogens can form iron halide compounds.

# Question90

Given below are two statements.

Statement I The E  $^{\circ}$  value of  $CC^{4+}/Ce^{3+}$  is +1.74V.

Statement II Ce is more stable in Ce<sup>4+</sup> state than Ce<sup>3+</sup> state.

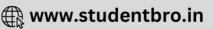
In the light of the above statements, choose the most appropriate answer from the options given below.

[16 Mar 2021 Shift 1]

#### **Options:**

- A. Both statement I and statement II are correct.
- B. Statement I is incorrect but statement II is correct.





- C. Both statement I and statement II are incorrect.
- D. Statement I is correct but statement II is incorrect.

**Answer: D** 

### **Solution:**

#### **Solution:**

 $\rm E_{\rm Ce^{4+}/Ce^{+3}}^{\circ} = 1.74$  (data given)

 $\mathrm{Ce}^{4+}$  can be reduced to  $\mathrm{Ce}^{3+}$  and can oxidise water.

 ${\rm Ce}^{4+}$  is having noble gas configuration [Xe]  $4{\rm f}^{\,0}$  but it is strong oxidant and gets converted to  ${\rm Ce}^{3+}$ . Statement I is correct but statement II is incorrect.

# Question91

Given below are two statements. One is labelled as Assertion (A) and the other is labelled as Reason (R).

Assertion (A) Size of  $Bk^{3+}$  ion is less than  $Np^{3+}$  ion.

Reason (R) The above is a consequence of the lanthanoid contraction. In the light of the above statements, choose the correct answer from the options given below

[16 Mar 2021 Shift 1]

#### **Options:**

- A. A is false but R is true
- B. Both A and R are true but R is not the correct explanation of A.
- C. Both A and R are true and R is the correct explanation of A.
- D. A is true but R is false.

**Answer: D** 

### **Solution:**

 $_{97}{
m Bk}$  (Berkelium) and  $_{93}{
m N}$  p (Neptunium) are present in actinide series. Due to actinoid contraction, there is gradual decrease in size of M <sup>3+</sup> ions across the period due to poor shielding of 5f -electrons. So, Bk<sup>3+</sup> is smaller than that of N p<sup>3+</sup> due to actinoid contraction. So, A is true and R is false.

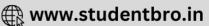
# Question 92

The spin only magnetic moments (in BM) for free T  $i^{3+}$ ,  $V^{2+}$  and  $Sc^{3+}$ ions respectively are

(At.No. Sc : 21, Ti : 22, V: 23)

[25 Jul 2021 Shift 2]





### **Options:**

A. 3.87, 1.73, 0

B. 1.73, 3.87, 0

C. 1.73, 0, 3.87

D. 0, 3.87, 1.73

**Answer: B** 

#### **Solution:**

$$\mu = \sqrt{n(n+2)}BM$$
 
$$T \, 1^{+3} = [Ar]3d^{\,1} \quad n=1 \qquad \mu = 1.73BM$$
 
$$V^{\,+2} = [Ar]3d^{\,3} \quad n=3 \qquad \mu = 3.87BM$$
 
$$Sc^{+3} = [Ar]3d^{\,0}4s^0 \quad n=0 \qquad \qquad \mu = 0$$

.....

# **Question93**

The correct order of following 3d metal oxides, according to their oxidation numbers is :

(a)  $CrO_3$ 

(b)  $Fe_2O_3$ 

(c) M nO<sub>2</sub>

(d)  $V_2O_5$ 

(e)  $Cu_2O$ 

[25 Jul 2021 Shift 1]

## **Options:**

A. 
$$(d) > (a) > (b) > (c) > (e)$$

B. 
$$(a) > (c) > (d) > (b) > (e)$$

C. (a) 
$$>$$
 (d)  $>$  (c)  $>$  (b)  $>$  (e)

D. (c) 
$$>$$
 (a)  $>$  (d)  $>$  (e)  $>$  (b)

**Answer: C** 

## **Solution:**

(a) 
$$\overset{+6}{\text{CrO}}_3$$

(b) 
$$F_{e_2}^{+3}O_3$$

(c) 
$$M nO_2$$

(d) 
$$\overset{+5}{\mathrm{V}}_{2}\mathrm{O}_{5}$$



The set having ions which are coloured and paramagnetic both is -[22 Jul 2021 Shift 2]

**Options:** 

B. 
$$Cu^{2+}$$
,  $Z n^{2+}$ ,  $M n^{4+}$ 

C. 
$$Sc^{3+}$$
,  $V^{5+}$ ,  $Ti^{4+}$ 

D. N 
$$i^{2+}$$
, M  $n^{7+}$ , H  $g^{2+}$ 

**Answer: A** 

**Solution:** 

**Solution:** 

 $Cu^{2+}$ : [Ar]3d  $^{9}4s^{0}$  $Cr^{3+}$ : [Ar]3d  $^{3}4s^{0}$ Sc<sup>+</sup>: [Ar]3d  $^{1}4s^{1}$ 

All are coloured and paramagnetic due to presence of unpaired electrons

**Question95** 

Number of electrons that Vanadium (Z = 23) has in p-orbitals is equal [22 Jul 2021 Shift 2]

**Answer: 12** 

**Solution:** 

 $_{23}V:1s^22s^22p^63s^23p^63d^34s^2$ 

Number of electrons in p-orbitals is equal to 12.00



Number of electrons present in 4f orbital of H o <sup>3+</sup> ion is (Given Atomic No. of Ho = 67) [25 Jul 2021 Shift 2]	•
[25 Jul 2021 Shift 2]	

**Answer: 10** 

**Solution:** 

#### **Solution:**

```
H o = [X e]4f^{11}6s^2
H o<sup>3+</sup> = [X e]4f^{10}
so number of e<sup>-</sup>present in 4f is 10.
```

\_\_\_\_\_

# **Question97**

The nature of oxides  $V_2O_3$  and CrO is indexed as 'X' and 'Y' type respectively. The correct set of X and Y is [27 Aug 2021 Shift 1]

### **Options:**

A. X = basic Y = amphoteric

B. X = amphoteric, Y = basic

C. X = acidic, Y = acidic

D. X = basic, Y = basic

**Answer: D** 

**Solution:** 

#### **Solution:**

 $V_2O_3$  is a basic oxide as on dissolving in acids, give solution of vanadium (III) complexes. In case of transition metal oxide, the metal with lower oxidation states are basic. So, CrO is also basic as oxidation state of Cr is  $\pm 2$ .

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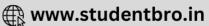
## **Question98**

Which one of the following when dissolved in water gives coloured solution in nitrogen atmosphere? [26 Aug 2021 Shift 1]

#### **Options:**

A. CuCl<sub>2</sub>

B. AgCl



C. ZnCl<sub>2</sub>

D. Cu<sub>2</sub>Cl<sub>2</sub>

**Answer: A** 

## **Solution:**

**Solution:** 

 $CuCl_2$  will dissolve in water to give blue colour  $Cu^{2+}$  ions. The colour is due to the presence of unpaired  $e^-$  in  $Cu^{2+}$  ions.  $CuCl_2 + nH_2O \rightarrow Cu^{2+}(aq)$ 

AgCl and  $Cu_2Cl_2$  are insoluble in water while  $ZnCl_2$  on dissolving in water forms colourless solution due to formation of  $Zn^{2+}$  ions, which has no unpaired  $e^-$ .

------

# **Question99**

# In which one of the following sets all species show disproportionation reaction [31 Aug 2021 Shift 2]

**Options:** 

A.  $\text{ClO}_2^{-}$ ,  $\text{F}_2$ ,  $\text{MnO}_4^{\ 2-}$  and  $\text{Cr}_2\text{O}_7^{\ 2-}$ 

B.  $\operatorname{Cr_2O_7}^{2-}$ ,  $\operatorname{MnO_4}^{-}$ ,  $\operatorname{ClO_2}^{-}$  and  $\operatorname{Cl_2}$ 

C.  $ClO_2^-$ ,  $Cl_2$  and  $Mn^{3+}$ 

D.  $\text{ClO}_4^-$ ,  $\text{MnO}_4^{\ 2-}$ ,  $\text{ClO}_2^-$  and  $\text{F}_2^-$ 

**Answer: B** 

## **Solution:**

#### Solution

Disproportionation reactions are redox reaction in which a compound undergoes oxidation as well as reduction. The element of reacting species is in an intermediate oxidation state and simultaneously gets oxidised and reduced.

$$3\text{ClO}_2^{-3} \rightarrow \text{Cl}^- + 2\text{ClO}_3^{-5}$$

$$0$$

$$\text{Cl}_2 + \text{OH}^- \rightarrow \text{Cl}^- + \text{ClO}_3^- + \text{H}_2\text{O}$$

$$2\text{Mn}^{3+} + 2\text{H}_2\text{O} \rightarrow \text{Mn}^{2+} + \text{MnO}_2 + 4\text{H}^+$$
But  $\text{MnO}_4^-$  does not get disproportonated.
Hence, option (c) is the correct answer.

\_\_\_\_\_

# Question100

Potassium permanganate on heating at 513K gives a product which is [27 Aug 2021 Shift 2]

**Options:** 



- A. paramagnetic and colourless
- B. diamagnetic and green
- C. diamagnetic and colourless
- D. paramagnetic and green

**Answer: A** 

## **Solution:**

#### **Solution:**

When KMnO<sub>4</sub> is heated at 513K, it forms  $K_2MnO_4$  and  $MnO_2$  and  $O_2$ .  $2KMnO_4 \xrightarrow{+6} K_2MnO_4 + MnO_2 + O_2$   $513K \xrightarrow{(Green)} (Black residue)$   $Mn_{25} = 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 4s^2, 3d^5$   $Mn^{+6} = 1s^2, 2s^2, 2p^6, 3s^2, 3p^6, 4s^1; 1$  unpaired electron, so paramagnetic.

So, option (d) is correct.

# Question101

# The addition of dilute NaOH to Cr<sup>3+</sup> salt solution will give [27 Aug 2021 Shift 2]

### **Options:**

- A. a solution of  $[Cr(OH)_4]^-$
- B. precipitate of  $Cr_2O_3(H_2O)_n$
- C. precipitate of  $[Cr(OH)_6]^{3-}$
- D. precipitate of  $Cr(OH)_3$

**Answer: B** 

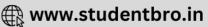
#### **Solution:**

#### Solution:

The addition of dil. NaOH to  ${\rm Cr^{3+}}$  solution will produce ppt. of  ${\rm Cr_2O_3(H_2O)_n}$ .  ${\rm Cr^{3+}}$  + NaOH  $\rightarrow$   ${\rm Cr_2O_3(H_2O)_n}_{\rm (Precipitate)}$  Therefore, the option (b) is correct.

# Question102

The  $Eu^{2+}$  ion is a strong reducing agent in spite of its ground state electronic configuration (outermost): [Atomic number of Eu = 63] [31 Aug 2021 Shift 2]



### **Options:**

A.  $4f^{7}6s^{2}$ 

B. 4f<sup>6</sup>

C. 4f<sup>7</sup>

D.  $4f^{6}6s^{2}$ 

**Answer: C** 

#### **Solution:**

#### Solution:

The electronic configuration of Eu and  $Eu^{2+}$  ion is as follows:

 $Eu(Z = 63) = [Xe]4f^76s^2$  $Eu^{2+}(Z = 63) = [Xe]4f^7$ 

\_\_\_\_\_\_

# Question103

Which one of the following lanthanides exhibits +2 oxidation state with diamagnetic nature ?

(Given, Z for Nd = 60, Yb = 70, La = 57, Ce = 58) [31 Aug 2021 Shift 1]

### **Options:**

A. Nd

B. Yb

C. La

D. Ce

**Answer: B** 

#### **Solution:**

#### Solution:

Ytterbium shows +2 oxidation state and it is diamagnetic in nature.

Yb = [Xe],  $4f^{14}$ ,  $6s^2$ Yb<sup>2+</sup> = [Xe],  $4f^{14}$ Yb<sup>2+</sup> = [Xe], 111111111

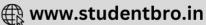
Since, there is no unpaired electron, it is diamagnetic in nature.

\_\_\_\_\_

# Question104

The number of f - electrons in the ground state electronic configuration of Np(Z = 93) is ........





## (Nearest integer) [27 Aug 2021 Shift 1]

**Answer: 18** 

**Solution:** 

#### Solution:

The electronic configuration of Np = 93 is  $1s^22s^22p^63s^23p^63d^{10}4s^24p^64d^{10}5s^25p^64f^{14}5d^{10}6s^26p^65f^46d^{17}s^2$ In ground state, total number of f electrons = 14 + 4 = 18

# Question 105

The number of 4f electrons in the ground state electronic configuration of Gd<sup>2+</sup> is [Atomic number of Gd is 64.] [26 Aug 2021 Shift 1]

**Answer: 7** 

**Solution:** 

#### **Solution:**

The electronic configuration of Gd is as follows  $Gd = [Xe] 4f^7, 5d^1, 6s^2$ So, electronic configuration of Gd<sup>2+</sup> will be  $Gd^{2+} = [Xe], 4f^7, 5d^1, 6s^0$  $\therefore$  The number of 4 f electrons in ground state configuration of  $\mathrm{Gd}^{2+}$  is 7.

# **Question 106**

In the given chemical reaction, colours of the Fe<sup>2+</sup> and Fe<sup>3+</sup> ions, are respectively

 $5Fe^{2+} + MnO_4^- + 8H^+ \longrightarrow Mn^{2+} + 4H_2O + 5Fe^{3+}$ [1 Sep 2021 Shift 2]

**Options:** 

A. yellow, orange

B. yellow, green

C. green, orange



D. green, yellow

**Answer: D** 

#### **Solution:**

The  ${\rm Fe}^{2+}$  ion has green colour while  ${\rm Fe}^{3+}$  is yellow in colour.

\_\_\_\_\_

## Question 107

# The third ionization enthalpy is minimum for: [Jan. 08,2020 (I)]

**Options:** 

A. Co

B. Fe

C. N

D. Mn

**Answer: B** 

#### **Solution:**

#### Solution:

 $_{26}$ F e = [Ar]3d  $^6$ 4s $^2$ . Third ionisation results into stable d  $^5$  configuration.

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# Question108

For the following Assertion and Reason, the correct option is: Assertion: For hydrogenation reactions, the catalytic activity increases from Group 5 to Group 11 metals with maximum activity shown by Group 7 – 9 elements.

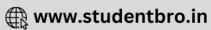
Reason: The reactants are most strongly adsorbed on group 7-9 elements.

[Jan. 08,2020 (II)]

#### **Options:**

- A. The assertion is true, but the reason is false.
- B. Both assertion and reason are false.
- C. Both assertion and reason are true and the reason is the correct explanation for the assertion.
- D. Both assertion and reason are true but the reason is not the correct explanation for the assertion.





**Answer: A** 

**Solution:** 

**Solution:** 

Reactant should not be adsorbed strongly which might result into immobilisation that inhibit further adsorption on the catalyst's surface.

-----

**Question109** 

The atomic radius of Ag is closest to: [Jan. 07,2020 (I)]

**Options:** 

A. Au

B. Ni

C. Cu

D. Hg

Answer: A

**Solution:** 

**Solution:** 

Atomic size of elements of 4d and 5d transition series are nearly same due to lanthanide contraction.

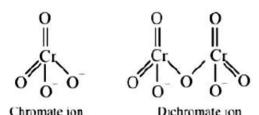
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**Question110** 

The sum of the total number of bonds between chromium and oxygen atoms in chromate and dichromate ions is \_\_\_\_\_. [NV, Jan. 09, 2020 (II)]

**Answer: 12** 

**Solution:** 



⇒ Total number of Cr and O bonds is 12.

Consider the following reactions: NaCl +  $K_2Cr_2O_7$  +  $H_2SO_4$ (Conc.)  $\rightarrow$  (A) + Side products (A) + NaOH  $\rightarrow$  (B) + Side products (B) +  $H_2SO_4$  (dilute) +  $H_2O_2 \rightarrow$  (C) + Side products The sum of the total number of atoms in one molecule each of (A), (B) and (C) is

[NV, Jan. 07, 2020 (II)]

**Answer: 18** 

### **Solution:**

-----

# **Question112**

The electronic configurations of bivalent europium and trivalent cerium are:

(atomic number : X e = 54, Ce = 58, E u = 63 [Jan. 09,2020 (I)]

## **Options:**

A. [X e]4f $^2$  and [X e]4f $^7$ 

B. [X e]4f  $^7$  and [X e]4f  $^1$ 

C. [X e]4f $^7$ 6s $^2$  and [X e]4f $^2$ 6s $^2$ 

D. [X e]4f  $^4$  and [X e]4f  $^9$ 

**Answer: B** 

## **Solution:**

**Solution:** 

 $Eu^{2+}: [Xe]4f^{7}; Ce^{3+}: [Xe]4f^{1}$ 



The highest possible oxidation states of uranium and plutonium, respectively, are:
[Jan. 09,2020 (I)]

### **Options:**

A. 6 and 7

B. 6 and 4

C. 7 and 6

D. 4 and 6

**Answer: A** 

#### **Solution:**

#### **Solution:**

Maximum oxidation state shown by uranium is +6 and plutonium is 7.

-----

# **Question114**

The incorrect statement(s) among (1) - (3) is (are):

- (1) W(VI) is more stable than Cr(VI).
- (2) in the presence of H  $\mbox{Cl}$  , permanganate titrations provide satisfactory results.
- (3) some lanthanoid oxides can be used as phosophorus. [Sep. 4,2020 (II)]

#### **Options:**

- A. (2) and (3) only
- B. (1) and (2) only
- C. (2) only
- D. (1) only

**Answer: C** 

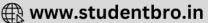
#### **Solution:**

#### Solution:

- (i) W(VI) is more stable than Cr(VI) due to smaller size of atoms and also due to lanthanide contraction.
- (ii) Permanganate titrations in presence of H Cl are unsatisfactory as H Cl is oxidised to Cl ,
- (iii) Lanthanoid oxides are used as phosphors.

\_\_\_\_\_





# The incorrect statement is: [Sep. 03,2020(II)]

### **Options:**

A. Manganate and permanganate ions are tetrahedral

B. In manganate and permanganate ions, the  $\pi$  -bonding takes place by overlap of p -orbitals of oxygen and d – orbitals of manganese

C. Manganate and permanganate ions are paramagnetic

D. Manganate ion is green in colour and permanganate ion is purple in colour

**Answer: C** 

### **Solution:**

#### **Solution:**

Maganate	Permanganate
$MnO_4^{\ 2-}$	$MnO_4^-$
	$O = \bigcup_{Mn^{+7}}^{O} O^{-}$
Paramagnetic, green in colour, Tetrahedral & contains $p\pi - d\pi$ bond	Diamagnetic, purple in colour, Tetrahedral & contains $p\pi - d\pi$ bond

Manganate ion is paramagnetic while permanganate ion is diamagnetic.

\_\_\_\_\_

# **Question116**

# The lanthanoid that does NOT show+ 4 oxidation state is: [Sep. 06,2020(I)]

#### **Options:**

A. Dy

B. Ce

C. Eu

D. Tb

**Answer: C** 



## Mischmetal is an alloy consisting mainly of: [Sep. 06,2020(II)]

#### **Options:**

- A. lanthanoid metals
- B. actinoid and transition metals
- C. lanthanoid and actinoid metals
- D. actinoid metals

**Answer: A** 

### **Solution:**

#### Solution:

Mischmetal is an alloy consisting mainly of lanthanoid metals. Lan. metal  $\Rightarrow \approx 95\%$ 

Iron  $\Rightarrow$  ≈ 5%

S, C, Ca, Al ⇒ traces

## Question118

The correct electronic configuration and spin-only magnetic moment (BM) of Gd $^{3+}$ (Z = 64), respectively, are: [Sep. 05,2020(I)]

#### **Options:**

- A. [Xe]  $4f^7$  and 8.9
- B. [Xe]  $4f^7$  and 7.9
- C.  $[X e]5f^7$  and 8.9
- D. [Xe] 5f and 7.9

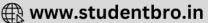
**Answer: B** 

## **Solution:**

#### **Solution:**

Electronic configuration of 64  $Gd = [X e]4f^{7}5d^{1}6s^{2}$ 





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# **Question119**

The pair of metal ions that can give a spin only magnetic moment of 3.9BM for the complex [M (H  $_2$ O) $_6$ ]Cl  $_2$ , is:

[Jan. 12,2019(I)]

### **Options:**

A.  $V^{2+}$  and  $Co^{2+}$ 

B.  $V^2$ + and  $Fe^{2+}$ 

C.  $Co^{2+}$  and  $Fe^{2+}$ 

D.  $Cr^{2+}$  and  $Mn^{2+}$ 

**Answer: A** 

#### **Solution:**

#### **Solution:**

Given  $\mu = 3.9BM$ 

 $\mu = \sqrt{n(n+2)}B \cdot M$ ;  $3.9 = \sqrt{n(n+2)}$ ; n = 3

So, the central metal ion has 3 unpaired electrons.

 $\therefore$  Configuration is either d  $^3$  or d  $^7$ 

As H  $_2$ O is a weak field ligand. V  $^{2+}$  and Co $^{2+}$  will have 3 unpaired electrons. V  $^{2+}$  has d  $^3$  configuration; Co $^{2+}$  has d  $^7$  configuration.

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# Question120

The element that usually does NOT show variable oxidation states is: [Jan. 11, 2019(I)]

#### **Options:**

A. Cu

B. Ti

C. Sc

D. V

**Answer: C** 

## **Solution:**

#### Solution:

Sc shows oxidation state of +3 only.



The highest value of the calculated spin only magnetic moment (in BM) among all the transition metal complexes is:
[[an. 9,2019(I)]

#### **Options:**

A. 5.92

B. 6.93

C. 3.87

D. 4.90

**Answer: A** 

#### **Solution:**

#### **Solution:**

Magnetic moment,  $\mu = \sqrt{n(n+2)}BM$  (where, n = no. of unpaired electrons)

As transition metal atom/ion in a complex may have unpaired electrons ranging from zero to 5 . So, maximum number of unpaired electrons that may be present in a complex is 5.

: Maximum value of magnetic moment among all the transition metal complexes is  $\sqrt{5(5+2)} = \sqrt{35} = 5.92 \, \mathrm{BM}$ 

## **Question122**

# The transition element that has lowest enthalpy of atomisation is: [Jan. 9,2019(II)]

#### **Options:**

A. Fe

B. Cu

C. V

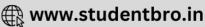
D. Zn

**Answer: D** 

#### **Solution:**

#### Solution:

As zinc has no unpaired of electrons to take part in the bond, it has least enthalpy of atomisation amongst the given transition elements.



# The correct order of atomic radii is : [Jan. 12, 2019 (II)]

**Options:** 

A. N > Ce > Eu > Ho

B. Ho > N > Eu > Ce

C. Ce > Eu > Ho > N

D. Eu > Ce > Ho > N

**Answer: D** 

### **Solution:**

#### **Solution:**

Atomic radii follows the order E u > Ce > H o > N 185pm 182pm 177pm 71pm

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# Question124

The effect of lanthanoid contraction in the lanthanoid series of elements by and large means:
[Jan. 10,2019 (I)]

**Options:** 

- A. increase in both atomic and ionic radii
- B. decrease in atomic radii and increase in ionic radii
- C. decrease in both atomic and ionic radii
- D. increase in atomic radii and decrease in ionic radii

**Answer: C** 

### **Solution:**

#### Solution:

Due to lanthanoid contraction, size of atoms as well as ions of lanthanoid decreases.

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$$\mathbf{A} \xrightarrow{\text{4H Cl}} \mathbf{2B} + \mathbf{2H}_{2}\mathbf{O}$$

$$\mathbf{B} \xrightarrow{\text{4H Cl}} \mathbf{2C} + \mathbf{M}_{2}\mathbf{NO}_{2} + \mathbf{2H}_{2}\mathbf{O}$$

$$\mathbf{2C} \xrightarrow{\text{H}_{2}O, \text{KI}} \mathbf{2A} + \mathbf{K}_{2}\mathbf{O} + \mathbf{D}$$

## In the above sequence of reactions, A and D, respectively, are: [Jan. 11,2019(II)]

**Options:** 

A. K I and K M  $nO_4$ 

B.  $M nO_2$  and  $K I O_3$ 

C. KIO<sub>3</sub> and MnO<sub>2</sub>

D. KI and K<sub>2</sub>M nO<sub>4</sub>

**Answer: B** 

#### **Solution:**

**Solution:** 

Solution:
$$2M \text{ nO}_{2} \xrightarrow{\text{KOH, O}_{2}} 2K_{2}M \text{ nO}_{4} + 2H_{2}O$$

$$(Green) \text{ (B)}$$

$$K_{2}M \text{ nO}_{4} \xrightarrow{\text{4H Cl}} 2K M \text{ nO}_{4} + M \text{ nO}_{2} + 2H_{2}O$$

$$(Purple) \text{ (C)}$$

$$2K M \text{ nO}_{4} \xrightarrow{\text{H}_{2}O, \text{ KI}} 2M \text{ nO}_{2} + 2K \text{ OH} + KIO_{3} \text{ (D)}$$

# **Question126**

## The pair that has similar atomic radii is: [April 12,2019 (II)]

**Options:** 

A. Mn and Re

B. Ti and Hf

C. Sc and Ni

D. Mo and W

**Answer: D** 

#### **Solution:**

#### **Solution:**

Mo and W belong to group-6 and period 5 (4d series) and 6(5d series) respectively. Due to lanthanoid contraction, radius of Mo and W are almost same i.e. 0.140nm and 0.141nm respectively.



Consider the hydrated ions of Ti<sup>2+</sup>, V<sup>2+</sup>, Ti<sup>3+</sup>, and Sc<sup>3+</sup>. The correct order of their spin-only magnetic moments is: [April 12,2019 (II)]

## **Options:**

A. 
$$V^{2+} < Ti^{2+} < Ti^{3+} < Sc^{3+}$$

B. 
$$Sc^{3+} < Ti^{3+} < Ti^{2+} < V^{2+}$$

C. 
$$Ti^{3+} < Ti^{2+} < Sc^{3+} < V^{2+}$$

D. 
$$Sc^{3+} < Ti^{3+} < V^{2+} < Ti^{2+}$$

**Answer: B** 

### **Solution:**

#### **Solution:**

Electronic configuration of the given transition metal ions are :

$$Sc^{3}(Z = 21)l s^{2}2s^{2}2p^{6}3s^{2}3p^{6}$$

$$Sc^{3}(Z = 21)l s^{2}2s^{2}2p^{6}3s^{2}3p^{6}$$
  
 $T i^{2+}(Z = 22)1s^{2}2s^{2}2p^{6}3s^{2}3p^{6}3d^{2}$ 

$$Ti^{3+}(Z = 22)1s^22s^22p^63s^23p^63d^1$$

$$V^{2+}(Z = 23)1s^22s^22p^63s^23p^63d^3$$

Since, magnetic moment is directly proportional to the number of unpaired electrons. The correct increasing order of magnetic moment is

 $Sc^{3+} < Ti^{3+} < Ti^{2+} < V^{2+}$  because they have 0,1,2 and 3 unpaired electrons respectively.

# Question 128

## The correct order of the first ionization enthalpies is: [April 10,2019 (II)]

## **Options:**

A. 
$$Ti < Mn < Zn < Ni$$

B. 
$$Ti < Mn < Ni < Zn$$

C. 
$$M n < T i < Z n < N i$$

**Answer: B** 

#### **Solution:**

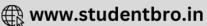
## Solution:

I.E. increases on moving left to right in a period.  $\because \ T\,i < M\,n < N\,i < Z\,n$ 

# **Question129**

## The INCORRECT statement is:





## [April 10,2019 (II)]

### **Options:**

- A. the gemstone, ruby, has  $Cr^{3+}$  ions occupying the octahedral sites of beryl.
- B. the spin-only magnetic moment of  $[N i(N H_3)_4(H_2O)_2]^{2+}$  is 2.83BM
- C. the color of [CoCl (N H  $_3$ ) $_5$ ] $^{2+}$  is violet as it absorbs the yellow light.
- D. the spin-only magnetic moments of  $[Fe(H_2O)_6]^{2+}$  and  $[Cr(H_2O)_6]^{2+}$  are nearly similar.

#### **Answer: A**

#### **Solution:**

#### **Solution:**

Ruby is aluminium oxide (Al  $_2$ O $_3$ ) containing about 0.5-1%Cr $^{3+}$  ions, which are randomly distributed in place of Al  $^{3+}$  ions.

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# Question130

## Match the catalysts (Column I) with products (Column II).

Column I	Column II
Catalyst	Product
V <sub>2</sub> O <sub>5</sub>	Polyethylene
TiCI <sub>4</sub> /AI(Me) <sub>3</sub>	ethanol
PdCI <sub>2</sub>	H <sub>2</sub> SO <sub>4</sub>
Iron Oxide	NH <sub>3</sub>

## [April 9,2019 (I)]

## **Options:**

A. 
$$(A) - (iii)$$
;  $(B) - (iv)$ ;  $(C) - (i)$ ;  $(D) - (ii)$ 

B. 
$$(A) - (ii)$$
;  $(B) - (iii)$ ;  $(C) - (i)$ ;  $(D) - (iv)$ 

C. (A) 
$$-$$
 (iii); (B)  $-$  (i); (C)  $-$  (ii); (D)  $-$  (iv)

D. (A) 
$$-$$
 (iv); (B)  $-$  (iii); (C)  $-$  (ii); (D)  $-$  (i)

**Answer: C** 

# **Solution:**

#### Solution:

(A)V  $_2O_5 \rightarrow$  Preparation of H  $_2SO_4$  in contact process

- (B) TiCl<sub>4</sub> + Al (Me)<sub>3</sub>→ Polyethylene (Ziegler-Natta catalyst)
- (C) Pd Cl<sub>2</sub>→ Ethanol (Wacker's process)





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# **Question131**

# The statement that is INCORRECT about the interstitial compounds is: [April 8,2019 (II)] $\[ \]$

### **Options:**

- A. they are chemically reactive.
- B. they are very hard.
- C. they have metallic conductivity.
- D. they have high melting points.

**Answer: A** 

#### **Solution:**

#### **Solution:**

Interstitial compounds are inert, i.e., they are chemicallynon-reactive.

\_\_\_\_\_

## **Question132**

Thermal decomposition of a Mn compound (X) at 513K results in compound Y, M  $nO_2$  and a gaseous product. M  $nO_2$  reacts with N aCl and concentrated H  $_2SO_4$  to give a pungent gas Z . X, Y, and Z, respectively, are :

[April 12, 2019 (II)]

## **Options:**

- A. K M nO $_4$ , K  $_2$ M nO $_4$  and Cl  $_2$
- B. K  $_2\mathrm{M}~\mathrm{nO}_4$  , K M  $\mathrm{nO}_4$  and  $\mathrm{SO}_2$
- C. K  $_3{\rm M}~{\rm nO}_4$  , K  $_2{\rm M}~{\rm nO}_4$  and Cl  $_2$
- D. K  $_2\mathrm{M}$  nO  $_4$  , K M nO  $_4$  and Cl  $_2$

**Answer: A** 

#### **Solution:**

$$2K \ M \ nO_4 \xrightarrow{513K} K_2 M \ nO_4 + M \ nO_2 + O_2$$

$$M \ nO_2 + 4N \ aCl + 4H_2 SO_4 \rightarrow M \ nCl_2 + 4N \ aH \ SO_4 + Cl_2 + 2H_2 O$$

$$Conc.$$

# The maximum number of possible oxidation states of actinoides are shown by:

[April 9, 2019 (II)]

### **Options:**

- A. Nobelium (No) and lawrencium (Lr)
- B. Actinium (Ac) and thorium (Th)
- C. Berkelium (Bk) and californium (Cf)
- D. Neptunium (Np) and plutonium (Pu)

**Answer: D** 

#### **Solution:**

#### **Solution:**

Actinoids Oxidation state shown Th + 3, +4Ac + 3Pu + 3, +4, +5, +6, +7Np + 3, +4, +5, +6, +7Bk + 3. + 4Cm + 3, +4, +5Ir + 3

 $\therefore$  Maximum oxidation state is shown by N p and Pu.

# Question134

## The lanthanide ion that would show colour is: [April 8, 2019 (II)]

### **Options:**

A. Gd<sup>3</sup>

B. Sm<sup>3</sup>

C. La<sup>3+</sup>

D. Lu<sup>3</sup>

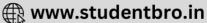
**Answer: B** 

## **Solution:**

#### **Solution:**

 $Sm = 4f^66s^2$  $Sm^{3+} = 4f^5 = Partially filled f orbital$ ∴Sm<sup>3+</sup> will be coloured





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# Question135

When  $XO_2$  is fused with an alkali metal hydroxide in presence of an oxidizing agent such as  $KNO_3$ , a dark green product is formed which disproportionate in acidic solution to afford a dark purple solution. X is:

[Online April 16, 2018]

### **Options:**

- A. Mn
- B. Cr
- C. V
- D. Ti

**Answer: A** 

## **Solution:**

**Solution:** 

$$\begin{array}{l} {\rm M~nO_2+K~OH~+K~N~O_3} \longrightarrow {\rm K~_2~M~nO_4~+K~N~O_2+H~_2O} \\ {\rm M~nO_4^{~2^-}+4H^{~+}} \xrightarrow{\rm dispropor-} {\rm 2~M~nO_4^{~-}+M~nO_2+2H~_2O} \\ \end{array}$$

\_\_\_\_\_

# **Question136**

Which of the following ions does not liberate hydrogen gas on reaction with dilute acids?
[Online April 9, 2017]

## **Options:**

- A.  $Ti^{2+}$
- B. V<sup>2+</sup>
- C. Cr<sup>2+</sup>
- D.  $M n^{2+}$

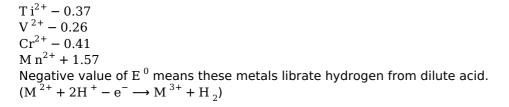
**Answer: D** 

## **Solution:**

#### **Solution:**

Ions  $E^{0}(V)$ 





In the following reactions, Z nO is respectively acting as a/an:

- (i)  $Z nO + N a_2O \rightarrow N a_2Z nO_2$
- (ii)  $Z nO + CO_2 \rightarrow Z nCO_3$ [2017]

**Options:** 

- A. base and acid
- B. base and base
- C. acid and acid
- D. acid and base

**Answer: D** 

**Solution:** 

Solution:

(i) 
$$Z \text{ nO} + N \text{ a}_{2\text{base}} \rightarrow N \text{ a}_{2}Z \text{ nO}_{2\text{salt}}$$

(ii)  $Z \underset{\text{base}}{\text{nO}} + CO_2 \rightarrow Z \underset{\text{salt}}{\text{nCO}_3}$ 

# Question138

The reaction of zinc with dilute and concentrated nitric acid, respectively, produces: [2016]

**Options:** 

- A. N O and N  $_2$ O
- B.  $NO_2$  and  $N_2O$
- C. N $_2$ O and N $O_2$
- D. N $\rm O_2$  and N $\rm O$

Answer: C

Reaction of Z n with different concentration of H N  $O_3$  are as follows: Z n + 4H N  $O_3$ (60%)  $\longrightarrow$  Z n(N  $O_3$ )<sub>2</sub> + 2N  $O_2$  ↑ +2H  $_2$ O 3Z n + 8H N  $O_3$ (30%)  $\longrightarrow$  3Z n(N  $O_3$ )<sub>2</sub> + 2N O ↑ +4H  $_2$ O 4Z n + 10H N  $O_3$ (20%)  $\longrightarrow$  4Z n(N  $O_3$ )<sub>2</sub> + N  $_2$ O ↑ +5H  $_2$ O 5Z n + 12H N  $O_3$ (10%)  $\longrightarrow$  5Z n(N  $O_3$ )<sub>2</sub> + N  $_2$  ↑ +6H  $_2$ O 4Z n + 10H N  $O_3$ (3%)  $\longrightarrow$  4Z n(N  $O_3$ )<sub>2</sub> + N H  $_4$ N  $O_3$  + 3H  $_2$ O Hence option (c) is correct.

-----

# Question139

# Which one of the following species is stable in aqueous solution? [Online April 9,2016]

**Options:** 

A. Cr<sup>2+</sup>

B.  $M nO_4^{2}$ 

C.  $M nO_4^{3}$ 

D. Cu<sup>+</sup>

**Answer: B** 

### **Solution:**

**Solution:** 

I  $nM nO_4^{\ 2-}$  manganese is in +6 oxidation state which is having highest stability.

\_\_\_\_\_

# Question 140

# The transition metal ions responsible for colour in ruby and emerald are, respectively: [Online April 10,2016]

**Options:** 

A.  $Co^{3+}$  and  $Cr^{3+}$ 

B.  $Co^{3+}$  and  $Co^{3+}$ 

C. Cr<sup>3+</sup> and Cr<sup>3+</sup>

D.  $Cr^{3+}$  and  $Co^{3+}$ 

**Answer: C** 



A ruby is a crystal of alumina, aluminum oxide (Al  $_2O_3$ ) containing a trace of chromium (III) ions replacing some of the aluminum ions. In ruby, each Al't ion and  ${\rm Cr}^{3^+}$  ion is surrounded by six oxide ions in an octahedral arrangement. The origin of the color of emeralds is similar to that of the color of rubies. However, the bulk of an emerald crystal is composed of beryl, beryllium aluminum silicate (Be $_3$ Al  $_2$ (SiO $_3$ ) $_6$ ) instead of the alumina which forms rubies. The color is produced by chromium (III) ions, which replace some of the aluminum ions in the crystal. In emeralds, the  ${\rm Cr}^{3^+}$  is surrounded by six silicate ions, rather than the six oxide ions in ruby. Therefore, the color (green) of emeralds is different from that of ruby.

-----

## **Question141**

# Which of the following compounds is metallic and ferromagnetic? [2016]

### **Options:**

 $A. VO_2$ 

B.  $M nO_2$ 

C. TiO<sub>2</sub>

 $D. CrO_2$ 

**Answer: D** 

#### **Solution:**

#### **Solution:**

Out of all the four given metallic oxides,  ${\rm CrO_2}$  is attracted by magnetic field very strongly. The effect persists even when the magnetic field is removed. Thus  ${\rm CrO_2}$  is metallic and ferromagnetic in nature.

\_\_\_\_\_

## Question 142

When concentrated H Cl is added to an aqueous solution of CoCl<sub>2</sub>, its colour changes from reddish pink to deep blue. Which complex ion gives blue colour in this reaction? [Online April 11, 2015]

## **Options:**

A. [CoCl<sub>4</sub>]<sup>2-</sup>

B. [CoCl <sub>6</sub>]<sup>3-</sup>

C. [CoCl <sub>6</sub>]<sup>4</sup>

D.  $[Co(H_2O)_6]^{2+}$ 

**Answer: A** 



Aqueous solution of  $\mathrm{CoCl}_2$  contains  $[\mathrm{Co(H}_2\mathrm{O)}_6]^{2^+}$  which is pinkish in colour so option (d) is incorrect. Reduction potential of  $\mathrm{Co}^{3^+} \to \mathrm{Co}^{2^+}$  is high so option(b) is incorrect.  $\mathrm{Co}^{2^+}$  does not oxidises easily to  $\mathrm{Co}^{3^+}$ . It is general case that symmetrical substituted octahedral complexes are less deeper in colour than tetrahedral complexes. So  $[\mathrm{CoCl}_4]^{2^-}$  is deep blue in colour.

\_\_\_\_\_

## **Question143**

## Which of the following statements is false? [Online April 11, 2015]

#### **Options:**

A. N  $a_2Cr_2O_7$  is less soluble than  $K_2Cr_2O_7$ 

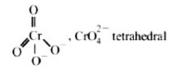
B. N a<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub> is primary standard in volumetry

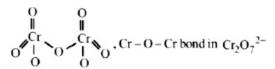
C. CrO<sub>4</sub><sup>2-</sup> is tetrahedral in shape

D.  $CrO_7^{2-}$  has a Cr - O - Cr bond

**Answer: A** 

#### **Solution:**





 $N a_2 Cr_2 O_7$  is used as a primary standard in volumetry.

 $\text{Hydration energy of } N \text{ a}^{\oplus} \text{ is greater than } K \text{ $^{\oplus}$. Because of smaller size of } N \text{ a}^{\oplus}, N \text{ a}_2 Cr_2 O_7 \text{ is more soluble than } K \text{ }_2 Cr_2 O_7 \text{ is more soluble than } K \text{ }_2 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ is more soluble than } K \text{ }_3 Cr_2 O_7 \text{ } \text{ }_3 Cr_2 O_7 \text{ }_3 Cr_2 O_7$ 

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## Question144

Chloro compound of Vanadium has only spin magnetic moment of 1.73BM. This Vanadium chloride has the formula: [Online April 9,2014]

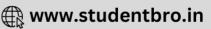
### **Options:**

A. V Cl <sub>2</sub>

B. VCl<sub>4</sub>

C. V Cl  $_3$ 

D. V Cl<sub>5</sub>



**Answer: B** 

#### **Solution:**

#### **Solution:**

Magnetic moment  $= \sqrt{n(n+2)}$  where n = number of unpaired electrons  $\sqrt{n(n+2)} = 1.73$   $\therefore n = 1$ 

Electronic configuration of V is [Ar]4 s<sup>2</sup>3d<sup>3</sup>.

For only one unpaired electron it has to be  $V^{4+}$ .

Hence, the formula of vanadium chloride  $isVCl_4$ .

#### \_\_\_\_\_

## **Question145**

## Which one of the following exhibits the large number of oxidation states?

[Online April 12,2014]

#### **Options:**

A. Ti(22)

B. V (23)

C. Cr(24)

D. M n(25)

**Answer: D** 

#### **Solution:**

#### **Solution:**

Maganese exhibits the large number of oxidation states. The most common oxidation states of M  $\rm n$  are +2,+3,+4,+6 and +7

### \_\_\_\_\_

## **Question146**

## The equation which is balanced and represents the correct product(s) is:

### [2014]

#### **Options:**

A. 
$$\text{Li}_2\text{O} + 2\text{K Cl} \longrightarrow 2\text{LiCl} + \text{K}_2\text{O}$$

B. 
$$[CoCl (N H_3)_5]^+ + 5H^+ \longrightarrow Co^{2+} + 5N H_4^+ + Cl^-$$

C. 
$$[M g(H_2O)_6]^{2+} + (E DT A)^{4-} \xrightarrow{\text{excess NaOH}} [M g(E DT A)]^{2+} + 6H_2O$$

D. 
$$CuSO_4 + 4KCN \rightarrow K_2[Cu(CN)_4] + K_2SO_4$$



**Answer: B** 

#### **Solution:**

#### **Solution:**

The equation in option (b) is correct since both charges as well as atoms are balanced. For the rest,

- (a) Given reaction in unfavourable in the forward direction (  $\rm K_2O$  is unstable, while  $\rm Li_2O$  is stable)
- (c) Given reaction is not balanced w.r.t. charge.
- (d) Given reaction will give  $K_3[Cu(CN)_4]$  as product instead of  $K_2[Cu(CN)_4]$

\_\_\_\_\_

## **Question147**

# Which series of reactions correctly represents chemical reactions related to iron and its compound? [2014]

#### **Options:**

A. 
$$F e^{\frac{d \operatorname{il} H_2 SO_4}{\bullet}} F eSO_4^{\frac{H_2 SO_4}{\bullet}} F e_2(SO_4)_3^{\frac{heat}{\bullet}} F e$$

B. 
$$F e^{\frac{O_2, \text{ heat}}{\bullet}} F eO^{\frac{\text{dil.H}_2SO_4}{\bullet}} F eSO_4^{\frac{\text{heat}}{\bullet}} F e$$

C. 
$$F \stackrel{\text{Cl}_2}{\longrightarrow} F eCl_3 \stackrel{\text{heat, air}}{\longrightarrow} F eCl_2 \stackrel{\text{Z n}}{\longrightarrow} F e$$

D. 
$$F e^{\stackrel{O_2, \text{ heat}}{\longrightarrow}} F e_3 O_4 \xrightarrow{\text{CO, } 600^{\circ}\text{C}} F eO \xrightarrow{\text{CO, } 700^{\circ}\text{C}} F e$$

**Answer: D** 

#### **Solution:**

#### Solution

In equation (a)  $Fe_2(SO_4)_3$ , and in equation (b)  $FeSO_4$  on decomposition will form oxide instead of Fe. In equation (c)  $FeCl_3$  cannot be reduced when heated in air. Hence equation (d ) is correct.

-----

## Question148

Which of the following is not formed when H  $_2$ S reacts with acidic K  $_2$ Cr $_2$ O $_7$  solution? [Online April 9, 2014]

#### **Options:**

B. 
$$Cr_2(SO_4)_3$$

C. K 
$$_2$$
SO $_4$ 



D. S

**Answer: A** 

#### **Solution:**

**Solution:** 

$$\mathrm{K_{2}Cr_{2}O_{7} + 4H_{2}SO_{4} + 3H_{2}S \longrightarrow \mathrm{K_{2}SO_{4} + Cr_{2}(SO_{4})_{3} + 7H_{2}O + 3S}$$

-----

## **Question149**

# Which of the following arrangements does not represent the correct order of the property stated against it? [2013]

#### **Options:**

A.  $V^{2+} < Cr^{2+} < M n^{2+} < F e^{2+}$ : paramagnetic behaviour

B. N  $i^{2+}$  < Co<sup>2+</sup> < F  $e^{2+}$  < M  $n^{2+}$ : ionic size

C.  $Co^{3+} < Fe^{3+} < Cr^{3+} < Sc^{3+}$ : stability in aqueous solution

D. Sc < Ti < Cr < Mn: number of oxidation states

**Answer: A** 

#### **Solution:**

#### **Solution:**

 $V=3d~^34s^2$ ;  $V^{2+}=3d~^3=3$  unpaired electrons  $Cr=3d~^54s^1$ ;  $Cr^{2+}=3d~^4=4$  unpaired electrons  $M~n=3d~^54s^2$ ;  $M~n^{2+}=3d~^5=5$  unpaired electrons  $F~e=3d~^64s^2$ ;  $F~e^{2+}=3d~^6=4$  unpaired electrons Hence the correct order of paramagnetic behaviour  $V^{2+}< Cr^{2+}=F~e^{2+}< M~n^{2+}$ 

(b) For the same oxidation state, the ionic radii generally decreases as the atomic number increases in a particular transition series, hence the order is

 $M n^{2+} > F e^{2+} > Co^{2+} > N i^{2+}$ 

(c) Larger size, least hydrated more stable in aqueous solution. As we move across the period ( $Sc^{3+} \rightarrow Cr^{3+} \rightarrow Fe^{3+} \rightarrow Co^{3+}$ ), the ionic size usually decreases.  $Sc^{3+}$  with the large size as least hydrated and hence more stable. (d) Sc - (+2), (+3)

(d) Sc - (+2), (+3) Ti - (+2), (+3), (+4) Cr - (+2), (+3), (+4), (+5), (+6) Mn - (+2), (+3), (+4), (+5), (+6), (+7) i.e. Sc < Ti < Cr < Mn

## Question150

Four successive members of the first row transition elements are listed below with atomic numbers. Which one of them is expected to have the highest E  $_{M^{\,3+}/\,M^{\,2+}}$  value?

[2013]



#### **Options:**

A. Cr(Z = 24)

B. M n(Z = 25)

C. Fe(Z = 26)

D. Co(Z = 27)

**Answer: D** 

#### **Solution:**

#### **Solution:**

 $E \, ^{\circ}Cr^{3+} / \, Cr^{2+} = -0.41V$   $E_{Fe^{3+}/Fe^{2+}} = +0.77V$   $E \, ^{\circ}M \, n^{3+} / \, M \, n^{2+} = +1.57V$ ,  $E \, ^{\circ}Co^{3+} / \, Co^{2+} = +1.97V$ 

\_\_\_\_\_

## **Question151**

The element with which of the following outer electron configuration may exhibit the largest number of oxidation states in its compounds: [Online April 9,2013]

#### **Options:**

A.  $3d^{5}4s^{2}$ 

B. 3d <sup>8</sup>4s<sup>2</sup>

C.  $3d^{7}4s^{2}$ 

D.  $3d^64s^2$ 

**Answer: A** 

#### **Solution:**

#### **Solution:**

The element with outer electron configuration  $3d^54s^2$  is M n which exhibits oxidation states from +2 to +7.

-----

## Question 152

Potassium dichromate when heated with concentrated sulphuric acid and a soluble chloride, gives brown-red vapours of:
[Online April 9, 2013]

#### **Options:**

A. CrO<sub>3</sub>



B. CrCl <sub>3</sub>

C.  $CrO_2Cl_2$ 

D.  $Cr_2O_3$ 

**Answer: C** 

#### **Solution:**

#### **Solution:**

Solid potassium dichromate when heated with concentrated sulphuric acid and a soluble chloride gives orange red vapours of a volatile oily liquid

 $CrO_2Cl_2 K_2Cr_2O_7 + 4NaCl + 6H_2SO_4 \rightarrow 2KHSO_4 + 4NaHSO_4 + 2CrO_2Cl_2$ 

## Question153

When a small amount of K M  $nO_4$  is added to concentrated H  $_2SO_4$ , a green oily compound is obtained which is highly explosive in nature. Compound may be: [Online April 23, 2013]

#### **Options:**

A. MnSO<sub>4</sub>

B. M  $n_2O_7$ 

C. M  $nO_2$ 

D.  $M n_2 O_3$ 

Answer: B

#### **Solution:**

#### Solution:

K M  $nO_4$  reacts with H  $_2SO_4$  to form M  $n_2O_7$  which is highly explosive substance. 2K M  $nO_4$  + H  $_2SO_4$   $\longrightarrow$  K  $_2SO_4$  + M  $n_2O_7$  + H  $_2O$ 

## Question154

## Identify incorrect statement: [Online April 23, 2013]

#### **Options:**

A. Cu<sub>2</sub>O is colourless.

B. Copper (I) compounds are colourless except when colour results from charge transfer.



C. Copper (I) compounds are diamagnetic.
D. Cu <sub>2</sub> S is black.
Answer: A
Solution:
<b>Solution:</b> $\mathrm{Cu_2O}$ is yellow in colour.
Question155
Iron exhibits +2 and +3 oxidation states. Which of the following statements about iron is incorrect? [2012]
Options:
A. Ferrous oxide is more basic in nature than the ferric oxide.
B. Ferrous compounds are relatively more ionic than the corresponding ferric compounds.
C. Ferrous compounds are less volatile than the corresponding ferric compounds.
D. Ferrous compounds are more easily hydrolysed than the corresponding ferric compounds.
Answer: D
Solution:
<b>Solution:</b> F $e^{3+}$ is easily hydrolysed than F $e^{2+}$ due to more positive charge.
Question156
Which pair of elements with the given atomic numbers is expected to have similar properties? [Online May 19,2012]
Options:
A. 40,72
B. 20,36
C. 10,28
D. 11,12
Answer: A
Solution:

CLICK HERE >>>

Which is not the correct statement? (At. nos. Ce = 58, Lu = 71, La = 57, Yb = 70) [Online May 7,2012]

#### **Options:**

A. Colour of Y b<sup>3+</sup> ion is pink.

B. La<sup>3+</sup> is diamagnetic.

C. Ce<sup>4+</sup> has f <sup>0</sup> configuration.

D.  $Lu^{3+}$  had f  $^{14}$  configuration.

**Answer: A** 

#### **Solution:**

**Solution:** 

Option (a) is incorrect as Y b<sup>3+</sup> is colorless.

## Question158

Magnetic moment of  $Gd^{3+}$  ion (Z = 64) is [Online May 12,2012]

### **Options:**

A. 3.62BM

B. 9.72BM

C. 7.9BM

D. 10.60BM

**Answer: C** 

Gd (64) = [X e]4f 
$$^{7}5d$$
  $^{1}6s^{2}$   
Gd  $^{3+}$  = [X e]4f  $^{7}5d$   $^{0}$  as  $^{0}$   
i.e. no. of unpaired electrons = 7  
 $\mu = \sqrt{n(n+2)} = \sqrt{7(7+2)}$   
=  $\sqrt{63}$  = 7.93BM

Which of the following forms stable +4 oxidation state? [Online May 26,2012]

#### **Options:**

A. La(Z = 57)

B. E u(Z = 63)

C. Ce(Z = 58)

D. Gd(Z = 64)

**Answer: C** 

#### **Solution:**

#### **Solution:**

Only Ce(Z = 58) shows stable (+4) oxidation state.

-----

## Question 160

The number of unpaired electrons in Gadolinium [Z = 64] is [Online May 26,2012]

#### **Options:**

A. 3

B. 8

C. 6

D. 2

**Answer: B** 

#### **Solution:**

#### **Solution:**

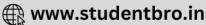
Gd (64) =  $[X e]4f^75d^16s^2$  $\therefore$  No. of unpaired electrons = 8

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## Question161

The correct order of E  $^{\circ}$ M  $^{2+}$  / M  $^{2+}$  values with negative sign for the four successive elements Cr, M n, F e and Co is [2010]

**Options:** 



A.	M n	>	Cr	>	F	е	>	Со

B. 
$$Cr < Fe > Mn > Co$$

D. 
$$Cr > Mn > Fe > Co$$

**Answer: A** 

#### **Solution:**

#### **Solution:**

Across the first transition series, the negative values for standard electrode potential decrease except for M n due to the stable d  $^5$  configuration. Thus, correct order is M  $n^{2+} > Cr^{2+} > F \, e^{2+} > Co^{2+}$ 

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## **Question162**

Knowing that the chemistry of lanthanoids(Ln) is dominated by the state oxidation state, which of the following statements is incorrect? [2009]

#### **Options:**

- A. The ionic size of Ln (III) decreases in general with increasing atomic number
- B. Ln (III) compounds are generally colourless.
- C. Ln (III) hydroxides are mainly basic in character.
- D. Because of the large size of the Ln(III) ions the bonding in its compounds is predominantly ionic in character.

**Answer: B** 

#### **Solution:**

#### **Solution:**

Most of the  $Ln^{3+}$  compounds except  $La^{3+}$  and  $Lu^{3+}$  are coloured due to the presence of unpaired f -electrons.

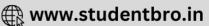
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## Question 163

Amount of oxalic acid present in a solution can be determined by its titration with K M  $nO_4$  solution in the presence of H  $_2SO_4$ . The titration gives unsatisfactory result when carried out in the presence of H Cl , because H Cl [2008]

#### **Options:**

A. gets oxidised by oxalic acid to chlorine



- B. furnishes H + ions in addition to those from oxalic acid
- C. reduces permanganate to M n<sup>2+</sup>
- D. oxidises oxalic acid to carbon doxide and water

**Answer: C** 

#### **Solution:**

#### **Solution:**

The titration of oxalicacid with K M  $nO_4$  in presence of H Cl gives unsatisfactory result because of the fact that K M  $nO_4$  can also oxidise H Cl along with oxalic acid. H Cl on oxidation gives Cl  $_2$  and H Cl reduces K M  $nO_4$  to M  $n^{2+}$ , thus the correct answer is (c).

.....

## Question164

# Larger number of oxidation states are exhibited by the actinoids than those by the lanthanoids, the main reason being [2008]

#### **Options:**

- A. 4f orbitals more diffused than the 5 forbitals
- B. leasser energy difference between 5f and 6d than between 4f and 5d orbitals
- C. more energy difference between 5f and 6d than between 4f and 5d orbitals
- D. more reactive nature of the actionids than the lanthanoids

**Answer: B** 

#### **Solution:**

#### **Solution:**

**Note:** The main reason for exhibiting larger number of oxidation states by actinoids as compared to lanthanoids is lesser energy difference between 5f and 6d orbitals as compared to that between 4f and 5d orbitals. In case of actinoids we can remove electrons from 5f as well as from 6d and due to this actinoids exhibit larger number of oxidation state than lanthanoids.

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### Question 165

## Identify the incorrect statement among the following: [2007]

#### **Options:**

- A. 4f and 5 forbitals are equally shielded.
- B. d -Block elements show irregular and erratic chemical properties among themselves.
- C. La and Lu havepartially filled d -orbitals and no other partially filled orbitals.





D. The chemistry of various lanthanoids is very similar.
Answer: A
Solution:
<b>Solution:</b> $4f$ Orbital is nearer to nucleus as compared to $5f$ orbital therefore, shielding of $4f$ is more than $5f$ .
Question166
The actinoids exhibit more number of oxidation states in general than the lanthanoids. This is because [2007]
Options:
A. the 5f orbitals extend further from the nucleus than the 4f orbitals
B. the 5f orbitals are more buried than the 4f orbitals
C. there is a similarity between 4f and 5 forbitals in their angular part of the wave function
D. the actinoids are more reactive than the lanthanoids.
Answer: A
Solution:
<b>Solution: Note:</b> More the distance between nucleus and outer orbitals, lesser will be force of attraction on them. Distance betwee nucleus and $5f$ orbitals is more as compared to distance between $4f$ orbital and nucleus. So actinoids exhibit more number of oxidation states in general than the lanthanoids.
Question167
The "spin-only" magnetic moment [in units of Bohr magneton, $(\mu_B)$ ] of
N i <sup>2</sup> in aqueous solution would be (At. No. N i = 28 ) [2006]
Options:
A. 6
B. 1.73
C. 2.82
D. 4.90
Answer: C
Solution:

The number of unpaired electrons in N i<sup>2+</sup>(aq) = 2. Water is a weak ligand, hence no pairing will take place  $\therefore$  spin magnetic moment =  $\sqrt{n(n+2)} = \sqrt{2(2+2)} = \sqrt{8} = 2.82$ 

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## **Question168**

## Lanthanoid contraction is caused due to [2006]

#### **Options:**

- A. the same effective nuclear charge from Ce to Lu
- B. the imperfect shielding on outer electrons by 4f electrons from the nuclear charge
- C. the appreciable shielding on outer electrons by 4f electrons from the nuclear charge
- D. the appreciable shielding on outer electrons by 5d electrons from the nuclear charge

**Answer: B** 

#### **Solution:**

#### Solution:

The configuration of lanthanides shows that the additional electron enters the 4f subshell. The shielding of one 4f electron by another is very little or imperfect. The imperfect shielding of f electrons is due to the shape of f orbitals which is very much diffused. Thus as the atomic number increases, the nuclear charge increases by unity at each step, while no comparable increase in the mutual shielding effect of f foccurs. This causes a contraction in the size of the f subshell as a result of which atomic and ionic radii decrease gradually from La to Lu.

\_\_\_\_\_

## Question169

# The oxidation state of chromium in the final product formed by the reaction between K1 and acidified potassium dichromate solution is: [2005]

#### **Options:**

A. +3

B. +2

C. +6

D. +4

**Answer: A** 

#### **Solution:**

$$\text{Cr}_2\text{O}_7^{2-} + 6\text{I}^- + 14\text{H}^+ \longrightarrow 2\text{Cr}^{3+} + 7\text{H}_2\text{O} + 3\text{I}_2$$





The value of the 'spin only' magnetic moment for one of the following configurations is 2.84BM. The correct one is [2005]

#### **Options:**

A. d <sup>5</sup> (in strong ligand field)

B. d<sup>3</sup> (in weak as well as in strong fields)

C. d 4 (in weak ligand fields)

D. d 4 (in strong ligand fields)

**Answer: D** 

#### **Solution:**

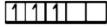
#### **Solution:**

 $d^5 \rightarrow$  strong ligand field



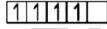
$$\mu = \sqrt{n(n+2)} = \sqrt{1(1+2)} = \sqrt{3} = 1.73BM$$

d3- in weak as well as in strong field



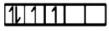
$$\mu = \sqrt{3(3+2)} = \sqrt{15} = 3.87B \cdot M$$

d4- in weak ligand field



$$\mu = \sqrt{4(4+2)} = \sqrt{24} = 4.89$$

d4- in strong ligand field



$$\mu = \sqrt{2(2+2)} = \sqrt{8} = 2.82$$

## Question171

# Which of the following factors may be regarded as the main cause of lanthanide contraction? [2005]

#### **Options:**

- A. Greater shielding of 5d electrons by 4f electrons
- B. Poorer shiel ding of 5d electrons by 4f electrons
- C. Effective shielding of one of the 4f electrons by another in the subshell
- D. Poor shielding of one of the 4f electrons by another in the subshell

**Answer: D** 





#### **Solution:**

#### Solution:

Lanthanide contraction is associated with the intervention of the 4f orbitals which must be filled before the 5d series of elements begin. The filling of 7f before 5d orbitals results in a regular decrease in atomic radii called lanthanoid contraction which essentially compensates for the expected increase in atomic size with increasing atomic number. The factor responsible for the lanthanoid contraction is the imperfect shelding of one electron by another in the same set of orbitals.

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## **Question172**

## The lanthanide contraction is responsible for the fact that [2005]

#### **Options:**

- A. Zr and Zn have the same oxidation state
- B. Zr and Hf have about the same radius
- C. Zr and Nb have similar oxidation state
- D. Zr and Y have about the same radius

**Answer: B** 

#### **Solution:**

#### Solution:

**Note:** In vertical columns of transition elements, there is an increase in size from first member to second member as expected but from second member to third member, there is very small chang in size and some times sizes are same. This is due to lanthanide contraction this is the reason for  $Z\,r$  and  $H\,f$  to have same radius.

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## Question173

## Heating mixture of $Cu_2O$ and $Cu_2S$ will give [2005]

#### **Options:**

- A.  $Cu_2SO_3$
- B. CuO + CuS
- C.  $Cu + SO_3$
- D. Cu +  $SO_2$

Answer: D





## Calomel (H g<sub>2</sub>Cl<sub>2</sub>) on reaction with ammonium hydroxide gives [2005]

**Options:** 

- A. HgO
- B.  $Hg_2O$
- C. N H  $_2$  H g H g Cl
- D. H gN H <sub>2</sub>Cl

**Answer: D** 

**Solution:** 

**Solution:** 

 $H g_2Cl_2 + 2N H_4OH \longrightarrow H g + H gN H_2Cl + N H_4Cl + 2H_2O$ 

## Question 175

### Of the following outer electronic configurations of atoms, the highest oxidation state is achieved by which one of them? [2004]

**Options:** 

- A.  $(n 1)d^{3}ns^{2}$
- B.  $(n 1)d^{5}ns^{1}$
- C.  $(n 1)d^{8}ns^{2}$
- D.  $(n-1)d^{5}ns^{2}$

**Answer: D** 

**Solution:** 

**Solution:** 

 $(n-1)d^{5}ns^{2}$  attains the maximum O.S. of +7

## Question 176



# The correct order of magnetic moments (spin only values in B.M.) among the following is [2004]

**Options:** 

A. 
$$[Fe(CN)_6]^{4-} > [MnCl_4]^{2-} > [CoCl_4]^{2-}$$

B. 
$$[M \, nCl_4]^{2-} > [F \, e(CN)_6]^{4-} > [CoCl_4]^{2-}$$

C. 
$$[M \, nCl_4]^{2-} > [CoCl_4]^{2-} > [F \, e(CN)_6]^{4-}$$

D. 
$$[Fe(CN)_6]^{4-} > [CoCl_4]^{2-} > [MnCl_4]^{2-}$$
 (Atomic nos. : Mn = 25, Fe = 26, Co = 27)

**Answer: C** 

#### **Solution:**

#### **Solution:**

 $[Fe(CN)_6]^4 \rightarrow$ 

No of unpaired electron = 0

 $[MnCl_4]^{2-} \rightarrow$ 

No of unpaired electrons = 5

 $[CoCl_4]^{2-} \rightarrow$ 

No of unpaired electrons = 3

Note : The greater the number of unpaired electrons, greater the magnitude of magnetic moment. Hence the correct order will be  $[M \ nCl_4]^{2^-} > [CoCl_4]^{2^-} > [F \ e(CN)_6]^4$ 

## Question177

# Cerium (Z = 58) is an important member of the lanthanoids. Which of the following statements about cerium is incorrect? [2004]

**Options:** 

- A. The +4 oxidation state of cerium is not known in solutions
- B. The +3 oxidation state of cerium is more stable than the +4 oxidation state
- C. The common oxidation states of cerium are +3 and +4
- D. Cerium (IV) acts as an oxidizing agent

**Answer: A** 



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## **Question178**

Ammonia forms the complex ion  $[Cu(N H_3)_4]^{2+}$  with copper ions in alkaline solutions but not in acidic solutions. What is the reason for it? [2003]

#### **Options:**

- A. In acidic solutions, protons coordinate with ammonia molecules forming N H  $_4^{\,+}$  ions and thus N H  $_3$  molecules are not available
- B. In alkaline solutions insoluble Cu(OH)<sub>2</sub> is precipitated which is soluble in excess of any alkali
- C. Copper hydroxide is an amphoteric substance
- D. In acidic solutions hydration protects copper ions

**Answer: A** 

#### **Solution:**

Solution:

 $\overset{..}{N}$  H  $_3$  + H  $^+$  (acid medium)  $\rightleftharpoons$   $\overset{+}{N}$  H  $_4$ 

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## Question 179

A red solid is insoluble in water. However it becomes soluble if some KI is added to water. Heating the red solid in a test tube results in liberation of some violet coloured fumes and droplets of a metal appear on the cooler parts of the test tube. The red solid is [2003]

#### **Options:**

- A. H  $gL_2$
- B. H gO
- C.  $Pb_3O_4$
- D. (N H<sub>4</sub>)<sub>2</sub>Cr<sub>2</sub>O<sub>7</sub>

**Answer: A** 

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## Question180

What would happen when a solution of potassium chromate is treated with an excess of dilute nitric acid? [2003]

**Options:** 

- A.  $Cr_2O_7^{\ 2-}$  and H  $_2O$  are formed
- B. CrO<sub>4</sub><sup>2-</sup> is reduced to +3 state of Cr
- C. CrO<sub>4</sub><sup>2-</sup> is oxidized to +7 state of Cr
- D.  $Cr^{3+}$  and  $Cr_2O_7^{\ 2-}$  are formed

**Answer: A** 

**Solution:** 

**Solution:** 

 $2 \text{K} \ _2 \text{CrO}_4 + 2 \text{H N O}_3 \longrightarrow \text{K} \ _2 \text{Cr}_2 \text{O}_7 + 2 \text{K N O}_3 + \text{H} \ _2 \text{O}$ 

## Question181

Which one of the following nitrates will leave behind a metal on strong heating?
[2003]

**Options:** 

- A. Copper nitrate
- B. Manganese nitrate
- C. Silver nitrate
- D. Ferric nitrate

**Answer: C** 

**Solution:** 

**Solution:** 

AgN O<sub>3</sub> on heating till red hot decomposes as follows:

$$AgN O_3 \rightarrow Ag + N O_2 + \frac{1}{2}O_2$$

The radius of  $La^{3+}$  (Atomic number of La = 57) is 1.06Å. Which one of the following given values will be closest to the radius of  $\mathrm{Lu}^{3+}$  (Atomic number of Lu = 71)? [2003]

#### **Options:**

A. 1.40Å

B. 1.06Å

C. 0.85Å

D. 1.60Å

**Answer: C** 

#### **Solution:**

#### Solution:

Ionic radii  $\propto \frac{1}{2}$ Thus,  $\frac{z_2}{z_1} = \frac{71}{57} = \frac{1.06}{\text{(lonic radii of } Lu^{3+})}$  $\therefore$  lonic radii of Lu<sup>3+</sup> = 0.85Å

## **Question183**

A reduction in atomic size with increase in atomic number is a characteristic of elements of [2003]

#### **Options:**

A. d -block

B. f -block

C. radioactive series

D. high atomic masses

**Answer: B** 

#### **Solution:**

f -Block elements show a regular decrease in atomic size due to lanthanide\/actinide contraction.

## Question184





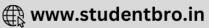


Which of the following ions has the maximum magnetic moment? 2002]
ptions:
. M n <sup>+2</sup>
. F e <sup>+2</sup>
. T i <sup>+2</sup>
. Cr <sup>+2</sup>
nswer: A
olution:
<b>Dlution:</b> $n^{2+} - 5 \text{ unpaired electrons}$ $e^{2+} - 4 \text{ unpaired electrons}$ $e^{2+} - 2 \text{ unpaired electrons}$ $e^{2+} - 4 \text{ unpaired electrons}$ $e^{2+} - 4 \text{ unpaired electrons}$ ote: Magnetic moment $\alpha$ Number of unpaired electrons
Question185
he most stable ion is 2002]
ptions:
. $[Fe(OH)_5]^3$
. $[Fe(Cl)_6]^3$
. $[Fe(CN)_6]^{3-}$
. $[Fe(H_2O)_6]^{3+}$
nswer: C
olution:
<b>Dolution:</b> ne cyano and hydroxo complexes are far more stable than those formed by halide ion. This is due to the fact that CN $^-$ and OH $^-$ are strong Lewis bases (nucleophiles). Further [F e(OH) $_5$ ] $^{3-}$ is not formed. hence most stable ion is

 $[Fe(CN)_6]^{3-}$ 

## **Question186**

When  $\mathbf{K}\,\mathbf{M}\,\mathbf{nO_4}$  acts as an oxidising agent and ultimately forms



[M  $nO_4$ ]<sup>-2</sup>, M  $nO_2$ , M  $n_2O_3$ , M  $n^{+2}$  then the number of electrons transferred in each case respectively is [2002]

**Options:** 

A. 4,3,1,5

B. 1,5,3,7

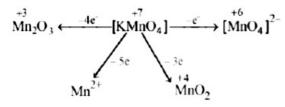
C. 1, 3, 4, 5

D. 3,5,7,1.

**Answer: C** 

**Solution:** 

**Solution:** 



## **Question187**

Most common oxidation states of Ce (cerium) are [2002]

**Options:** 

A. +2,+3

B. +2,+4

C. +3, +4

D. +3, +5.

**Answer: C** 

**Solution:** 

Solution:

Common oxidation states of Ce(cerium) are +3 and +4

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## Question188

Arrange  $Ce^{+3}$ ,  $La^{+3}$ ,  $Pm^{+3}$  and  $Yb^{+3}$  in increasing order of their ionic radii.

#### [2002]

#### **Options:**

A. 
$$Y b^{+3} < Pm^{+3} < Ce^{+3} < La^{+3}$$

B. 
$$Ce^{+3} < Yb^{+3} < Pm^{+3} < La^{+3}$$

$$C. Y b^{+3} < Pm^{+3} < La^{+3} < Ce^{+3}$$

D. 
$$Pm^{+3} < La^{+3} < Ce^{+3} < Yb^{+3}$$
.

**Answer: A** 

#### **Solution:**

In lanthanides there is a regular decrease in the atomic radii as well as ionic radii of trivalent ions as the atomic number increases from Ce to Lu. This decrease in size of atoms and ions is known as **lanthanide contraction**. Although the atomic radii do show some irregularities but ionic radii decrease from La to Lu. Thus the correct order is.

 $Y b^{3+} < Pm^{3+} < Ce^{3+} < La^{3+}$ 

86.8pm 97pm 102pm 103pm

