## JEE Main 2020 Paper

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Subject: Chemistry

1. Determine the wavelength of electron in the 4th Bohr's orbit:
a. $4 \pi a_{0}$
b. $2 \pi a_{0}$
c. $8 \pi \mathrm{a}_{0}$
d. $6 \pi \mathrm{a}_{0}$

Answer: c
Solution: $\mathrm{n}=4$
$\mathrm{Z}=1$
$\Lambda=$ ?


Circumference ( $2 \pi r$ ) $=n \lambda$
$\frac{2 \pi \mathrm{a}_{0} \mathrm{n}^{2}}{\mathrm{z}}=\mathrm{n} \lambda$
On solving, we get $8 \pi a_{0}$
2. Which of the following species have one unpaired electron each?
a. $\mathrm{O}_{2}, \mathrm{O}_{2}^{-}$
b. $\mathrm{O}_{2}, \mathrm{O}_{2}^{+}$
c. $\mathrm{O}_{2}^{+}, \mathrm{O}_{2}^{-}$
d. $\mathrm{O}_{2}, \mathrm{O}_{2}^{2-}$

Answer: c
Solution:
$\mathrm{O}_{2}: \sigma_{1 \mathrm{~s}^{2}} \sigma_{1 \mathrm{~s}^{2}}^{*} \sigma_{2 \mathrm{~s}^{2}} \sigma_{2 \mathrm{~s}^{2}}^{*} \sigma_{2 p_{\mathrm{z}}^{2}} \pi_{2 \mathrm{p}_{\mathrm{x}}^{2}}=\pi_{2 \mathrm{py}^{2}} \pi_{2 p_{\mathrm{x}}^{1}}^{*}=\pi_{2 p_{\mathrm{y}}^{1}}^{*}$
$\mathrm{O}_{2}^{-}: \sigma_{1 \mathrm{~s}^{2}} \sigma_{1 \mathrm{~s}^{2}}^{*} \sigma_{2 \mathrm{~s}^{2}} \sigma_{2 \mathrm{~s}^{2}}^{*} \sigma_{2 \mathrm{p}_{\mathrm{z}}^{2}} \pi_{2 \mathrm{p}_{\mathrm{x}}^{2}}=\pi_{2 \mathrm{py}^{2}} \pi_{2 \mathrm{p}_{\mathrm{x}}^{2}}^{*}=\pi_{2 \mathrm{p}_{\mathrm{y}}^{1}}^{*}$
$\mathrm{O}_{2}^{+}: \sigma_{1 \mathrm{~s}^{2}} \sigma_{1 \mathrm{~s}^{2}}^{*} \sigma_{2 \mathrm{~s}^{2}} \sigma_{2 \mathrm{~s}^{2}}^{*} \sigma_{2 \mathrm{p}_{\mathrm{z}}^{2}} \pi_{2 \mathrm{p}_{\mathrm{x}}^{2}}=\pi_{2 \mathrm{py}^{2}} \pi_{2 \mathrm{p}_{\mathrm{x}}^{1}}^{*}$
3. $\quad$ For $\mathrm{Br}_{2}(\mathrm{l})$, the enthalpy of atomisation $=\mathrm{xkJ} / \mathrm{mol}$ and the bond dissociation enthalpy of bromine $=y \mathrm{~kJ} / \mathrm{mol}$. Then,
a. $x>y$
b. $\mathrm{x}<\mathrm{y}$
c. $x=y$
d. Relation does not exist.

Answer: a
Solution:

$\Delta \mathrm{H}_{\text {atomisation }}=\Delta \mathrm{H}_{\text {vap }}+\mathrm{y}$
$x-y=\Delta H_{v a p}$
4. Which of the following oxides are acidic, basic and amphoteric, respectively?
a. $\mathrm{MgO}, \mathrm{P}_{4} \mathrm{O}_{10}, \mathrm{Al}_{2} \mathrm{O}_{3}$
b. $\mathrm{N}_{2} \mathrm{O}_{3}, \mathrm{Li}_{2} \mathrm{O}, \mathrm{Al}_{2} \mathrm{O}_{3}$
c. $\mathrm{SO}_{3}, \mathrm{Al}_{2} \mathrm{O}_{3}, \mathrm{Na}_{2} \mathrm{O}$
d. $\mathrm{P}_{4} \mathrm{O}_{10}, \mathrm{Al}_{2} \mathrm{O}_{3}, \mathrm{MgO}$

Answer: b
Solution:
Non-metallic oxides are acidic in nature, metallic oxides are basic in nature and $\mathrm{Al}_{2} \mathrm{O}_{3}$ is amphoteric in nature
5. The complex $\mathrm{Cr}\left(\mathrm{H}_{2} \mathrm{O}\right)_{6} \mathrm{Cl}_{\mathrm{n}}$, shows geometrical isomerism and also reacts with $\mathrm{AgNO}_{3}$ solution. Given: The spin only magnetic moment $=3.8 \mathrm{~B}$. M. What is the IUPAC name of the complex?
a. Hexaaquachromium(III) chloride
b. Tetraaquadichloridochromium(III) chloride dihydrate
c. Hexaaquachromium(IV) chloride
d. Tetraaquadichloridochromium(IV) chloride dehydrate

Answer: b
Solution:
Spin only magnetic moment $=3.8$ B. M. This implies, $\mu=\sqrt{ }(n(n+2))$ B.M.
( $\sqrt{16}=4$ implies that $\sqrt{15}$ should be less than four.
This means, $\mathrm{n}=3$ as $\sqrt{15}=\sqrt{ }(3(3+2))$
$\operatorname{Cr}(24)=[\operatorname{Ar}] 4 \mathrm{~s}^{1} 3 \mathrm{~d}^{5}$
(g.s)

For 3 unpaired electrons, the oxidation state of Cr should be +3
$\mathrm{Cr}^{3+}$ can be attained if the complex has a structure that looks like: $\left[\mathrm{Cr}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4} \mathrm{Cl}_{2}\right] \mathrm{Cl}_{2} 2 \mathrm{H}_{2} \mathrm{O}$ $\left[\mathrm{Cr}\left(\mathrm{H}_{2} \mathrm{O}\right)_{4} \mathrm{Cl}_{2}\right] \mathrm{Cl} .2 \mathrm{H}_{2} \mathrm{O}$ has the IUPAC name : Tetraaquadichloridochromium(III) chloride dihydrate
6. The electronic configuration of bivalent Europium and trivalent Cerium, respectively is:
(Atomic Number: $\mathrm{Xe}=54, \mathrm{Ce}=58, \mathrm{Eu}=63$ )
a. $[\mathrm{Xe}] 4 \mathrm{f}^{7},[\mathrm{Xe}] 4 \mathrm{f}^{1}$
b. $[\mathrm{Xe}] 4 \mathrm{f}^{7} 6 \mathrm{~s}^{2},[\mathrm{Xe}] 4 \mathrm{f}^{1}$
c. $[\mathrm{Xe}] 4 \mathrm{f}^{7} 6 \mathrm{~s}^{2},[\mathrm{Xe}] 4 \mathrm{f}^{1} 5 \mathrm{~d}^{1} 6 \mathrm{~s}^{2}$
d. $[\mathrm{Xe}] 4 \mathrm{f}^{7},[\mathrm{Xe}] 4 \mathrm{f}^{1} 5 \mathrm{~d}^{1} 6 \mathrm{~s}^{2}$

Answer: a
Solution:
Ce (58): [Xe] $6 \mathrm{~s}^{2} 4 \mathrm{f}^{2}$
(g.s)
$\mathrm{Ce}^{3+}:[\mathrm{Xe}] 4 \mathrm{f}^{1}$
$\operatorname{Eu}(63):[\mathrm{Xe}] 6 \mathrm{~s}^{2} 4 \mathrm{f}^{7}$
(g.s)
$\mathrm{Eu}^{2+}:[\mathrm{Xe}] 4 \mathrm{f}^{7}$
7. $\mathrm{K}_{\text {sp }}$ of $\mathrm{PbCl}_{2}=1.6 \times 10^{-5}$. On mixing $300 \mathrm{~mL}, 0.134 \mathrm{M} \mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}$ (aq.) and $100 \mathrm{~mL}, 0.4 \mathrm{M}$ $\mathrm{NaCl}(\mathrm{aq}):$.
a. $\quad \mathrm{Q}>\mathrm{K}_{\mathrm{sp}}$
b. $\quad \mathrm{Q}<\mathrm{K}_{\text {sp }}$
c. $\quad \mathrm{Q}=\mathrm{K}_{\mathrm{sp}}$
d. Relation does not exist

Answer: a
Solution: Given $\mathrm{K}_{\text {sp }}$ of $\mathrm{PbCl}_{2}=1.6 \times 10^{-5}$
$\mathrm{Pb}\left(\mathrm{NO}_{3}\right)_{2}:$ mmoles $=300 \mathrm{~mL} \times 0.134 \mathrm{M}=40.2$
$\mathrm{NaCl}:$ mmoles $=100 \mathrm{~mL} \times 0.4 \mathrm{M}=40$
This implies, $[\mathrm{Pb}]^{2+}=\frac{40.2}{400} \approx 0.1 \mathrm{M}$
$[\mathrm{Cl}]^{-}=\frac{40}{400}=0.1 \mathrm{M}$
$\mathrm{Q}_{\mathrm{sp}}=\left[\mathrm{Pb}^{2+}\right]\left[2 \mathrm{Cl}^{-}\right]^{2}=4 \times 10^{-3}>\mathrm{K}_{\mathrm{sp}}$
8. Which of the following cannot act as both oxidising and reducing agent?
a. $\mathrm{H}_{2} \mathrm{SO}_{3}$
b. $\mathrm{HNO}_{2}$
c. $\mathrm{H}_{3} \mathrm{PO}_{4}$
d. $\mathrm{H}_{2} \mathrm{O}_{2}$

Answer: c
Solution:
When the oxidation state is maximum it acts like a strong oxidising agent
When the oxidation state is minimum it acts like a strong reducing agent
When the oxidation state is between its maximum and minimum, it acts like both an oxidizing and as a reducing agent

In $\mathrm{H}_{3} \mathrm{PO}_{4}$, P has a +5 oxidation state and hence can act like a strong oxidising agent. In the rest, the oxidation state is between their maximum and minimum.
9. The first Ionisation energy of Be is higher than that of Boron. Select the correct statements regarding this:
(i) It is easier to extract electron from 2 p orbital than 2 s orbital
(ii) Penetration power of 2 s orbital is greater than 2 p orbital
(iii) Shielding of 2 p electron by 2 s electron
(iv) Radius of Boron atom is larger than that of Be
a. (i), (ii), (iii), (iv)
b. (i), (iii), (iv)
c. (ii), (iii), (iv)
d. (i), (ii), (iii)

Answer: d
Solution:
Be (4): $1 \mathrm{~s}^{2} 2 \mathrm{~s}^{2}$
B (5): $1 \mathrm{~s}^{2} 2 \mathrm{~s}^{2} 2 \mathrm{p}^{1}$
The electron in $2 \mathrm{p}^{1}$ can easily be extracted.
The penetrating power is of the order: $s>p>d>f$
The shielding power order: $s>p>d>f$
As we move along the period, the size decreases, as $\mathrm{Z}_{\text {eff }}$ increases. Hence the radius of B is smaller than the radius of Be.
10. For $[\mathrm{PdFClBrI}]^{2-}$, the number of geometrical Isomers $=n$. Determine the spin only magnetic moment and CFSE for $\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{\mathrm{n}-6}$ (Ignore pairing energy).
a. 1.73 B. M.,$-2 \Delta_{0}$
b. 2.84 B. M.,$-1.6 \Delta_{0}$
c. $0,-1.6 \Delta_{0}$
d. 5.92 B. M.,$-2.4 \Delta_{0}$

Answer: a
Solution:


Number of geometrical isomers ( n ) = 3
$\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{\mathrm{n}-6}=\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{3-6}=\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{-3}$
This implies, that Iron is in its +3 oxidation state.
$\mathrm{Fe}^{3+}(26):[\mathrm{Ar}] 3 \mathrm{~d}^{5}$
$\mathrm{CN}^{-}$is a strong ligand in $\left[\mathrm{Fe}(\mathrm{CN})_{6}\right]^{-3}$ and causes pairing. Hence, according to CFT, the configuration will be $t_{2 g}^{5} e_{g}^{0}$.

Hence, there is only 1 unpaired electron, i.e, $n=1$ in $\sqrt{n(n+2)}=\sqrt{3}=1.73$ B.M

$$
\begin{aligned}
\text { CFSE } & =\left(-0.4 \times \mathrm{n}_{\mathrm{t} 2 \mathrm{~g}}+0.6 \times \mathrm{n}_{\mathrm{eg}}\right) \Delta_{0} \\
& =(-0.4 \times 5+0.6 \times 0) \Delta_{0} \\
& =-2 \Delta_{0}
\end{aligned}
$$

11. A can reduce $\mathrm{BO}_{2}$ under which conditions?

a. $>1400^{\circ} \mathrm{C}$
b. $<1400^{\circ} \mathrm{C}$
c. $>1200{ }^{\circ} \mathrm{C}$
d. $<1200^{\circ} \mathrm{C}$

Answer: a
Solution: In Ellingham's diagram, the line of the element that lies below can reduce the oxide of the element which lies above it. Therefore, for A to reduce $\mathrm{BO}_{2}$, the temperature when the line for element A is below that of $\mathrm{BO}_{2}$, according to the graph when $\mathrm{T}>1400^{\circ} \mathrm{C}$.

For $\mathrm{T}>1400^{\circ} \mathrm{C}, \Delta \mathrm{G}_{\mathrm{r}}<0$ for $\mathrm{A}+\mathrm{BO}_{2} \rightarrow \mathrm{~B}+\mathrm{AO}_{2}$
12. $\mathrm{A} \rightarrow \mathrm{B} ; 700 \mathrm{~K}$

A $\xrightarrow{\mathrm{C}} \mathrm{B} ; 500 \mathrm{~K}$
Rate of the reaction in absence of catalyst at 700 K is same as in presence of catalyst at 500 K . If catalyst decreases the activation energy barrier by $30 \mathrm{~kJ} / \mathrm{mol}$, determine the activation energy in presence of catalyst. (Assume ' A ' factor to be same in both cases)
a. 75 kJ
b. 135 kJ
c. 105 kJ
d. 125 kJ

Answer: c
Solution:
$K=A e^{\left(-\frac{E_{a}}{R T}\right)}$
$\mathrm{K}_{\text {catalyst }}=\mathrm{K}_{\text {without catalyst }}$


Progress of Reaction

$$
A e^{\left(-\frac{\left(\mathrm{E}_{\mathrm{a}}\right) \mathrm{c}}{\mathrm{RT}_{500 \mathrm{~K}}}\right)}=A \mathrm{e}^{\left(-\frac{\left(\mathrm{E}_{\mathrm{a}}\right)}{\mathrm{RT}_{700 \mathrm{k}}}\right)}
$$

$$
\left.e^{\left(-\frac{\left(\mathrm{E}_{\mathrm{a}}\right) \mathrm{c}}{\mathrm{RT}} 500 \mathrm{k}\right.}\right)=\mathrm{e}^{\left(-\frac{\left(\mathrm{E}_{\mathrm{a}}\right)}{\mathrm{RT}} \mathrm{~T}_{700 \mathrm{~K}}\right)}
$$

$$
-\frac{\left(E_{a}\right) c}{\mathrm{RT}_{500 \mathrm{k}}}=-\frac{\left(\mathrm{E}_{\mathrm{a}}\right)}{\mathrm{RT}_{700 \mathrm{k}}}
$$

$$
\left(E_{a}\right) c=E_{a}-30
$$

$$
-\frac{\left(\mathrm{E}_{\mathrm{a}}-30\right)}{\mathrm{T}_{500 \mathrm{k}}}=-\frac{\left(\mathrm{E}_{\mathrm{a}}\right)}{\mathrm{T}_{700 \mathrm{k}}}
$$

On Solving, $\mathrm{E}_{\mathrm{a}}=105 \mathrm{kJmol}^{-1}$
13. A substance ' X ' having low melting point, does not conduct electricity in both solid and liquid state. ' X ' can be :
a. Hg
b. SiC
c. ZnS
d. $\mathrm{CCl}_{4}$

Answer: d
Solution: $\mathrm{CCl}_{4}$ is non polar and does not conduct in either solid or liquid state.
14.


The major product for above sequence of reaction is:
a.


c. $\quad \mathrm{NO}_{2}$
b.

d.


Answer: c
Solution:



Hence, major product formed is that of option b.
15. Which of the following can give the highest yield in Friedel-Craft's reaction?
a.

c.

b.

d.


Answer: b
Solution: Out of the four options given, only aniline and phenol show strong +R effects, but as we know, aniline is a Lewis base and can react with a Lewis acid that is added during the reaction. Hence, Phenol gives the highest yield in Friedel-Craft's reaction.
16.


What will be the major product?
a.

c.

b.

d.


Answer: a
Solution:



17. Which of the following is the correct order for heat of combustion?
(A)

(B)

(C)

a. $\mathrm{C}>\mathrm{B}>\mathrm{A}$
b. $\mathrm{B}>\mathrm{A}>\mathrm{C}$
c. $\mathrm{A}>\mathrm{B}>\mathrm{C}$
d. $C>A>B$

Answer: c
Solution: Heat of combustion $\alpha \frac{1}{\text { stability }}$
The trans-isomer is more stable than the cis-isomer. More the number of trans forms in a structure, higher the stability.

18. Write the correct order of basicity:

(A)

$: \bar{C} \equiv N$ :

$\overline{\mathrm{C}} \equiv \mathrm{CH}$
(B)
(C)
(D)
(E)
a. $\mathrm{A}>\mathrm{B}>\mathrm{D}>\mathrm{E}>\mathrm{C}$
b. $\mathrm{B}>\mathrm{A}>\mathrm{D}>\mathrm{C}>\mathrm{E}$
c. $\mathrm{A}>\mathrm{B}>\mathrm{E}>\mathrm{D}>\mathrm{C}$
c. $\mathrm{C}>\mathrm{E}>\mathrm{D}>\mathrm{B}>\mathrm{A}$

Answer: a
Solution: As we know weaker the conjugate base, stronger the acid.
The order of stability of conjugate base:


Hence, the order of basicity or acidic strength is:
A $>$ B $>$ D $>\mathrm{E}>\mathrm{C}$
19. A, B, C, and D are four artificial sweetners.
(i) $\mathrm{A} \& \mathrm{D}$ give positive test with ninhydrin.
(ii) C form precipitate with AgNO 3 in the lassaigne extract of the sugar.
(iii) $\mathrm{B} \& \mathrm{D}$ give positive test with sodium nitroprusside.

Correct option is :
a. A - Saccharine, B - Aspartame, C - Sucralose, D - Alitame
b. A - Aspartame, B - Saccharine, C - Sucralose, D - Alitame
c. A - Saccharine, B - Aspartame, C - Alitame , D - Sucralose
d. A - Aspartame, B - Sucralose, C - Saccharine, D - Alitame

Answer: b
Solution:


It has a free amine group and hence reacts with ninhydrin to give a purple colour known as Ruhemann's purple.

## B- Saccharine



It has Sulphur, therefore, it will give a positive test with sodium nitroprusside.


It has chlorine and hence it forms a precipitate with $\mathrm{AgNO}_{3}$ in the Lassaigne's extract of the sugar.

D-Alitame


It has a free amine group and hence reacts with ninhydrin to give purple colour known as Ruhemann's purple. Also, it has Sulphur, therefore, it will give positive test with sodium nitroprusside.
20.


Predict the compound (P) on the basis of above sequence of the reactions, where compound (P) gives positive Iodoform test:

b.

a.

c.

d.


Answer: b
Solution:

is a methyl ketone, which gives positive Iodoform test.

21. Given a solution of $\mathrm{HNO}_{3}$ of density $1.4 \mathrm{~g} / \mathrm{mL}$ and $63 \% \frac{\mathrm{w}}{\mathrm{w}}$. Determine molarity of $\mathrm{HNO}_{3}$ solution. Answer: 14.00

Solution: $\% \frac{\mathrm{w}}{\mathrm{w}}=63 \%$

$$
\begin{aligned}
& \rho=1.4 \mathrm{~g} / \mathrm{mL} \\
& M=\frac{\left(\% \frac{\mathrm{w}}{\mathrm{w}} \times \rho \times 10\right)}{\mathrm{MM}} \\
& M=\frac{(63 \times 1.4 \times 10)}{63} \\
& M=14 \mathrm{~mol} / \mathrm{L}
\end{aligned}
$$

22. Determine the degree of hardness in terms of ppm of $\mathrm{CaCO}_{3}$ of $10^{-3}$ molar $\mathrm{MgSO}_{4}(\mathrm{aq})$.

Answer: 100.00
Solution:
Hardness of water is measured in ppm in terms $\mathrm{CaCO}_{3}$.
$\mathrm{n}_{\mathrm{CaCO}_{3}=}=\mathrm{n}_{\mathrm{MgSO}_{4}}$
ppm is the parts (in grams) present per million i.e, $10^{6}$
1000 mL has $10^{-3}$ moles of $\mathrm{MgSO}_{4}$.
Grams of $\mathrm{CaCO}_{3}$ in $1000 \mathrm{~mL}=10^{-3} \times 100$ grams
Grams of $\mathrm{CaCO}_{3}$ in $1 \mathrm{~mL}=\frac{10^{-3} \times 100}{1000 \mathrm{~mL}}$ grams
Hardness $=\frac{10^{-3} \times 100}{1000 \mathrm{~mL}} \times 10^{6}=100$
23. Determine the amount of NaCl to be dissolved in 600 g of $\mathrm{H}_{2} \mathrm{O}$ to decrease the freezing point by $0.2^{\circ}$. Given : $\mathrm{k}_{\mathrm{f}}$ of $\mathrm{H}_{2} \mathrm{O}=2 \mathrm{Km}^{-1}$

Answer: 1.76
Solution: NaCl is strong electrolyte and gives 2 ions in the solution. This implies, $\mathrm{i}=2$.
Molatility $=\frac{\mathrm{w} \times 1000}{58.5 \times 600}$
$\Delta \mathrm{T}_{\mathrm{f}}=0.2^{\circ} \mathrm{C}$
$\Delta \mathrm{T}_{\mathrm{f}}=\mathrm{i} \times \mathrm{k}_{\mathrm{f}} \times \mathrm{m}$
On solving we get,
$\mathrm{w}=1.76 \mathrm{grams}$
24. On passing a particular amount of electricity in $\mathrm{AgNO}_{3}$ solution, 108 g of Ag is deposited. What will be the volume of $\mathrm{O}_{2}(\mathrm{~g})$ in litres liberated at 1 bar, 273 K by the same quantity of electricity?

Answer: 5.68
Solution: On applying Faraday's $1^{\text {st }}$ law,
Moles of Ag deposited= 108/108=1 mol.
$\mathrm{Ag}^{+}+\mathrm{e}^{-} \rightarrow \mathrm{Ag}$
1Faraday is required to deposit 1 mole of Ag .
$\mathrm{H}_{2} \mathrm{O} \rightarrow 2 \mathrm{H}^{+}+\frac{1}{2} \mathrm{O}_{2}+2 \mathrm{e}^{-}$
$\frac{1}{2}$ moles of $\mathrm{O}_{2}$ are deposited by 2 F of charge.
This implies, 1 F will deposit $\frac{1}{4}$ moles of $\mathrm{O}_{2}$
Using $\mathrm{PV}=\mathrm{nRT}$
$\mathrm{P}=1 \mathrm{bar}$
$\mathrm{T}=273 \mathrm{~K}$
$\mathrm{R}=0.0823 \mathrm{Lbar} \mathrm{mol}^{-1} \mathrm{~K}^{-1}$
On solving we get,
$\mathrm{V}=5.68 \mathrm{~L}$
25. Find percentage nitrogen by mass in Histamine?

Answer: 37.84
Solution:
Molecular mass of Histamine $=111$
In Histamine, 3 Nitrogens are present (42g)
The percentage of Nitrogen by mass in Histamine $=\frac{42}{111} \times 100=37.84 \%$


