2020 TJC Promo H2 Chemistry MCQ Worked Solutions

1 Answer: B

Since J has a larger angle of deflection than ¹⁶O²⁻, it would have a greater charge/mass ratio than ¹⁶O²⁻.

2 Answer: C

P+: 1s² 2s² 2p⁶ 3s² 3p² Cu³⁺: 1s² 2s² 2p⁶ 3s² 3p⁶ 3d⁸

Ca: 1s² 2s² 2p⁶ 3s² 3p⁶ 4s²

S: 1s² 2s² 2p⁶ 3s² 3p⁴

3 Answer: A

No. of moles of ethanol molecules = $\frac{23}{24+16+6}$ = 0.500 mol

No. of moles of atoms in g of ammonia = $\frac{2.125}{17}$ x 4 = 0.500 mol

No. of moles of molecules in 11.35 dm³ of sulfur dioxide = $\frac{11.35}{22.7}$ = 0.500 mol

No. of moles of sodium ions = $\frac{250}{1000}$ x 2 x 2 = 1.00 mol

4 Answer: B

	V	С	Н	0
%	19.21	45.3	5.29	30.2
Ar	50.9	12	1	16
Amt	0.377	3.77	5.29	1.88
rati o	1	10	14	5

Empirical formula = VC₁₀H₁₄O₅

$$Z = (5 - 1)/2 = 2$$

5 Answer: A

 $H_2S \rightarrow S + 2H^+ + 2e$ $SO_2 + 4H^+ + 4e \rightarrow S + 2H_2O$ Reacting ratio: $2H_2S \equiv SO_2$

6 Answer: D

Bond angle 1 is around a sp² carbon atom with 3 bp, 0 lp = 120°

Bond angle 2 is around a sp³ carbon atom with 4 bp, 0 lp = 109.5°

Bond angle 3 is around a N-atom with 3 bp, 1 lp = 107°

7 Answer: C

There are no hydrogen bonds existing between methanal molecules. Methanal is a polar hence pd-pd interactions exist between methanol which is stronger than id-id interactions between non polar ethane molecules.

8 Answer: A

 $\Delta H_r = 4(-241.8) - 2(50.6) - (9.2) = -1077.6 \text{ kJ mol}^{-1}$

9 Answer: D

 $\sqrt{\ }$ 2: Combustion reactions are exothermic in nature.

 $\sqrt{4}$: The stoichiometric ratio of sodium oxide is defined as 1 and sodium oxide is formed from its elements.

10 Answer: B

Heat absorbed by water

$$= 80.2 \times \frac{80}{100} = 64.16 \text{ kJ} = 64 160 \text{ J}$$

Heat absorbed by water = $mc\Delta T$ 64 160 = m (4.18)(80 – 25)

Mass of water = 279 g

11 Answer: D

Since K_c remains constant, there is no change in position of equilibrium when temperature changes and the enthalpy change, ΔH is zero.

12 Answer: B

 $K_c = [NH_3]^2[H_2SO_4] = 2.57 \text{ x } 10^{-5}$

 $(2x)^2(x) = 2.57 \times 10^{-5} \text{ hence } x = 0.0186$

13 Answer: B

- $\sqrt{1}$ Vol of O₂ (product) increases from 0 to 50cm³ (½ of max vol) in 20 s Vol of O₂ (product) increases from 50 to 75cm³ (½ of maxi vol to ¾ of max vol) in 20 s half-life is a constant at 20 s.
- $\sqrt{2}$ Since half-life is a constant at 20s, the reaction is first order w.r.t. [H₂O₂].

14 Answer: C

Both k_f and k_b will increase while K_c remains constant as K_c is only temperature dependent.

15 Answer: A

When p increases $\frac{5}{2}$ times, the V should decrease $\frac{5}{2}$ times to 38.4 cm³ if the gas is ideal.

But V of the gas only decreases to 46 cm³ → V_{real} > V_{ideal}

16 Answer: B

 $(0.68)V_T = (0.7)(1.0) + (0.5)(2.5) + (1.0)(1.5)$ $V_T = 5.074 \text{ dm}^3$

Vol of connecting tube = $5.074 - (1.0 + 2.5 + 1.5) = 0.074 \text{ dm}^3$

17 Answer: C

An electrophile is an electron pair acceptor.

They are positively or partial positively charged.

They are attracted to electron rich sites or regions with high electron density.

18 Answer: A

H• radicals are not formed in free radical substitution reactions. Hence H₂ will not be formed.

HC*l* is formed in the propagation step.

 CH_2Cl_2 is formed from further substitution of CH_3Cl .

 CH_3CH_2Cl is formed from $\bullet CH_3$ and $\bullet CH_2Cl$ in a termination step.

19 Answer: D

$$+$$
 Br_2 \longrightarrow Br Or $+$ Br

20 Answer: B

Prismane undergoes free radical substitution with bromine in uv light. 2 hydrogen atoms are substituted with bromine atoms.

There are 3 possible constitutional isomers:

SECTION B (STRUCTURED)

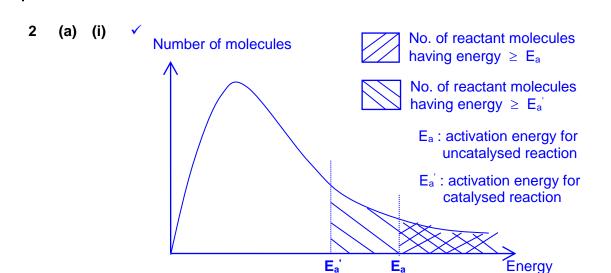
- 1 (a) (i) Comparing expt 1 & 2, when [NO] is constant and [O₂] doubles,

 rate of reaction doubles. The order of reaction wrt O₂ is 1.
 - (ii) Comparing expt 2 & 3, when [NO] doubles and [O₂] constant,

 rate increases by 4 times. The order of reaction wrt NO is 2.
 - (iii) Rate = $k [O_2][NO]^2$
 - (iv) $[NO] = 3.90 \times 10^{-2} \text{ mol dm}^{-3}$
 - (b) (i) $K_c = \frac{[NO]^2}{[NO]^2[O_2]}$ $= \frac{[\frac{0.02}{2}]^2}{[\frac{0.44}{2}]^2[\frac{0.34}{2}]}$ $= 0.0122 \text{ mol}^{-1} \text{ dm}^3$
 - (ii) By Le Chatelier's Principle, the equilibrium position will shift left OR towards the backward reaction in order to produce greater no of moles of gas to increase pressure.

Amount of NO₂ decreases and the amount of NO and O₂ increases.

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A catalyst provides an alternative pathway with lower

activation energy or Ea.

There are more molecules with energy greater than or equal

to the lowered Ea, Frequency of effective collisions increases

Rate constant increases and hence rate of reaction increases.

(ii) Iron is a heterogeneous catalyst that provides a surface for H₂ and

N₂ to be adsorbed. This increases the concentration

of H₂ and N₂ at the catalyst surface. The

intramolecular OR covalent bonds in H₂ and N₂ are weakened.

Product NH₃ desorbed from the iron surface and diffuses away.

- (iii) Porous solid provides a larger surface area for catalysis.
- (b) (i) ICl is polar (has a permanent dipole) hence the electron rich C=C bond attacks the Iδ+ in ICl more readily.

(ii) Elimination Alcoholic KOH, heat

(iii)
HO OH

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3 (a) (i) Element Y is in Group 16.

Large increase in IE when the 7th electron is removed

implies that 7th electron is removed from the inner shell

(ii) I

(iii) J has larger nuclear charge as it has more protons.

Due to its additional electron shell, J has greater screening effect.

The valence electrons of J are further from the nucleus and less

strongly attracted by the nucleus. Less energy is required to

remove the electron of J than that of B.

(b) (i)
$$NO_3^- + Sn^{2+} + 2H^+ \rightarrow NO_2^- + Sn^{4+} + H_2O$$

(ii) No. of moles of
$$S_2O_3^{2-} = 0.12 \times 0.022$$

= 0.00264 mol

No. of moles of
$$Sn^{4+} = 0.00264/2$$

= 0.00132 mol

No. of moles of Sn^{2+} in 50 cm³ solution = 0.00132 × 50/10 = 0.00660 mol

(iii) Percentage purity of tin = $(0.00660 \times 118.7)/2.5 \times 100\% = 31.3\%$

4 (a) (i) Carbon 1: sp

Carbon 2: sp³

(ii) Carbon 1: Linear, 180°

Oxygen: Bent, 104.5°

(b) (i) $C_4H_6O(I) + 5O_2(g) \rightarrow 4CO_2(g) + 3H_2O(I)$

(ii) Energy taken in for bond breaking
=
$$6(410) + 1(350) + 1(840) + 2(360) + 5(496)$$

= 6850 kJ mol^{-1}

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Energy given out for bond forming

= 8(805) + 6(460)

 $= 9200 \text{ kJ mol}^{-1}$

 $\Delta H_c = 6850 - 9200 = -2350 \text{ kJ mol}^{-1}$

(iii) Compound X is a liquid whereas the

bond energies in Data Booklet correspond to breaking covalent

bonds in the gaseous state.

$$\begin{array}{c|c} & -2444 \\ C_4H_6O & (I) + 5O_2 & (g) \rightarrow 4CO_2 & (g) + 3H_2O & (I) \\ \Delta H_{vap} & & & & & & & \\ & & -2350 & & & & \\ C_4H_6O & (g) + 5O_2 & (g) \rightarrow 4CO_2 & (g) + 3H_2O & (g) \end{array}$$

$$\Delta H_{\text{vap}} = (-2444) + 3(+41) - (-2350) = +29.0 \text{ kJmol}^{-1}$$

(c) (i) Compound Y exhibits cis-trans isomerism as there are restricted

rotation about the carbon-carbon double bonds and each carbon of the double bond is bonded to 2 different substituents.

Total number of stereoisomers: 4

$$H$$
 $C=C$
 H
 H_3C
 $C=C$
 H
 $O-CH_3$
cis-trans

$$C = C$$
 $C = C$
 $C =$

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2020 TJC Promo Section C Suggested Answers

Section C Free Response

- 1 (a) $2H_2S + 2O_2 \rightarrow S + SO_2 + 2H_2O$
 - (b) (i) ΔH_c^e , is the heat energy evolved when one mole of the substance is completely burnt in excess oxygen at 298 K and 1 bar.

(ii)
$$\Delta H_c = \frac{(-242)+(-283)}{2} = -262.5 = \frac{-263 \text{ kJ mol}^{-1}}{2}$$

(iii) No. of moles of methane = 1500/890 = 1.69 molVolume of methane = $1.69 \times 24 = 40.6 \text{ dm}^3$

No. of moles of water gas = 1500/262.5 = 5.71 mol Volume of water gas = $5.71 \times 24 = 137$ dm³

(iv) From (b)(iii), a smaller volume of methane (in natural gas) is required to produce the same amount of energy.

Water gas contains highly toxic carbon monoxide gas that binds strongly to haemoglobin in blood. OR

Water gas contains hydrogen gas which is highly explosive. OR Water gas must be synthetically produced (using superheated steam) increases costs and resources needed.

(c) (i)
$$\begin{array}{c} -128 \\ 3/2 \ O_2(g) + CO(g) + 2H_2(g) \rightarrow CH_3OH(I) + 3/2 \ O_2(g) \\ -283 \ +2(-242) \\ CO_2(g) + 2H_2O(I) \end{array}$$

$$\Delta H_c = -639 \ \text{kJ mol}^{-1}$$

- (ii) Both have simple molecular structures. Hydrogen bonding present between methanol molecules is stronger and requires more energy to break than the weak instantaneous dipole-induced dipole interactions in methane molecules. Methanol has a larger electron cloud than methane which is more easily distorted. This causes the id-id forces of attraction to be stronger between methanol molecules. Thus, boiling point of methanol is higher and it exists as a liquid.
- (d) (i) ΔH_n^{θ} , is the enthalpy change when an acid and a base react under infinitely dilute conditions to form one mole of water at 298 K and 1 bar.
 - (ii) Limiting reagent is NaOH(aq).

No. of moles of water produced
$$= \frac{30}{1000} \times 1 = 0.0300 \text{ mol}$$

(iii)
$$\Delta H = -\frac{(30+30)\times4.18\times6.8}{0.03} = -56.8 \text{ kJ mol}^{-1}$$

No. of moles of water produced in second reaction $=\frac{15}{1000} \times 2 = 0.03$ mol

$$\Delta H = -\frac{(15+15)\times4.18\times\Delta T}{0.03} = -56.8 \text{ kJ mol}^{-1}$$

 $\Delta T = 13.6 \text{ °C}$

(iv) No. of moles of remaining $H_2SO_4 = 0.03 - 0.015 = 0.015$ mol

No. of moles of H_2SO_4 in 25 cm³ of diluted solution = 0.015/4 = 0.00375 mol

[KOH] =(0.00375×2) $0.0075 \div (18.60/1000) = 0.403 \text{ mol dm}^{-3}$

2 (a) (i) Electrophilic addition

CH(CH₃)₂
$$\mu$$

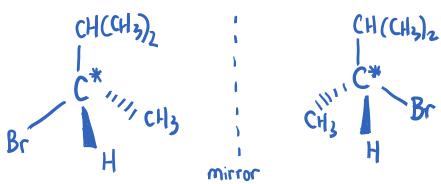
Slow $H - C - C - H + Br$

CH(CH₃)₂ μ

CH(CH₃)₂ μ
 $H - C - C - H + Br$
 $H - C - C$
 $H - C - C$

(ii) In step 2 of the mechanism, there is planar arrangement about the carbon with the positive charge. The carbocation can be attacked by Breither from the top or the bottom of the plane, with equal probability.

This leads to the formation of a pair of enantiomers which are non-superimposable (as there is a chiral centre).



(iii) Optically inactive

CH₃ H H

CH₃ CH

CH₃ H

H

Br

Br

Br

Decolourisation of reddish brown bromine

- (v) The entropy will increase as the gaseous state has more ways to arrange the particles as compared to the liquid state.
- **(b) (i)** Pressure = $1 \times 8.31 \times 298 / 0.25 \times 10^{-3} = 9.91 \times 10^{6} \text{ Pa}$
 - (ii) $p = (1)(8.31)(298)/(0.25 \times 10^{-3} 1 \times 4.28 \times 10^{-5}) (1)^2(0.228)/(0.25 \times 10^{-3})^2$ = 8.30 x 10⁶ Pa
 - (iii) There is significant instantaneous dipole–induced dipole interactions between CH₄ molecules.
- (c) (i) $\Delta H_{soln}^{\theta} = (-390) + (-384) (-779) = +5.00 \text{ kJ mol}^{-1}$
 - (ii) Mg²⁺ has a higher charge and smaller ionic radius OR has a higher charge density as compared to Na⁺, hence a large magnitude of hydration energy.
 - (iii) ΔG_{soln}^{e} (NaCl) = +5 298(43.2/1000) = -7.87 kJ mol⁻¹ <0 Reaction is spontaneous, NaCl is soluble in water

 ΔG_{soln}^{e} (AgC*l*) = +65.7-298(34.3/1000) = +55.5 kJ mol⁻¹>0 Reaction is not spontaneous, AgC*l* is not soluble in water