## DAILY PRACTICE PROBLEM OF PHYSICAL CHEMISTRY FOR NEET

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## THERMOCHEMISTRY



Plot No. 38, Near Union Bank of India, Rajeev Gandhi Nagar, Kota, Rajasthan – 324005 Mob. : 9214233303 DPP-1

| 1.   | If the heat of formation of heat of reaction $2NO_{(g)}$ + | $^{\circ}NO_2$ is 'x' $[^{1}/_2 N_2(g) + O_2(g)$<br>$O_{2(g)} \rightarrow 2NO_{2(g)}$ is z, then | $\rightarrow NO_{2(g)}$ ] the heat of react  | ion $N_{2(g)} + O_{2(g)} \rightarrow 2NO_{2(g)}$ is y and the |
|------|--|--|--|---|
|      | (1)2x+z=y  | (2) 2y + z = x   | (3) 2x - z = y   | (4) 2z + x = y  |
| Ans. | (3)  |  |  |   |
| 2.   | $S (rhombic) + O_2(g) \rightarrow SO_2(g)$                 | $D_2(g); \Delta H = -297.5  kJ$  |  |   |
|      | S (monoclinic) + $O_2(g) \rightarrow$                      | $SO_2(g); \Delta H = -300 \text{ kJ}$  |  |   |
|      | The above data can pred                                    | ict that   |  |   |
|      | (1) Rhombic sulphur is y                                   | ellow in colour.   | (2) Monoclinic sulphur h   | as metallic lustre  |
|      | (3) Monoclinic sulphur                                     | is more stable   | (4) $\Delta H_{(Transition)}$ of S (R) to  | S (M) is endothermic process.                                 |
| Ans. | (4)  |  |  |   |
| 3.   | If $S + O_2 \rightarrow SO_2$ ;                            | $\Delta H = -298.2 \text{ kJ}$   |  |   |
|      | $SO_2 + 1/2 O_2 \rightarrow SO_3;$                         | $\Delta H = -98.7  kJ$   |  |   |
|      | $SO_3 + H_2O \rightarrow H_2SO_4;$                         | $\Delta H = -130.2 \text{ kJ}$   |  |   |
|      | $H_2 + 1/2 O_2 \rightarrow H_2O;$                          | $\Delta H = -287.3 \text{ kJ}$   |  |   |
|      | then the enthalpy of form                                  | ation of $H_2SO_4$ at 298 K is   |  |   |
|      | (1)-814.4 kJ   | (2)-650.3 kJ   | (3)-320.5 kJ   | (4)-433.5 kJ  |
| Ans. | (1)  |  |  |   |
| 4.   | When 1 g of anhydrous or<br>acid : $C_2H_2O_4$ )           | xalic acid is burnt at 25°C, th  | ne amount of heat liberated  | is 2.835 kJ. $\Delta H$ combustion is (oxalic                 |
|      | (1)–255.15 kJ  | (2)-445.65 kJ  | (3)–295.24 kJ  | (4)–155.16 kJ   |
| Ans. | (1)  |  |  |   |
| 5.   | Which one of the following                                 | ng is not applicable for a the   | ermochemical equation :  |   |
|      | (1) It tells about physical                                | state of reactants and prod  | lucts  |   |
|      | (2) It tells whether the re                                | action is spontaneous  |  |   |
|      | (3) It tells whether the rea                               | ction is exothermic or endo  | thermic  |   |
|      | (4) It tells about the allot                               | opic form (if any) of the rea  | octants  |   |
| Ans. | (2)  |  |  |   |
| 6.   | The enthalpy changes of                                    | formation of the gaseous or  | xide of nitrogen $(N_2O \text{ and } P_2O  an$ | NO) are positive because of :                                 |
|      |  | of the nitrogen molecule   | (2) The high electron aff  |   |
|      | (3) The high electron affi                                 | nity of nitrogen atoms   | (4) The tendency of oxyg   | en to form O <sup>2-</sup>                                    |
| Ans. | (1)  |  |  |   |
| 7.   | Heat of formation, $\Delta H_{\rm f}^{\rm o}$              | of an explosive compound l   | ike $NCl_3$ is –   |   |
|      | (1) Positive   | (2) Negative   | (3) Zero   | (4) Positive or negative                                      |
| Ans. | (1)  |  |  |   |
| 8.   | Enthalpy of a compound                                     | is equal to its :- (Who  | en it is formed from constit   | uent particles)   |
|      | (1) Heat of combustion                                     | (2) Heat of formation  | (3) Heat of reaction   | (4) Heat of solution  |
| Ans. | (2)  |  |  |   |

| ).         | The enthalpy of formation of ammonia is $-46.0 \text{ KJ mol}^{-1}$ . The enthalpy change for the reaction $2\text{NH}_3(g) \rightarrow \text{N}_3(g) + 3\text{H}_3(g)$ is:  |  |   |   |  |  |
|------------|--|--|---|---|--|--|
|            | $N_2(g) + 3H_2(g)$ is :<br>(1) 46.0 KJ mol <sup>-1</sup>   | (2) 92.0 KJ mol <sup>-1</sup>  | $(3) - 23.0 \mathrm{KJ}\mathrm{mol}^{-1}$ | $(4) - 92.0 \text{ KJ mol}^{-1}$                          |  |  |
| Ans.       | (1) 40.0 KS mor<br>(2)   | (2) 72.0 KJ mor  | (5) = 25.0 KJ mor                         | (+) = 52.0 K3 mor   |  |  |
| 0.         | Standard enthalpy of   | formation is zero for  |   |   |  |  |
| 0.         |  | (2) $\operatorname{Br}_{(g)}$  | (3) C                                     | (4) O <sub>3(g)</sub>                                     |  |  |
|            | $(1) C_{diamond}$  | $(2) \operatorname{Dr}_{(g)}$  | (3) C <sub>graphite</sub>                 | (+) O <sub>3(g)</sub>                                     |  |  |
| Ans.<br>1. | (3)<br>Civen standard on the   | $1_{\rm max}$ of formation of CO ( $-110$  | V L m a = 1 and $C O (204)$               | KImal-1) The heat of combustion when                      |  |  |
| 1.         | one mole of graphite   |  | <b>KJ</b> more thank $CO_2(-394)$         | KJ mol <sup>-1</sup> ). The heat of combustion when       |  |  |
|            | (1)–110 KJ   | (2) – 284 KJ   | (3) – 394 KJ                              | (4) – 504 KJ  |  |  |
| .ns.       | (3)  |  |   |   |  |  |
| 2.         |  | luring the combustion of I   | 12 litre of water gas at STF              | $P(\text{mixture of equal volume of H}_2 \text{ and CO})$ |  |  |
|            | $H_2(g) + \frac{1}{2} O_2(g) = H_2$  | $O(g); \Delta H = -241.8 \text{ KJ}$   |   |   |  |  |
|            | $CO(g) + \frac{1}{2} O_2(g) = CO(g)$   |  |   |   |  |  |
|            | (1)241.8 KJ  | (2) 283 KJ   | (3) 1312 KJ                               | (4) 1586 KJ   |  |  |
| ns.        | (3)  |  |   |   |  |  |
| 3.         |  |  |   |   |  |  |
|            | (1) 23.5 K cals  | (2) 2.35 K cals  | (3) 94.0 K cals                           | (4) 31.3 K cals   |  |  |
| ns.        | (1)  |  |   |   |  |  |
| 4.         | Given $C(s) + O_2(g)$ —  | $\rightarrow CO_2(g) + 94.2 \text{ Kcal}$  |   |   |  |  |
|            | $H_2(g) + \frac{1}{2} O_2(g) \longrightarrow H_2O(\ell) + 68.3 \text{ Kcal}$   |  |   |   |  |  |
|            | 2 2  | $CO_{2}(g)+2H_{2}O(\ell)+210.8$ Kcal   |   |   |  |  |
|            | 4 2  | of methane in Kcal will be   |   |   |  |  |
|            | (1)-45.9   | (2)-47.8   | (3)-20.0                                  | (4)-47.3  |  |  |
| ns.        | (3)  |  | (0) 2000                                  |   |  |  |
| 5.         |  | HCl(g); $\Delta$ H <sup>0</sup> = −44 Kcal<br>$\Rightarrow$ 2NaCl(s) + H <sub>2</sub> (g); |   |   |  |  |
|            | Then, $Na(s) + 0.5 Cl_2$   | $g) \longrightarrow \text{NaCl}(s); \Delta H^0 = ?$  |   |   |  |  |
|            | (1) 108 Kcal   | (2) 196 Kcal   | (3)-98 Kcal                               | (4) 54 Kcal   |  |  |
| ns.        | (3)  |  |   |   |  |  |
| 6.         | Given that :   |  |   |   |  |  |
|            | $Zn + \frac{1}{2} O_2 \rightarrow ZnO + 84000 cal$ 1   |  |   |   |  |  |
|            | $Hg + \frac{1}{2}$ $O_2 \rightarrow HgO + 21700 cal$ 2   |  |   |   |  |  |
|            | The heat of reaction ( $\Delta$ H) for,  |  |   |   |  |  |
|            | $Zn + HgO \rightarrow ZnO + ZnO +$ | Hg is :-   |   |   |  |  |
|            | (1) 105700 cal   | (2) 62300 cal  | (3)-105700 cal                            | (4) - 62300 cal   |  |  |
| ns.        | (4)  |  |   |   |  |  |
|            |  | or   |   |   |  |  |

|      | $AO + \frac{1}{2} O_2 \rightarrow AO_2$ is 1   | 00 Kcal. The heat of reaction            | on for $A + O_2 \rightarrow AO_2$ is:- |                 |  |  |  |  |
|------|--|--|--|-----------------|--|--|--|--|
|      | (1) - 50 K cal.  | (2) + 50 K cal.                          | (3) 100 K cal.                         | (4) 150 K cal.  |  |  |  |  |
| Ans. | (2)  |  |  |                 |  |  |  |  |
| 18.  | Using the following therr  | nochemical data:                         |  |                 |  |  |  |  |
|      | $C(S) + O_2(g) \rightarrow CO_2(g),$   | $\Delta H = -94.0$ Kcal                  |  |                 |  |  |  |  |
|      | $\mathrm{H_2(g)} + 1/2\mathrm{O_2(g)} \rightarrow \mathrm{H_2C}$   | $O(\ell), \Delta H = -68.0 \text{ Kcal}$ |  |                 |  |  |  |  |
|      | CH <sub>3</sub> COOH ( $\ell$ ) + 2O <sub>2</sub> (g) → 2CO <sub>2</sub> (g) + 2H <sub>2</sub> O( $\ell$ ), $\Delta$ H = - 210.0 Kcal<br>The heat of formation of acetic acid is:- |  |  |                 |  |  |  |  |
|      |  |  |  |                 |  |  |  |  |
|      | (1)116.0 Kcal  | (2) - 116.0 Kcal                         | (3) - 114.0 Kcal                       | (4)+114.0 K cal |  |  |  |  |
| Ans. | (3)  |  |  |                 |  |  |  |  |

| 1.   | Consider the following r<br>C (graphite) + $O_2(g) \rightarrow$   |  |   |  |  |  |
|------|---|--|---|--|--|--|
|      | $C (diamond) + O_2 (g) \rightarrow CO_2(g);  \Delta H = -x, cal$  |  |   |  |  |  |
|      | -   | sition of graphite into diam           | nond?                                     |  |  |  |
|      | $(1) x_1 + x_2$   | $(2) x_2 - x_1$                        | $(3) x_1 - x_2$                           | $(4) x_1 x_2$                          |  |  |
| Ans. | (2)   |  |   |  |  |  |
| 2.   | The heat of combustion of yellow P and red P are –9.91 kJK and –8.78 kJ respectively. The heat of transition of yellow to red phosphorus is |  |   |  |  |  |
|      | (1)–18.69 kJ  | (2)+1.13 kJ                            | (3)+18.69 kJ                              | (4)–1.13 kJ                            |  |  |
| Ans. | (4)   |  |   |  |  |  |
| 3.   | $\Delta H_{f} C_{2} H_{4} = 12.5 \text{ kcal}$  |  |   |  |  |  |
|      | Heat of atomisation of C  |  |   |  |  |  |
|      | Bond energy of $H_2 = 104$  |  |   |  |  |  |
|      | Bond energy of $C - H =$  |  |   |  |  |  |
|      | What is $C = C$ bond energy   | rgy?                                   |   |  |  |  |
|      | (1) 140.9 kcal  | (2) 49 kcal                            | (3) 40 kcal                               | (4) 76 kcal                            |  |  |
| Ans. | (1)   |  |   |  |  |  |
| 4.   | $H_2(g) + \frac{1}{2}O_2(g) \longrightarrow H_2(g)$   | D ( <i>l</i> )                         |   |  |  |  |
|      | $H_2O_{(l)} \rightarrow H_2O_{(g)}; \Delta H = x$   | 4                                      |   |  |  |  |
|      | Given,  |  |   |  |  |  |
|      | $E_{H-H} = x_1$   |  |   |  |  |  |
|      | $E_{0=0} = x_2$   |  |   |  |  |  |
|      | $E_{0-H} = x_3$   |  |   |  |  |  |
|      | $\Delta H_{\rm F}$ of H <sub>2</sub> O vapour is  |  |   |  |  |  |
|      |   |  |   |  |  |  |
|      | (1) $x_1 + \frac{x_2}{2} - x_3 + x_4$   | (2) $2x_3 - x_1 - \frac{x_2}{2} - x_4$ | $(3) x_1 + \frac{x_2}{2} - 2x_3 - x_4$    | $(4) x_1 + \frac{x_2}{2} - 2x_3 + x_4$ |  |  |
| Ans. | (4)   |  |   |  |  |  |
| 5.   | $H(g) + O(g) \rightarrow O - H(g);$   | $\Delta H$ for this reaction is        |   |  |  |  |
|      | (1) Heat of formation of $O - H$  |  | (2) Bond energy of $O - H$                |  |  |  |
|      | (3) Heat of combustion of H,  |  | (4) Zero at all temperatures              |  |  |  |
| Ans. | (2)   | 2                                      | ( ) _ · · · · · · · · · · · · · · · · · · |  |  |  |
| 6.   |   | nciate 4 g of gaseous H in             | to free gaseous atoms is 8                | 72 kJ at 25°C. The bond energy of      |  |  |
| 0.   | H–H bond will be  | -                                      |   |  |  |  |
|      | (1) 8.72 kJ   | (2) 4.36 kJ                            | (3) 436 kJ                                | (4) 43.6 kJ                            |  |  |
| Ans. | (3)   |  |   |  |  |  |
| 7.   | The enthalpy of reaction  | ,                                      |   |  |  |  |
|      | $2HC \equiv CH + 5O_2 \rightarrow 4$  | $CO_2 + 2H_2O$                         |   |  |  |  |
|      | If the bond energies of C–H, C=C, O=O, C=O and O–H bonds are p, q, r, s, t respectively.  |  |   |  |  |  |
|      | (1)[8s+4t]-[4p+q+5r]  |  | (2) $[4p+2q+5r]-[8s+4]$                   | 4t]                                    |  |  |
|      | (3) [4p + 2q + 5r + 8s + 4t]  | ]                                      | (4) [2p+q+5r] - [8s+4t]                   | ]                                      |  |  |
| Ans. | -(2)  | Diat Na 20 No Uniter P                 | uk of India. Raieev Gandhi                | Nagar Page#5                           |  |  |

| 8.   | Using bond energy data   | , calculate heat of formation            | on of isoprene                              |   |
|------|--|--|---|---|
|      | $5C(s) + 4H_2(g) \longrightarrow 1$  | $H_2C = C - CH = CH_2$                   |   |   |
|      |  | CH <sub>3</sub>                          |   |   |
|      | Given C–H, H–H, C–C,   | $C = C$ and $C(s) \rightarrow C(g)$ resp | ectively as 98.8 kcal, 104 kca              | l, 83 kcal, 147 kcal, 171 kcal          |
|      | (1)–21 kcal  | (2)21 kcal                               | (3) 40 kcal                                 | (4) 50 kcal                             |
| Ans. | (2)  |  |   |   |
| 9.   | Entropy of vaporisation  | of water at 100°C, if molar              | heat of vaporisation is 9710                | cal mol <sup>-1</sup> will be           |
|      | $(1) 20 \text{ cal mol}^{-1} \text{K}^{-1}$  | (2) 26.0 cal mol <sup>-1</sup> $K^{-1}$  | $(3)24 \text{ cal mol}^{-1} \text{ K}^{-1}$ | (4) 28.0 cal mol <sup>-1</sup> $K^{-1}$ |
| Ans. | (2)  |  |   |   |
| 10.  | If $\Delta H_{f}^{o}$ of $ICl_{(g)}$ , $Cl_{(g)}$ , ar   | nd $I_{(g)}$ is 17.57, 121.34 and 10     | 6.96 J mol <sup>-1</sup> respectively. Th   | en bond dissociation energy of ICl bond |
|      | is -   |  |   |   |
|      | (1) 35.15 J mol <sup>-1</sup>  | (2) 106.69 J mol <sup>-1</sup>           | $(3) 210.73 \text{ J mol}^{-1}$             | (4) 420.9 J mol <sup>-1</sup>           |
| Ans. | (3)  |  |   |   |
| 11.  | The enthalpy of vapour   | isation of liquid water using            | g the data:                                 |   |
|      | $\mathrm{H}_{2}(\mathrm{g}) + 1/2 \mathrm{O}_{2}(\mathrm{g}) \longrightarrow \mathrm{H}_{2}\mathrm{O}$             | $(\ell); \Delta H = -285.77 K Jmol^{-1}$ |   |   |
|      | $\mathrm{H}_{2}(\mathrm{g}) + 1/2 \mathrm{O}_{2}(\mathrm{g}) \longrightarrow \mathrm{H}_{2}\mathrm{O}(\mathrm{g})$ | g);∆H=-241.84KJmol <sup>-1</sup>         |   |   |
|      | (1)+43.93 KJ mol <sup>-1</sup>   | (2) - 43.93 KJ mol <sup>-1</sup>         | $(3) + 527.61 \text{ KJ mol}^{-1}$          | (4) - 527.61 KJ mol <sup>-1</sup>       |
| Ans. | (1)  |  |   |   |
|      |  |  |   |   |
|      |  |  |   |   |
|      |  |  |   |   |

## **DPP - 3**

|   |   | 1  | /11-5   |   |  |  |  |
|---|---|--|---|---|--|--|--|
| 1.  | In the reactions  |  |   |   |  |  |  |
|   | $HCl + NaOH \rightarrow NaCl + H_2O + x cal$  |  |   |   |  |  |  |
|   | $H_2SO_4 + 2NaOH \rightarrow Na_2$  | $_{2}$ SO <sub>4</sub> + 2H <sub>2</sub> O + y cal     |   |   |  |  |  |
|   | (1)x = y  | (2) x = 2y   | $(3) x = \frac{y}{2}$   | (4) $x = \sqrt{y}$                            |  |  |  |
| Ans.  | (3)   |  |   |   |  |  |  |
| 2.  | The difference between  | $\Delta H$ and $\Delta E$ for the reaction             | $2C_6H_6(l) + 15O_2(g) \rightarrow 12$                                  | $CO_2(g) + 6H_2O(l)$ at 25°C in kJ is         |  |  |  |
|   | (1)-7.43 kJ   | (2)+3.72  kJ   | (3)-3.72 kJ   | (4)+7.43 kJ                                   |  |  |  |
| Ans.  | (1)   |  |   |   |  |  |  |
| 3.  | The heat of neutralization  | on of LiOH and HCl at 25°                              | C is 34.868 kJ mol <sup>-1</sup> . The he                               | eat of ionisation of LiOH will be             |  |  |  |
|   | (1)44.674 kJ  | (2) 22.232 kJ  | (3) 32.684 kJ   | (4) 96.464 kJ                                 |  |  |  |
| Ans.  | (2)   |  |   |   |  |  |  |
| 4.  | -   | absorb the maximum amou<br>C in kcal/mol of each solut |   | in the same amount of water ? (Integral       |  |  |  |
|   | $(1)$ HCl $(\Delta H = -17.74)$   |  | (2) $\text{HNO}_{3}(\Delta \text{H} = -7.85)$                           | )   |  |  |  |
|   | $(3) \text{NH}_4 \text{NO}_3 (\Delta \text{H} = +16.0)$   | )8)  | (4) NaCl ( $\Delta$ H = +1.02)  |   |  |  |  |
| Ans.  |   |  |   |   |  |  |  |
| 5. $HA + OH^- \rightarrow H_2O + A^- + q_1kJ$ |   |  |   |   |  |  |  |
|   | $H^+ + OH^- \rightarrow H_2O + q_2 kJ$  |  |   |   |  |  |  |
|   | The enthalpy of ionisation of HA is   |  |   |   |  |  |  |
|   | $(1)(q_1 + q_2)$  | $(2)(q_1 - q_2)$                                       | $(3)(q_2 - q_1)$  | $(4) - (q_1 + q_2)$                           |  |  |  |
| Ans.  | (3)   |  |   | · · · · · · · · · · · · · · · · · · ·         |  |  |  |
| 6.  | For strong acid strong base neutralisation energy for 1 mole $H_2O$ formation is -57.1 kJ. If 0.25 mole of strong monop acid is reacted with 0.5 mole of strong base then enthalpy of neutralisation of |  |   |   |  |  |  |
|   | (1) $-(0.25 \times 57.1)$   | $(2) 0.5 \times 57.1$                                  |   | -(0.5 × 57.1)                                 |  |  |  |
| Ans.  | (1)   |  |   |   |  |  |  |
| 7.  |   | n of solid benzoic acid at                             | constant volume is -321.3   | kJ at 27°C. The heat of combustion at         |  |  |  |
|   | The heat of combustion of solid benzoic acid at constant volume is -321.3 kJ at 27°C. The heat of combustion at constant pressure is  |  |   |   |  |  |  |
|   | (1)-321.3-300R  | (2) - 321.30 + 300R                                    | (3)-321.3-150R  | (4) - 321.3 + 900R                            |  |  |  |
| Ans.  | (3)   |  |   |   |  |  |  |
| 8.  | The standard entropies of $N_2(g)$ , $H_2(g)$ and $NH_3(g)$ are 191.5, 130.5, 192.6 JK <sup>-1</sup> mol <sup>-1</sup> . The value of $\Delta S^\circ$ of formation ammonia is                          |  |   |   |  |  |  |
|   | $(1)-98.9 \text{ JK}^{-1} \text{ mol}^{-1}$   | (2) Zero   | $(3) + 129.4  \text{JK}^{-1}  \text{mol}^{-1}$                          | $(4) - 29.4 \text{ JK}^{-1} \text{ mol}^{-1}$ |  |  |  |
| Ans.  | (1)   |  |   |   |  |  |  |
| 9.  | Given $S^{o}_{C_{2}H_{6}} = 225 \text{ J mol}^{-1} \text{ K}^{-1}$  |  |   |   |  |  |  |
|   | $S^{o}_{C_{2}H_{4}} = 220 \text{ J mol}^{-1} \text{ K}^{-1}, \ S^{o}_{H_{2}} = 130 \text{ J mol}^{-1} \text{ K}^{-1}$   |  |   |   |  |  |  |
|   | $S_{C_2H_4} = 220$ J mor K  | 2  | Then $\Delta S^{\circ}$ for the process<br>C H + H $\rightarrow$ C H is |   |  |  |  |
|   | Then $\Delta S^{\circ}$ for the proce   |  |   |   |  |  |  |
|   |   |  | (3) 135 J   | (4) 315 J                                     |  |  |  |

| 0.   | For the melting of NaCl heat required is 7.26 kcal mol <sup>-1</sup> and $\Delta S$ increases by 6.73 cal mol <sup>-1</sup> k <sup>-1</sup> . The melting point of the salt is |   |  |  |   |  |  |
|------|--|---|--|--|---|--|--|
|      |  |   | (2) 500 K  | (3) 1.77 K                                   | (4) 1.77°C  |  |  |
| Ans. | (1)  |   |  |  |   |  |  |
| 1.   |  | The $\Delta S$ for the reaction<br>$2H_2(g) + O_2(g) \rightarrow 2H_2O(1)$ at 300 K when            |  |  |   |  |  |
|      | $S_{H_2}^{o}(g) = 126.6$ , $S_{O_2}^{o}(g) = 201.20$ , $S_{H_2O}^{o}(l) = 68.0 \text{ JK}^{-1}\text{mol}^{-1}$ respectively is   |   |  |  |   |  |  |
|      | (1)-3  | 18.4 JK <sup>-1</sup> mol <sup>-1</sup>   | (2) 318.4 JK <sup>-1</sup> mol <sup>-1</sup>                       | (3) 31.84 JK <sup>-1</sup> mol <sup>-1</sup> | $(4) 3.184  \mathrm{JK}^{-1} \mathrm{mol}^{-1}$       |  |  |
| Ans. | (1)  |   |  |  |   |  |  |
| 2.   | Whic   | h of the following  | is correct?  |  |   |  |  |
|      |  | $\Delta H$  | $\Delta S$   | Nature of reaction                           |   |  |  |
|      | (1)  | (-)   | (+)  | Spontaneous only at high temp                | perature  |  |  |
|      | (2)  | (+)   | (-)  | Nonspontaneous regardless of                 | temperature   |  |  |
|      | (3)  | (+)   | (+)  | Spontaneous only at low temp                 | erature   |  |  |
|      | (4)  | (-)   | (-)  | Spontaneous at all temperature               | es  |  |  |
| Ans. | (2)  |   |  |  |   |  |  |
| 3.   | A particular reaction at 27°C for which $\Delta H > 0$ and $\Delta S > 0$ is found to be non-spontaneous. The reaction may proceed spontaneously if                            |   |  |  |   |  |  |
|      | (1) The temperature is decreased   |   | (2) The temperature is   | sincreased                                   |   |  |  |
|      | (3) The temperature is kept constant   |   | (4) It is carried in open vessel at 27°C                           |  |   |  |  |
| Ans. | (2)  |   |  |  |   |  |  |
| 4.   | It is ir   | npossible for a rea   | action to take place if  |  |   |  |  |
|      | (1) $\Delta H$ is +ve and $\Delta S$ is +ve  |   | (2) $\Delta H$ is –ve and $\Delta S$                               |  |   |  |  |
|      | (3) $\Delta$ H is +ve and $\Delta$ S is –ve  |   |  | (4) $\Delta H$ is –ve and $\Delta S$         | is –ve  |  |  |
| Ans. | (3)  |   |  |  |   |  |  |
| 5.   | The sole criterion for the spontaneity of a process is   |   |  |  |   |  |  |
|      | (1) Tendency to acquire minimum energy   |   |  |  |   |  |  |
|      | (2) Tendency to acquire maximum randomness   |   |  |  |   |  |  |
|      | (3) Tendency to acquire minimum energy and maximum randomness  |   |  |  |   |  |  |
|      | (4) Tendency to acquire maximum stability  |   |  |  |   |  |  |
| Ans. | (4)  |   |  |  |   |  |  |
| 6.   |  | If water is formed from H <sup>+</sup> ions and OH <sup>-</sup> the heat of formation of water is : |  |  |   |  |  |
|      | (1) - 1  | .3.7 Kcal   | (2)13.7 KCal   | (3)–63.4 Kcal                                | (4) More data required                                |  |  |
| Ans. | (1)  |   |  |  |   |  |  |
| 7.   |  |   | anhydrous CuSO <sub>4</sub> and<br>f anhydrous CuSO <sub>4</sub> ? |  | .80 Kcal mol <sup>-1</sup> respectively. What will be |  |  |
|      | (1)-1  | 8.69 KCal   | (2) 18.69 Kcal   | (3)-28.96 Kcal                               | (4) 28.96 Kcal  |  |  |
|      | (1)  |   |  |  |   |  |  |