## Yakeen NEET 2.0 2026

## Physical Chemistry by Amit Mahajan Sir Thermodynamics & Thermochemistry

DPP: 6

- **Q1**  $0.16 \, \mathrm{g}$  of methane was subjected to combustion at  $27^{\circ}\mathrm{C}$  in a bomb calorimeter system. The temperature of the calorimeter system (including water) was found to rise by  $0.5^{\circ}\mathrm{C}$ . Heat of combustion of methane at constant pressure is(Heat capacity of the calorimeter system is  $17.7 \mathrm{~kJ~K^{-1}}$  ).
  - (A) 890 kJ
  - (B) -885 kJ
  - (C) + 890 kJ
  - (D) +885 kJ
- Q2 Standard enthalpy of formation is zero for
  - (A)  $C_{diamond}$
  - (B) Br(g)
  - (C)  $C_{
    m graphite}$
  - (D)  $O_{3(g)}$
- $\mathbf{Q3}$  18 g of water is taken to prepare the tea. Find out the internal energy of vaporization at  $100^{\circ} C (\Delta_{vap} H \text{ for water at } 373 \text{ K})$  $40.66 \text{ kJ mol}^{-1}$ 
  - (A)  $37.56 \text{ kJ mol}^{-1}$
  - (B)  $-37.56 \text{ kJ mol}^{-1}$
  - (C)  $43.73 \text{ kJ mol}^{-1}$
  - (D)  $-43.76 \text{ kJ mol}^{-1}$
- **Q4** When 0.5~g of sulphur is burnt to  $SO_2, 4.6~kJ$ of heat is liberated. What is the enthalpy of formation of Sulphur dioxide.
  - (A) +147.2 kJ
  - (B) -147 kJ
  - (C) -294.4 kJ

- (D) +294.4 kJ
- Q5 The enthalpy change for the reaction,  $\mathrm{H}_2\mathrm{O}(\mathrm{s}) o \mathrm{H}_2\mathrm{O}(\ell)$  is called
  - (A) Enthalpy of formation
  - (B) Enthalpy of fusion
  - (C) Enthalpy of vaporisation
  - (D) Enthalpy of transition
- **Q6** The  $\Delta H^{\circ}$  for the reaction.  $4 \text{ S(s)} + 6 \text{O}_2(\text{g}) \rightarrow 4 \text{SO}_3(\text{g}) \text{ is - } 1583.2 \text{ kJ}.$ Standard enthalpy of formation of sulphur trioxide is:
  - (A) -3166.4 kJ
  - (B) 3166.4 kJ
  - (C) -395.8 kJ
  - (D) 395.8 kJ
- dissociation enthalpy Q7 Bond is used to defining enthalpy change of a reaction as
  - (A)  $\Delta H_r = \Sigma (Bond dissociation enthalpy)_{Reactant}$ Σ(Bond dissociation enthalpy)<sub>Product</sub>
  - (B)  $\Delta H_r = \Sigma (Bond dissociation enthalpy)_{Product}$  $-\Sigma$ (Bond dissociation enthalpy)<sub>Reactant</sub>
  - (C)  $\Delta H_r = \Sigma (Bond dissociation enthalpy)_{Product}$ +  $\Sigma$ (Bond dissociation enthalpy)<sub>Reactant</sub>
  - (D) None of these
- **Q8** The heat released in neutralization of HCl and NaOH is 13.7kcal/mol, the heat released on neutralization of NaOH with  $CH_3COOH$  is 3.7 kcal/mol. The  $\Delta H^{\circ}$  of ionization of CH<sub>3</sub>OOOH is
  - (A) 10.2kcal

- (B) 10kcal
- (C) 3.7kcal
- (D) 9.5kcal
- Q9 Heat of neutralization of strong acid by a strong base is equal to  $\Delta H$  of
  - (A)  $\mathrm{H}^+ + \mathrm{OH}^- \longrightarrow \mathrm{H}_2\mathrm{O}$
  - (B)  $H_2O + H^+ \longrightarrow H_3O^+$
  - (C)  $2H_2+O_2 \longrightarrow 2H_2O$
  - (D)  $NH_4OH + HCl \longrightarrow NH_4Cl + H_2O$
- Q10 The Enthalpy of neutralization of acetic acid and sodium hydroxide is  $-55.4 \mathrm{\ kJ}$ . What is the enthalpy of ionisation of acetic acid?
  - (A) -5.54 kJ
  - (B) +5.54 kJ
  - (C) +1.9 kJ
  - (D) -1.9 kJ
- Q11 Which of the following acid has the lowest value (magnitude) of heat neutralization?
  - (A)  $CH_3COOH$
  - (B) HCl
  - (C) HBr
  - (D) HI
- Q12 The enthalpy of neutralization of any strong acid and strong base is nearly equal to
  - (A) +57.3 kJ/mol
  - (B) -75.3 kJ/mol
  - (C) +75.3 kJ/mol
  - (D) -57.3 kJ/eq

<b>Answer Ke</b>	y
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Q1	(A)	<b>Q7</b>	(A)
Q2	(C)	Q8	(B)
Q3	(A)	Q7 Q8 Q9 Q10 Q11	(A)
Q4	(C)	Q10	(C)
Q5	(B)	Q11	(A)
Q6	(C)	Q12	(D)

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