



# Arjuna NEET (2024)

## Thermodynamics

**DPP-07**

1. Since the enthalpy of the elements in their standard states is taken to be zero. The heat of formation ( $\Delta H_f$ ) of compounds:
- (1) Is always negative
  - (2) Is always positive
  - (3) Is zero
  - (4) May be positive or negative
2. Which of the following equations represents standard heat of formation of  $\text{CH}_4$ ?
- (1)  $\text{C}_{(\text{diamond})} + 2\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$
  - (2)  $\text{C}_{(\text{graphite})} + 2\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$
  - (3)  $\text{C}_{(\text{diamond})} + 4\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$
  - (4)  $\text{C}_{(\text{graphite})} + 4\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$
3. Given enthalpy of formation of  $\text{CO}_2(\text{g})$  and  $\text{CaO}(\text{s})$  are  $-94.0 \text{ KJ}$  and  $-152 \text{ KJ}$  respectively and the enthalpy of the reaction:  
 $\text{CaCO}_3(\text{s}) \rightarrow \text{CaO}(\text{s}) + \text{CO}_2(\text{g})$  is  $42 \text{ KJ}$ .  
The enthalpy of formation of  $\text{CaCO}_3(\text{s})$  is
- (1)  $-42 \text{ KJ}$
  - (2)  $-202 \text{ KJ}$
  - (3)  $+202 \text{ KJ}$
  - (4)  $-288 \text{ KJ}$
4. The enthalpies of combustion of carbon and carbon monoxide are  $-393.5 \text{ KJ}$  and  $-283 \text{ KJ}$ , respectively the enthalpy of formation of carbon monoxide is:
- (1)  $-676.5 \text{ KJ}$
  - (2)  $-110.5 \text{ KJ}$
  - (3)  $110.5 \text{ KJ}$
  - (4)  $676.5 \text{ KJ}$
5. The heat of combustion of  $\text{CH}_4(\text{g})$ ,  $\text{C}(\text{s})$  and  $\text{H}_2(\text{g})$  at  $25^\circ\text{C}$  are  $-212.4 \text{ Kcal}$ ,  $-94.0 \text{ Kcal}$  and  $-68.4 \text{ Kcal}$  respectively, the heat of formation of  $\text{CH}_4$  will be-
- (1)  $+54.4 \text{ Kcal}$
  - (2)  $-18.4 \text{ Kcal}$
  - (3)  $-375.2 \text{ Kcal}$
  - (4)  $+212.8 \text{ Kcal}$
6. Standard enthalpy of formation is zero for.
- (1)  $\text{C}_{\text{diamond}}$
  - (2)  $\text{Br}_{(\text{g})}$
  - (3)  $\text{C}_{\text{graphite}}$
  - (4)  $\text{O}_{3(\text{g})}$
7. Heat of formation of  $\text{CO}_2$  is  $-94.0 \text{ Kcal}$ . What would be the quantity of heat liberated, when  $3 \text{ g}$  of graphite is burnt in excess of oxygen:-
- (1)  $23.5 \text{ Kcals}$
  - (2)  $2.35 \text{ Kcals}$
  - (3)  $94.0 \text{ Kcals}$
  - (4)  $31.3 \text{ Kcals}$
8. If  $\text{S} + \text{O}_2 \rightarrow \text{SO}_2$ ;  $\Delta H = -298.2 \text{ kJ mol}^{-1}$   
 $\text{SO}_2 + \frac{1}{2} \text{O}_2 \rightarrow \text{SO}_3$ ;  $\Delta H = -98.7 \text{ kJ mol}^{-1}$   
 $\text{SO}_3 + \text{H}_2\text{O} \rightarrow \text{H}_2\text{SO}_4$ ;  $\Delta H = -130.2 \text{ kJ mol}^{-1}$   
 $\text{H}_2 + \frac{1}{2} \text{O}_2 \rightarrow \text{H}_2\text{O}$ ;  $\Delta H = -287.3 \text{ kJ mol}^{-1}$   
Then the enthalpy of formation of  $\text{H}_2\text{SO}_4$  at  $298 \text{ K}$  will be-
- (1)  $-814.4 \text{ kJ mol}^{-1}$
  - (2)  $-650.3 \text{ kJ mol}^{-1}$
  - (3)  $-320.5 \text{ kJ mol}^{-1}$
  - (4)  $-433.5 \text{ kJ mol}^{-1}$
9. The heat of reaction for  $\text{A} + \frac{1}{2} \text{O}_2 \rightarrow \text{AO}$  is  $-50 \text{ kcal/mol}$  and  $\text{AO} + \frac{1}{2} \text{O}_2 \rightarrow \text{AO}_2$  is  $100 \text{ kcal/mol}$ . The heat of reaction (in  $\text{kcal/mol}$ ) for  $\text{A} + \text{O}_2 \rightarrow \text{AO}_2$  will be:
- (1)  $-50$
  - (2)  $+50$
  - (3)  $100$
  - (4)  $150$
10. Which of the reaction defines molar  $\Delta H_f^\circ$ ?
- (1)  $\text{CaO}(\text{s}) + \text{CO}_2(\text{g}) \rightarrow \text{CaCO}_3(\text{s})$
  - (2)  $\frac{1}{2} \text{Br}_2(\text{g}) + \frac{1}{2} \text{H}_2(\text{g}) \rightarrow \text{HBr}(\text{g})$
  - (3)  $\text{N}_2(\text{g}) + 2\text{H}_2(\text{g}) + \frac{3}{2} \text{O}_2(\text{g}) \rightarrow \text{NH}_4\text{NO}_3(\text{s})$
  - (4)  $\text{I}_2(\text{g}) + \text{H}_2(\text{g}) \rightarrow 2\text{HI}(\text{g})$

- 11.** The enthalpy of formation of ammonia is  $-46.0 \text{ kJ mol}^{-1}$ . The enthalpy change for following reaction is  $2\text{NH}_3 \rightarrow \text{N}_2 + 3\text{H}_2$
- $42.0 \text{ kJ mol}^{-1}$
  - $64.0 \text{ kJ mol}^{-1}$
  - $86.0 \text{ kJ mol}^{-1}$
  - $92.0 \text{ kJ mol}^{-1}$
- 12.** The molar heat of combustion of  $\text{C}_6\text{H}_6$  is  $-3250 \text{ kJ mol}^{-1}$ . When  $0.39 \text{ g}$  of  $\text{C}_6\text{H}_6$  is burnt in excess of oxygen in an open vessel, the amount of heat evolved is:
- $8.32 \text{ kJ}$
  - $12.36 \text{ kJ}$
  - $16.25 \text{ kJ}$
  - $20.74 \text{ kJ}$
- 13.** Heat of combustion  $\Delta H^\circ$  for  $\text{C}(\text{s})$ ,  $\text{H}_2(\text{g})$  and  $\text{CH}_4(\text{g})$  are  $-94$ ,  $-68$  and  $-213 \text{ kcal/mol}$ . Then,  $\Delta H^\circ$  for  $\text{C}(\text{s}) + 2\text{H}_2(\text{g}) \rightarrow \text{CH}_4(\text{g})$  is
- $-17 \text{ kcal/mol}$
  - $-111 \text{ kcal/mol}$
  - $-170 \text{ kcal/mol}$
  - $-85 \text{ kcal/mol}$
- 14.** The enthalpy and entropy change for the reaction,  $\text{Br}_2(\text{l}) + \text{Cl}_2(\text{g}) \rightarrow 2\text{BrCl}(\text{g})$  are  $30 \text{ kJ mol}^{-1}$  and  $105 \text{ JK}^{-1} \text{ mol}^{-1}$  respectively. The temperature at which the reaction will be in equilibrium is:
- $285.7 \text{ K}$
  - $273 \text{ K}$
  - $450 \text{ K}$
  - $300 \text{ K}$
- 15.** Assume each reaction is carried in open container. For which reaction  $\Delta H > \Delta E$ ?
- $\text{H}_2(\text{g}) + \text{I}_2(\text{g}) \rightarrow 2\text{HI}(\text{g})$
  - $\text{N}_2(\text{g}) + 3\text{H}_2(\text{g}) \rightarrow 2\text{NH}_3(\text{g})$
  - $\text{PCl}_5(\text{g}) \rightarrow \text{PCl}_3(\text{g}) + \text{Cl}_2(\text{g})$
  - $\text{SO}_2(\text{g}) + 1/2\text{O}_2(\text{g}) \rightarrow \text{SO}_3(\text{g})$
- 16.** The following reactions are given :
- $$\text{CH}_4(\text{g}) + 2\text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}) + 2\text{H}_2\text{O}(\text{g}); \Delta H = -x \dots \text{(i)}$$
- $$\text{C}(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}); \Delta H = -y \dots \text{(ii)}$$
- $$\text{H}_2(\text{g}) + 1/2\text{O}_2(\text{g}) \rightarrow \text{H}_2\text{O}(\text{g}); \Delta H = -z \dots \text{(iii)}$$
- Calculate the heat of formation of  $\text{CH}_4$  ?
- $x + y + z$
  - $y + 2z - x$
  - $x - y - 2z$
  - none of the above
- 17.**  $\Delta U^\circ$  of combustion of methane is  $-X \text{ kJ mol}^{-1}$ . The value of  $\Delta H^\circ$  is:
- $-X - 596R$
  - $-X + 596R$
  - $X + 596R$
  - $X - 596R$
- 18.** Consider the following process:
- $$\text{A} \rightarrow 2\text{B}; \Delta H = +150 \text{ kJ}$$
- $$3\text{B} \rightarrow 2\text{C} + \text{D}; \Delta H = -125 \text{ kJ}$$
- $$\text{G} + \text{A} \rightarrow 2\text{D}; \Delta H = +350 \text{ kJ}$$
- For  $\text{B} + \text{D} \rightarrow \text{G} + 2\text{C}$ ;  $\Delta H$  will be
- $+525 \text{ kJ}$
  - $+325 \text{ kJ}$
  - $-175 \text{ kJ}$
  - $-325 \text{ kJ}$
- 19.** The standard enthalpies of formation of  $\text{CO}$ ,  $\text{NO}_2$  and  $\text{CO}_2$  are  $-110.5 \text{ kJ mol}^{-1}$ ,  $-33.2 \text{ kJ mol}^{-1}$  and  $-393.5 \text{ kJ mol}^{-1}$  respectively. The standard enthalpy of the reaction is:
- $$4\text{CO}(\text{g}) + 2\text{NO}_2(\text{g}) \rightarrow 4\text{CO}_2(\text{g}) + \text{N}_2(\text{g})$$
- $-1065.6 \text{ kJ}$
  - $-200 \text{ kJ}$
  - $-700 \text{ kJ}$
  - $850 \text{ kJ}$
- 20.** Given  $\text{C}(\text{s}) + \text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}); \Delta H = -94.2 \text{ kcal}$   
 $\text{H}_2(\text{g}) + 1/2\text{O}_2(\text{g}) \rightarrow \text{H}_2\text{O}(\text{g}); \Delta H = -68.3 \text{ kcal}$   
 $\text{CH}_4(\text{g}) + 2\text{O}_2(\text{g}) \rightarrow \text{CO}_2(\text{g}) + 2\text{H}_2\text{O}(\text{g}); \Delta H = -210.8 \text{ kcal}$   
 What will be heat of formation of  $\text{CH}_4$  in (kcal)?
- $+45.9$
  - $-20.0$
  - $+47.8$
  - $-47.3$
- 21.** From the given information, what is the standard enthalpy of formation for  $\text{Al}_2\text{O}_3(\text{s})$ ?
- $$2\text{Al}_2\text{O}_3(\text{s}) \rightarrow 4\text{Al}(\text{s}) + 3\text{O}_2(\text{g}); \Delta H^\circ_{\text{rxn}} = 3352 \text{ kJ}$$
- $-6704 \text{ kJ/mol}$
  - $-3352 \text{ kJ/mol}$
  - $-1676 \text{ kJ/mol}$
  - $1676 \text{ kJ/mol}$



22. Consider the following reactions  
 $C(s) + O_2(g) \rightarrow CO_2(g) + X \text{ kJ}$   
 $CO(g) + 1/2O_2(g) \rightarrow CO_2(g) + Y \text{ kJ}$   
The heat of formation of  $CO(g)$  is
- (1)  $-(X + Y) \text{ kJ/mol}$
  - (2)  $(X - Y) \text{ kJ/mol}$
  - (3)  $(Y - X) \text{ kJ/mol}$
  - (4)  $(Y + X) \text{ kJ/mol}$
23. The enthalpy change for the following reaction is 368 kJ. Calculate the average O–F bond energy  
 $OF_2(g) \rightarrow O(g) + 2F(g)$
- (1) 184 kJ/mol
  - (2) 368 kJ/mol
  - (3) 536 kJ/mol
  - (4) 736 kJ/mol
24.  $N_2(g) + 2O_2(g) \longrightarrow 2NO_2(g) + xkJ$   
 $2NO(g) + O_2(g) \longrightarrow 2NO_2(g) + ykJ$   
The enthalpy of formation of  $NO$  is
- (1)  $x - 2y$
  - (2)  $x - y$
  - (3)  $\frac{1}{2}(y - x)$
  - (4)  $\frac{1}{2}(x - y)$



**Note: Kindly find the Video Solution of DPPs Questions in the DPPs Section.**

## Answer Key

1. (4)
2. (2)
3. (4)
4. (2)
5. (2)
6. (3)
7. (1)
8. (1)
9. (2)
10. (3)
11. (4)
12. (3)

13. (1)
14. (1)
15. (3)
16. (3)
17. (1)
18. (4)
19. (1)
20. (2)
21. (3)
22. (3)
23. (1)
24. (3)



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