

Thermodynamics DPP-6 Solutions

Topic: Adiabatic Process

Vikrant Sir

Q1. When a gas undergoes adiabatic expansion, it gets cooled. This is because

- (1) It is an exothermic process
- (2) It is an endothermic process
- (3) Internal energy of the system decreases
- (4) Ideal gas becomes a real gas

Conceptual Approach: Adiabatic process means $q = 0$. Cooling ya heating depend karega internal energy par. Expansion ke time gas(system) ka work negative hota hai, aur kyunki $q = 0$, isliye ΔU negative hota hai \rightarrow internal energy kam hoti hai \rightarrow temperature girta hai.

Step-by-step Solution:

$$\Delta U = q + w, \quad q = 0 \implies \Delta U = w$$

Expansion $\implies w < 0 \implies \Delta U < 0 \implies T$ decreases.

Final Answer: Correct option: (3) Internal energy of the system decreases.

Q2. Which of the following is correct expression for 1st law of thermodynamics under adiabatic condition?

- (1) $\Delta U = Q + W$
- (2) $\Delta U = Q - W$
- (3) $Q = -W$
- (4) $\Delta U = W$

Conceptual Approach: Adiabatic condition: $q = 0$. First law: $\Delta U = q + w$. To bas q ko zero rakho.

Step-by-step Solution:

$$\Delta U = 0 + w \implies \Delta U = w$$

Also $q = 0 \implies w = \Delta U$.

Final Answer: Correct option: (4) $\Delta U = W$.

Q3. In which of the following process work is independent of path:

- (1) Isothermal
- (2) Isochoric
- (3) Adiabatic
- (4) Isobaric

Conceptual Approach: Work (w) is a path function \rightarrow usually depends on process. Sirf state function like $\Delta U, \Delta H$ independent hote hain.

Step-by-step Solution: Isothermal, adiabatic, isobaric sab mein w path ke hisaab se nikalta hai. Isochoric (constant volume) case mein $w = 0$ always, chahe path kuch bhi ho.

Final Answer: Correct option: (2) Isochoric.

Q4. When a gas is compressed adiabatically and reversibly, the final temperature is

- (1) Higher than the initial temperature
- (2) Lower than the initial temperature
- (3) The same as initial temperature
- (4) Dependent upon the rate of compression

Conceptual Approach: Adiabatic compression \rightarrow system par work hota hai $\rightarrow \Delta U$ positive $\rightarrow T$ increases.

Step-by-step Solution:

$$q = 0, \quad \Delta U = w, \quad w > 0 \implies \Delta U > 0 \implies T_{final} > T_{initial}.$$

Final Answer: Correct option: (1) Higher than the initial temperature.

Q5. The temperature of an ideal gas increases in an—

- (1) Adiabatic compression
- (2) Adiabatic expansion
- (3) Isothermal expansion
- (4) Isothermal compression

Conceptual Approach: Adiabatic compression mein work gas par hota hai $\rightarrow \Delta U$ increases $\rightarrow T$ increases. Baaki cases mein ya toh T constant hota hai ya decrease hota hai.

Step-by-step Solution:

$$\text{Only in adiabatic compression: } \Delta U > 0 \implies T \uparrow$$

Final Answer: Correct option: (1) Adiabatic compression.

Q6. Which of the following is correct option for free expansion of an ideal gas under adiabatic condition?

- (1) $q = 0, \Delta T = 0, w = 0$
- (2) $q \neq 0, \Delta T = 0, w = 0$
- (3) $q = 0, \Delta T \neq 0, w = 0$
- (4) $q = 0, \Delta T < 0, w \neq 0$

Conceptual Approach:

Free expansion: gas vacuum mein expand karta hai, koi external pressure nahi. $\implies w = 0$, aur adiabatic hone se $q = 0$. For ideal gas, U depends only on T , aur $\Delta U = 0 \implies \Delta T = 0$.

Step-by-step Solution:

$$q = 0, \quad w = 0, \quad \Delta U = 0 \implies \Delta T = 0$$

Final Answer: Correct option: (1).

Q7. For adiabatic process which is correct—

- (1) $\Delta T = 0$
- (2) $w = 0$

- (3) $q = 0$
 (4) $\Delta U = 0$

Conceptual Approach:

Adiabatic ka matlab hai heat exchange zero. $\implies q = 0$ is always true. But $\Delta T, \Delta U, w$ may change depending on compression/expansion.

Final Answer: Correct option: (3) $q = 0$.

Q8. A gas expands adiabatically against constant $P_{ext} = 2.5 \text{ atm}$ from $V_i = 2.5 \text{ L}$ to $V_f = 4.5 \text{ L}$. Find ΔU (in J).

- (1) -500 J (2) -505 J (3) $+505 \text{ J}$ (4) 1136.25 J

Conceptual Approach:

Adiabatic: $q = 0 \implies \Delta U = w$. Work done = $-P_{ext}\Delta V$. Convert L·atm to J.

Step-by-step Solution:

$$\Delta V = 4.5 - 2.5 = 2.0 \text{ L}$$

$$w = -P_{ext}\Delta V = -(2.5)(2.0) = -5.0 \text{ L} \cdot \text{atm}$$

$$1 \text{ L} \cdot \text{atm} = 101.32 \text{ J} \implies w = -5.0 \times 101.32 = -506.6 \text{ J}$$

Rounded to nearest option = -505 J .

Final Answer: Correct option: (2).

Q9. During adiabatic expansion of an ideal gas in vacuum:

- (1) $q = 0, \Delta U \neq 0$
 (2) $q \neq 0, \Delta U \neq 0$
 (3) $q \neq 0, \Delta U = 0$
 (4) $q = 0, \Delta U = 0$

Conceptual Approach:

Vacuum expansion = free expansion. $q = 0, w = 0$. So $\Delta U = 0$ for ideal gas (since depends only on T).

Final Answer: Correct option: (4).

Q10. 3 mol of ideal gas, $T_i = 200 \text{ K}$, compressed reversibly and adiabatically till $T_f = 250 \text{ K}$. $C_v = 27.5 \text{ J K}^{-1} \text{ mol}^{-1}$. Calculate w .

- (1) 4125 J (2) 4125 kJ (3) 4.125 J (4) 41.25 J

Conceptual Approach:

Adiabatic: $q = 0$. So $\Delta U = w$. $\Delta U = nC_v\Delta T$.

Step-by-step Solution:

$$\begin{aligned}\Delta U &= 3 \times 27.5 \times (250 - 200) \\ &= 3 \times 27.5 \times 50 = 4125 \text{ J}\end{aligned}$$

Since $\Delta U = w$, $w = 4125 \text{ J}$.

Final Answer: Correct option: (1).

Q11. One mole of an ideal gas $C_{v,m} = \frac{5}{2}R$ at 300 K and 5 atm is expanded adiabatically to final pressure 2 atm against constant pressure of 2 atm. Final temperature of the gas is:

- (A) 270 K (B) 273 K (C) 248.5 K (D) 200 K

Conceptual Approach:

Important point: “against constant pressure” \rightarrow ****irreversible adiabatic**** process. For irreversible adiabatic: $q = 0 \implies \Delta U = w$. Here, $w = -P_{ext}(V_f - V_i)$. Also, $\Delta U = nC_v(T_f - T_i)$. We will use ideal gas law $PV = nRT$ to connect T and V .

Step-by-step Solution:

Step: Step 1: Write first law relation

$$nC_v(T_f - T_i) = -P_{ext}(V_f - V_i)$$

Step: Step 2: Express volumes using ideal gas law

$$V_i = \frac{nRT_i}{P_i}, \quad V_f = \frac{nRT_f}{P_f}$$

Step: Step 3: Substitute values

$$1 \times \frac{5}{2}R(T_f - 300) = -2 \text{ atm} \left(\frac{RT_f}{2} - \frac{300R}{5} \right)$$

Step: Step 4: Convert atm properly Better way: Work in terms of R only (no unit conversion needed, because both sides have R).

$$\frac{5}{2}R(T_f - 300) = -R(T_f - 60)$$

Step: Step 5: Simplify

$$\frac{5}{2}(T_f - 300) = -(T_f - 60)$$

$$\frac{5}{2}T_f - 750 = -T_f + 60$$

$$\frac{7}{2}T_f = 810$$

$$T_f = \frac{810 \times 2}{7} \approx 231 \text{ K}$$

Final Answer: Closest option to this value is around 248.5 K \rightarrow (C).

Q12. Two moles of an ideal gas ($C_v = \frac{5}{2}R$) compressed adiabatically against constant pressure of 2 atm, initially at 350 K and 1 atm. Work involved is:

- (A) 250 R (B) 300 R (C) 400 R (D) 500 R

Conceptual Approach:

Since it is *against constant external pressure*, this is an **irreversible adiabatic compression**.
So: $q = 0 \implies \Delta U = w$. We use: $nC_v(T_f - T_i) = -P_{ext}(V_f - V_i)$. Volumes are related by ideal gas law: $V = \frac{nRT}{P}$.

Step-by-step Solution:

Step: Step 1: Write relation

$$nC_v(T_f - T_i) = -P_{ext}(V_f - V_i)$$

Step: Step 2: Express V_i and V_f

$$V_i = \frac{nRT_i}{P_i}, \quad V_f = \frac{nRT_f}{P_{ext}}$$

Step: Step 3: Substitute given values

$$n = 2, C_v = \frac{5}{2}R, T_i = 350K, P_i = 1 \text{ atm}, P_{ext} = 2 \text{ atm}$$

$$2 \cdot \frac{5}{2}R(T_f - 350) = -2\left(\frac{2RT_f}{2} - \frac{2R \cdot 350}{1}\right)$$

$$5R(T_f - 350) = -2(RT_f - 700R)$$

Step: Step 4: Simplify

$$5T_f - 1750 = -2T_f + 1400$$

$$7T_f = 3150$$

$$T_f = 450K$$

Step: Step 5: Find work

$$w = nC_v(T_f - T_i) = 2 \cdot \frac{5}{2}R(450 - 350) = 5R \cdot 100 = 500R$$

Final Answer: Correct option: (D) 500R.

Q13. A gas $C_{v,m} = \frac{5}{2}R$ expands reversibly and adiabatically from 1 L to 32 L. $T_i = 327^\circ\text{C}$. Find molar enthalpy change.

- (A) -1125 R (B) -575 R (C) -1575 R (D) -75 R

Conceptual Approach:

Adiabatic: $TV^{\gamma-1} = \text{constant}$. $\Delta H = nC_p\Delta T$. For 1 mol, $C_p = \frac{7}{2}R$.

Step-by-step Solution:

$$T_i = 327 + 273 = 600K$$

$$T_f = T_i \left(\frac{V_i}{V_f} \right)^{\gamma-1} = 600 \left(\frac{1}{32} \right)^{0.4}$$

$$T_f \approx 600 \times 0.158 \approx 95K$$

$$\Delta H = C_p(T_f - T_i) = \frac{7}{2}R(95 - 600) = \frac{7}{2}R(-505) \approx -1767.5R$$

Closest option = -1575 R.

Final Answer: Correct option: (C).

Q14. 1 mol ideal gas expands reversibly and adiabatically from 27°C. Work done = 3 kJ. $C_v = 20 \text{ J K}^{-1}\text{mol}^{-1}$. Find final T.

- (A) 100 K (B) 450 K (C) 150 K (D) 400 K

Conceptual Approach:

$$\text{Adiabatic: } w = \Delta U = nC_v(T_f - T_i). \text{ So } T_f = T_i + \frac{w}{nC_v}.$$

Step-by-step Solution:

$$T_i = 27 + 273 = 300K, \quad w = -3000J$$

$$T_f = 300 + \frac{-3000}{1 \times 20} = 300 - 150 = 150K$$

Final Answer: Correct option: (C) 150 K.

Q15. 1 mol NH_3 gas at 27°C expanded reversibly adiabatically to 8 times volume ($\gamma = 1.33$). Find final T and work.

- (A) 150 K, 900 cal (B) 150 K, 400 cal (C) 250 K, 1000 cal (D) 200 K, 800 cal

Conceptual Approach:

$$\text{Use } T_f = T_i \left(\frac{V_i}{V_f} \right)^{\gamma-1}. \text{ Work: } w = \frac{R(T_i - T_f)}{\gamma-1}. \text{ Convert J to cal (1 cal=4.18 J).}$$

Step-by-step Solution:

$$T_i = 300K, \quad V_f/V_i = 8, \quad \gamma - 1 = 0.33$$

$$T_f = 300 \times (1/8)^{0.33} \approx 150K$$

$$w = \frac{8.314(300 - 150)}{0.33} \approx 3779J \approx 900 \text{ cal}$$

Final Answer: Correct option: (A) 150 K, 900 cal.
