

**Topic : Thermodynamics & Thermochemistry**

**Type of Questions**

Single choice Objective ('-1' negative marking) Q.1 to Q.7

(3 marks, 3 min.)

M.M., Min.

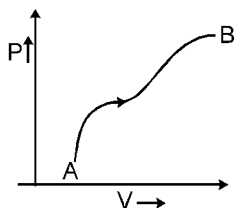
[21, 21]

Comprehension ('-1' negative marking) Q.8 to Q.10

(3 marks, 3 min.)

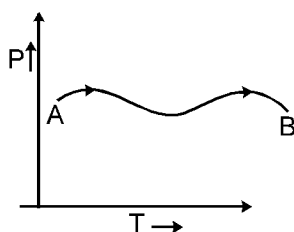
[9, 9]

1. The graph given below shows the P-V plot for a process on an ideal gas. Select the correct statement :



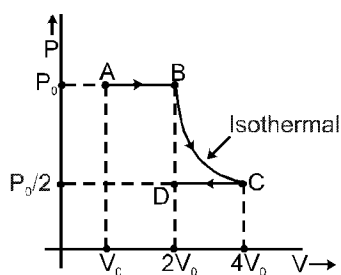
- (A) Enthalpy content of the gas is constantly increasing and the process is carried out slowly.
- (B) Enthalpy content of the gas first increases, then decreases and the process is carried out quasistatically.
- (C) Enthalpy content of the gas is constant and the process takes infinite time for completion.
- (D) Enthalpy content first decreases, then increases and the process is reversible.

2. The P-T graph, as given below, was observed for a process on an ideal gas. Which of the following statement is true :



- (A)  $W = +ve, \Delta H = +ve$
- (B)  $W = -ve, \Delta H = -ve$
- (C)  $W = -ve, \Delta H = +ve$
- (D)  $W = +ve, \Delta H = -ve$

3.  $q, W, \Delta E$  and  $\Delta H$  for the following process ABCD on a monoatomic gas are :



- (A)  $W = -2 P_0 V_0 \ln 2, \quad q = 2 P_0 V_0 \ln 2, \quad \Delta E = 0, \quad \Delta H = 0$
- (B)  $W = -2 P_0 V_0 \ln 2, \quad q = 2 P_0 V_0 \ln 2, \quad \Delta E = 0, \quad \Delta H = 2 P_0 V_0 \ln 2$
- (C)  $W = -P_0 V_0 (1 + \ln 2), \quad q = P_0 V_0 (1 + \ln 2), \quad \Delta E = 0, \quad \Delta H = 0$
- (D)  $W = -P_0 V_0 \ln 2, \quad q = P_0 V_0 \ln 2, \quad \Delta E = 0, \quad \Delta H = 0$

4. A system containing a real gas changes its state from state-1 to state-2.

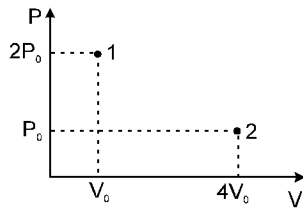
State-1 (3 atm, 2L, 300 K)

State-2 (5 atm, 4L, 500 K)

If change in internal energy = 30 L atm, then calculate change in enthalpy :

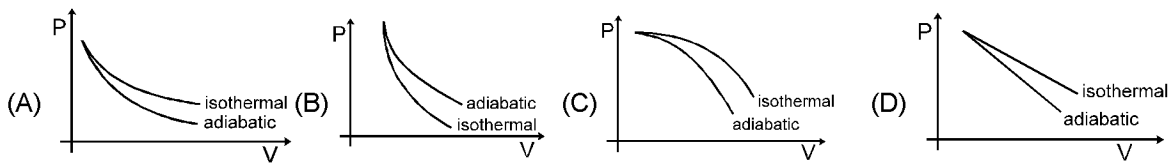
- (A) 44 L atm
- (B) 35 L atm
- (C) 40 L atm
- (D) None of these

5. A liquid which is confined inside an adiabatic piston is suddenly taken from state-1 to state-2 by a single stage irreversible process. If the piston comes to rest at point 2 as shown, then the enthalpy change for the process will be :

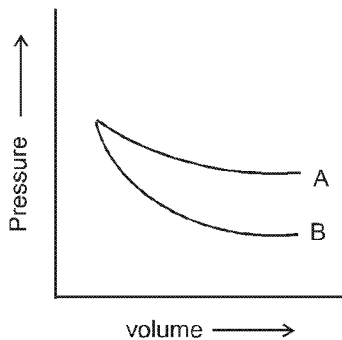


- (A)  $\Delta H = \frac{2\gamma P_0 V_0}{\gamma - 1}$       (B)  $\Delta H = \frac{3\gamma P_0 V_0}{\gamma - 1}$       (C)  $\Delta H = -P_0 V_0$       (D) None of these

6. The correct figure representing isothermal and adiabatic expansions of an ideal gas from a particular initial state is :



7. P-V plots for two gases during an adiabatic process are given in the figure :

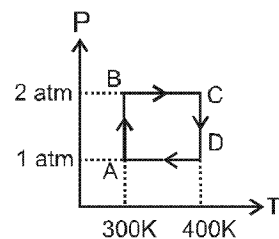


Plot A and plot B should correspond to :

- (A) He and O<sub>2</sub>      (B) He and Ar      (C) O<sub>2</sub> and He      (D) O<sub>2</sub> and F<sub>2</sub>

### Comprehension # (Q. Nos. 8 to 10)

One mole of Helium gas undergoes a reversible cyclic process ABCDA as shown in the figure. Assuming gas to be ideal, answer the following questions:



8. What is the value of  $\Delta H$  for the overall cyclic process :  
 (A)  $-100 R \ln 2$       (B)  $+100 R \ln 2$       (C)  $+200 R \ln 2$       (D) Zero
9. What is the value of 'q' for the overall cyclic process :  
 (A)  $-100 R \ln 2$       (B)  $+100 R \ln 2$       (C)  $+200 R \ln 2$       (D)  $-200 R \ln 2$
10. What is the net work involved in the cyclic process :  
 (A)  $-100 R \ln 2$       (B)  $+100 R \ln 2$       (C)  $+200 R \ln 2$       (D)  $-200 R \ln 2$

# Answer Key

## DPP No. # 47

- |        |        |        |        |         |
|--------|--------|--------|--------|---------|
| 1. (A) | 2. (C) | 3. (A) | 4. (A) | 5. (C)  |
| 6. (A) | 7. (C) | 8. (D) | 9. (B) | 10. (A) |

# Hints & Solutions

## DPP No. # 47

- The product  $PV$  is increasing so temperature will keep on increasing in the process, hence  $\Delta H = \Delta E + \Delta(PV)$  will increase constantly.
- From graph we know that  $V_B > V_A$ , so expansion has taken place so  $w$  will be with  $-ve$  sign and  $\Delta H$  will be  $+ve$  as both  $\Delta E$  and  $\Delta(PV)$  have increased.
- At A and D the temperatures of the gas will be equal, so  $\Delta E = 0$ ,  $\Delta H = 0$   
Now  $w = W_{AB} + W_{BC} + W_{CD} = -P_0 V_0 - 2P_0 V_0 \ln 2 + P_0 V_0 = -2P_0 V_0 \ln 2$   
and  $q = -W = 2P_0 V_0 \ln 2$
- Since liquid is expanding against external pressure  $P_0$  hence work done  
 $w = -P_0(4V_0 - V_0) = -3P_0 V_0$   
 $\Delta U = w = -3P_0 V_0$   
 $\Rightarrow \Delta H = \Delta U + P_2 V_2 - P_1 V_1 = -3P_0 V_0 + 4P_0 V_0 - 2P_0 V_0$
- $\gamma$  for  $O_2 = 1.44$   $\gamma$  for He = 1.66.
- Since,  $\Delta H$  is a state function, and the final state attained by the gas is same as its initial state, so value of  $\Delta H = 0$ .
- $q = q_{AB} + q_{BC} + q_{CD} + q_{DA}$   
 $= -1R \times 300 \ln 2 + 1 \times \frac{5R}{2} \times (400 - 300) + 1R \times 400 \ln 2 + 1 \times \frac{5R}{2} \times (300 - 400)$   
( $\because q_{AB} = -W_{AB} = -1R \times 300 \ln 2$  since process is reversible isothermal for which  $\Delta U = 0$ ).  
( $\because q_{BC} = \Delta H_{BC} = 1 \times \frac{5R}{2} \times (400 - 300)$  since process is reversible isobaric).  
( $\because q_{CD} = -W_{CD} = 1R \times 400 \ln 2$  since process is reversible isothermal for which  $\Delta U = 0$ ).  
( $\because q_{DA} = \Delta H_{DA} = 1 \times \frac{5R}{2} \times (300 - 400)$  since process is reversible isobaric).  
So,  $q = 100 R \ln 2$ .
- Since, for a cyclic process,  $\Delta U = 0$ .  
So,  $W = -q = -100 R \ln 2$ .

