11 Thermodynamics

TOPIC 1

Zeroth and First Law of Thermodynamics

01 1g of water, of volume 1 cm^3 at 100°C is converted into steam at same temperature under normal atmospheric pressure=($\simeq 1 \times 10^5$ Pa). The volume of steam formed equals 1671 cm³. If the specific latent heat of vaporisation of water is 2256 J/g, the change in internal energy is [NEET (Odisha) 2019] (a) 2423 J (b)2089J (c)167 J (d) 2256 J Ans. (b) Given, mass of water, m = 1g Volume of 1 g of water = $1 \text{ cm}^3 = 10^{-6} \text{ m}^3$ Volume of 1 g of steam = 1671 cm^3 $= 1671 \times 10^{-6} \text{ m}^{3}$ Pressure, $p = 1 \times 10^5$ Pa Latent heat of vaporization of water, L = 2256 J/gChange in volume, $\Delta V = (1671 - 1) \times 10^{-6} \text{ m}^3$ $= 1670 \times 10^{-6} \text{ m}^3$...(i) Heat supplied, $\Delta Q = mL = 1 \times 2256 = 2256 J$...(ii) As the steam expands, so the work done in expansion is $\Delta W = p \Delta V$ $= 1 \times 10^{5} \times 1670 \times 10^{-6}$ [from Eq. (i)] = 167 J ...(iii) According to first law of thermodynamics, $\Delta Q = \Delta U + \Delta W$ $\Rightarrow \Delta U = \Delta Q - \Delta W$ = 2256 - 167 [from Eq. (ii) and (iii)] =2089J

02 A sample of 0.1 g of water at 100°C and normal pressure $(1.013 \times 10^5 \text{ Nm}^{-2})$ requires 54 cal of heat energy to convert to steam at 100°C. If the volume of the steam produced is 167.1 cc, the change in internal energy of the sample, is **[NEET 2018]** (a) 42.2 J (b) 208.7 J (c) 104.3 J (d) 84.5 J

Ans. (b)

According to the question, Heat spent during the conversion of sample of water at 100° C to steam is, $\Delta Q = 54 \, \text{cal} = 54 \times 4.18 \, \text{J}$ = 225.72 J Normal pressure, $p = 1.013 \times 10^5$ Nm⁻² Net work done during the conversion would be given as $\Delta W = p \Delta V$ $= p[V_{\text{steam}} - V_{\text{water}}]$ Here, $V_{steam} = 167.1 \text{ cc} = 167.1 \times 10^{-6} \text{ m}^3$ $V_{water} = 0.1 \text{ g} = 0.1 \text{ cc} = 0.1 \times 10^{-6} \text{ m}^3$ $\therefore \Delta W = 1.013 \times 10^{5} [(167.1 - 0.1) \times 10^{-6}]$ $= 1.013 \times 167 \times 10^{-1}$ = 16.917 J Now, by the first law of thermodynamics, $\Delta Q = \Delta U + \Delta W$ where, ΔU is the change in internal energy of the sample. $\Rightarrow \Delta U = \Delta Q - \Delta W$ Substituting the values in the above equation, we get $\Delta U = 225.72 - 16.917 = 208.7 \text{ J}$

03 During an isothermal expansion, a confined ideal gas does -150 J of work against its surroundings. This implies that **[CBSE AIPMT 2011]** (a) 300 J of heat has been added to the gas

- (b) no heat is transferred because the process is isothermal
- (c) 150 J of heat has been added to the gas
- (d) 150 J of heat has been removed from the gas

Ans. (c)

Concept Apply first law of thermodynamics. According to first law of thermodynamics $\Delta U = \Delta Q + \Delta W$ For isothermal process, $\Delta U = 0$ So, $\Delta Q = -\Delta W$ Given, $\Delta W = -150 \text{ J}$ So, $\Delta Q = +150 \text{ J}$ When Q is positive, the heat is added to the gas.

04 If ΔU and ΔW represent the

increase in internal energy and work done by the system respectively in a thermodynamical process, which of the following is true? **[CBSE AIPMT 2010]** (a) $\Delta U = -\Delta W$, in an adiabatic process (b) $\Delta U = \Delta W$, in an isothermal process (c) $\Delta U = \Delta W$, in an adiabatic process (d) $\Delta U = -\Delta W$, in an isothermal process **Ans.** (a) From first law of thermodynamics

-rom first law	of thermodynamics
	$\Delta Q = \Delta U + \Delta W$

For adiabatic process, $\Delta Q = 0$

::	$\Delta Q = 0$	
So,	$\Delta U = -\Delta W$	

05 The internal energy change in a system that has absorbed 2 kcal of heat and done 500 J of work is [CBSE AIPMT 2009]

(a) 8900 J	(b)6400 J
(c)5400 J	(d)7900 J

Ans. (d)

According to first law of thermodynamics

$$\Delta Q = U + W$$

or $\Delta U = Q - W$

 ΔU = change in internal energy

Q = heat given to system

- W = work done $\therefore \quad \Delta U = 2 \times 4.2 \times 1000 - 500$
- = 8400 500 = 7900 J

06 If *Q*, *E* and *W* denote respectively

the heat added, change in internal energy and the work done in a closed cycle process, then

	Lobor VII IIII
(a)W=0	(b) $Q = W = 0$
(c) E = 0	(d) Q = 0

Ans. (c)

For a cyclic process, $\Delta U = 0$ or E = 0

07 We consider a thermodynamic system. If ΔU represents the increase in its internal energy and W the work done by the system, which of the following statements is true? [CBSE AIPMT 1998] (a) $\Delta U = -W$ in an adiabatic process (b) $\Delta U = W$ in an isothermal process (c) $\Delta U = -W$ in an isothermal process (d) $\Delta U = W$ in an adiabatic process

Ans.(a)

An isothermal process is a constant temperature process. In this process, T = constant or ΔT = 0.

$$\therefore \quad \Delta Q = \Delta U + \Delta W$$

$$\Rightarrow \Delta Q = \Delta W$$
$$\Delta U = nC_v \Delta T = 0$$

An adiabatic process is defined as one with no heat transfer into or out of a system. Therefore, Q = 0. From the first law of thermodynamics.

 $(\Delta U = 0)$

$$\Delta Q = \Delta U + \Delta W$$

or $\Delta U = -W$ [$\Delta Q = 0$]

08 110 J of heat is added to a gaseous system, whose internal energy is 40 J, then the amount of external work done is **[CBSE AIPMT 1993]** (a) 150 J (b) 70 J (c) 110 J (d) 40 J

Ans. (b)

Concept Apply first law of thermodynamics to calculate the required work done.

From first law of thermodynamics

$$\Delta Q = \Delta U + \Delta W$$
where, ΔQ = heat given
 ΔU = change in internal energy
 ΔW = work done
Here, ΔQ = 110 J
 ΔU = 40 J
 $\therefore \Delta W = \Delta Q - \Delta U$ = 110 - 40 = 70 J

09 First law of thermodynamics is a consequence of conservation of [CBSE AIPMT 1988]

	Fo= 0=
(a) work	(b)energy
(c) heat	(d)All of these

Ans. (b)

According to the first law of thermodynamics, when some quantity of heat (dQ) is supplied to a system capable of doing external work, then the quantity of heat absorbed by the system (dQ) is equal to the sum of the increase in the internal energy of the system (dU) due to rise in temperature and the external work done by the system (dW) in expansion,

i.e. dQ = dU + dW

This law, which is basically the law of conservation of energy applies to every process in nature.

TOPIC 2 Thermodynamic Process

10 The *p*-V diagram for an ideal gas in a piston cylinder assembly undergoing a thermodynamic process is shown in the figure. The process is **[NEET (Oct.) 2020]**



(a) adiabatic (c) isobaric



Ans. (c)



From the *pV* diagram, it is clear that pressure of ideal gas in the piston cylinder is constant during thermodynamic process. Hence, this process is isobaric.

11 Two cylinders *A* and *B* of equal capacity are connected to each other *via* a stop cock. A contains an ideal gas at standard temperature and pressure. *B* is completely evacuated. The entire system is thermally insulated. The stop cock is suddenly opened. The process is [NEET (Sep.) 2020]

(a) adiabatic (c) isobaric

(b)isochoric (d)isothermal

Ans. (a)

Since, the entire system is thermally insulated. Thus, their will be no transfer of heat between the system and the surrounding. Hence, when the stop cock is suddenly opened, a sudden process of expansion will takes place with no heat transfer. Thus, the given process will be adiabatic, as it is a process that occurs without transferring heat or mass between a thermodynamic system and its surroundings.

Hence, correct option is (a).

12 In which of the following processes, heat is neither absorbed nor released by a system? [NEET (National) 2019]

 (a) Adiabatic
 (b) Isobaric
 (c) Isochoric
 (d) Isothermal

Ans. (a)

In an adiabatic process, the system is completely insulated from the surroundings. Thus, heat is neither absorbed nor released by the system to the surroundings. So, $\Delta Q = 0$.

Sudden processes are adiabatic like bursting of cycle tyre, etc.

If the pressure of gas is kept constant, then the process is called isobaric, i.e. $\Delta p = 0$.

If the temperature of the system remains constant, then it is called isothermal process, i.e. $\Delta T = 0$. If the volume of gas is constant in a

system, then it is called isochoric process, i.e. $\Delta V = 0$.

13 The volume (V) of a monoatomic gas varies with its temperature (T), as shown in the graph. The ratio of work done by the gas, to the heat absorbed by it, when it undergoes a change from state A to state B, is [NEET 2018]



Ans. (c)

According to the given graph, Volume (V) \propto Temperature (T)

or
$$\frac{v}{\tau} = \text{constant}$$

Thus, the process is isobaric.



 \therefore Work done, $\Delta W = p \Delta V$

$$= nR\Delta T = nR(T_{B} - T_{A}) \dots (i)$$

Heat absorbed $\Delta Q = nC_{A}\Delta T$

 $= nC_{\rho}(T_{B} - T_{A}) \qquad \dots (ii)$ As, $C_{\rho} = \frac{\gamma R}{1} , \quad \left(\text{where, } \gamma = 1 + \frac{2}{4} \right)$

For a monoatomic gas,
$$f = 3$$

$$\Rightarrow \qquad C_{p} = \left(R + \frac{3}{2}R\right) = \frac{3}{2}R$$

Substituting the value of C_p in Eq. (ii), we get

$$\Delta 0 = n \left(\frac{5}{2}R\right) (T_B - T_A)$$

Hence, $\frac{\Delta W}{\Delta 0} = \frac{nR(T_B - T_A)}{n \left(\frac{5}{2}R\right) (T_B - T_A)} = \frac{2}{5}$

14 Thermodynamic processes are indicated in the following diagram [NEET 2017]



Match the following :

		-	
	Column-I		Column-II
Ρ.	Process I	a.	Adiabatic
Q.	Process II	b.	Isobaric
R.	Process III	с.	Isochoric
S.	Process IV	d.	lsothermal
-			

 $\begin{array}{l} (a)P \rightarrow a, Q \rightarrow c, R \rightarrow d, S \rightarrow b \\ (b)P \rightarrow c, Q \rightarrow a, R \rightarrow d, S \rightarrow b \\ (c)P \rightarrow c, Q \rightarrow d, R \rightarrow b, S \rightarrow a \\ (d)P \rightarrow d, Q \rightarrow b, R \rightarrow a, S \rightarrow c \end{array}$

Ans. (b)

In isochoric process, the curve is parallel to y-axis because volume is constant.lsobaric is parallel to x-axis because pressure is constant. Along the curve, it will be isothermal because temperature is constant.

So, $P \rightarrow c \Rightarrow Q \rightarrow a \Rightarrow R \rightarrow d$

 \Rightarrow S \rightarrow b

- **15** A gas is compressed isothermally to half its initial volume. The same gas is compressed separately through an adiabatic process until its volume is again reduced to half. Then **[NEET 2016]**
 - (a) compressing the gas through adiabatic process will require more work to be done.
 - (b) compressing the gas isothermally or adiabatically will require the same amount of work.
 - (c) which of the case (whether compression through isothermal or through adiabatic process) requires more work will depend upon the atomicity of the gas.
 - (d) compressing the gas isothermally will require more work to be done.

Ans. (a)

The solution of this question can be understood by plotting a *p*-V graph for the compression of a gas isothermally and adiabatically simultaneously to half of its initial volume. *i.e.*



Since, the isothermal curve is less steeper than the adiabatic curve. So, area under the *p*-V curve for adiabatic process has more magnitude than isothermal curve. Hence, work done in adiabatic process will be more than in isothermal process.

16 Figure below shows two paths that may be taken by a gas to go from a state *A* to a state *C*.





In process AB, 400 J of heat is added to the system and in process BC, 100 J of heat is added to the system. The heat absorbed by the system in the process AC will be (a) 380 J (b) 500 J (c) 460 J (d) 300 J

Ans. (c)

Since, initial and final points are same

So, $\Delta U_{A \to B \to c} = \Delta U_{A \to c}$...(i) Also $A \to B$ is isochoric process So dW = 0 and d0 = dU + dW

So
$$dW_{A \rightarrow B} = U$$
 and $dV = dU + dW$
So, $dQ_{A \rightarrow B} = dU_{A \rightarrow B} = 400 \text{ J}$



Next $B \rightarrow C$ is isobaric process

So, $dQ_{B \to C} = dU_{B \to C} + dW_{B \to C}$

 $= dU_{B \to C} + p\Delta V_{B \to C}$ $\Rightarrow \quad 100 = dU_{B \to C} + 6 \times 10^4 (2 \times 10^{-3})$

$$\Rightarrow dU_{B \to C} = 100 - 120 = -20 \text{ J}$$

From Eq. (i),
$$\therefore \Delta U_{A \to B \to C} = \Delta U_{A \to C}$$

$$\Rightarrow \Delta U_{A \to B} + \Delta U_{B \to C} = dQ_{A \to C} - dW_{A \to C}$$

$$\Rightarrow 400 + (-20) = dQ_{A \to C} - (p\Delta V_A + \text{ Area of } \Delta ABC)$$

$$\Rightarrow dQ_{A \to C} = 380 + \left(\frac{2 \times 10^4 \times 2 \times 10^{-3}}{4 \times 2 \times 10^{-3} \times 4 \times 10^4}\right)$$

$$= 380 + (40 + 40)$$

$$dQ_{A \to C} = 460 \text{ J}$$

17 One mole of an ideal diatomic gas undergoes a transition from *A* to *B* along a path *AB* as shown in the figure. **[CBSE AIPMT 2015]**



The change in internal energy of the gas during the transition is (a) 20 kJ (b) -20 kJ (c) 20 J (d) -12 kJ

Ans. (b)

For a diatomic gas,



The change in internal energy of gas in the transition from A to B is

$$\Delta U = nC_v dT = n\left(\frac{5R}{2}\right)(T_B - T_A)$$

= $nR\frac{5}{2}\left(\frac{p_B V_B}{nR} - \frac{p_A V_A}{nR}\right)$
= $\frac{5}{2}(2 \times 10^3 \times 6 - 5 \times 10^3 \times 4)$
= $\frac{5}{2} \times (-8 \times 10^3) = -\frac{4 \times 10^4}{2} = -20 \text{ kJ}$

18 An ideal gas is compressed to half its initial volume by means of several process. Which of the process results in the maximum work done on the gas? [CBSE AIPMT 2015] (a) Adiabatic (c) Isochoric

(b)Isobaric (d)Isothermal

Ans.(a)

Given, ideal gas is compressed to half its initial volume i.e.



The isochoric process is one in which volume is kept constant, meaning that work done by the system will be zero. i.e.

$$W_{\rm isochoric} = 0$$

As we know, work done on the gas = Area under curve, i.e.

$$W_{\rm adiabatic} > W_{\rm isothermal} > W_{\rm isobaric}$$

19 A monoatomic gas at a pressure *p*, having a volume *V* expands isothermally to a volume 2 *V* and then adiabatically to a volume 16 *V*. The final pressure of the gas is

$$\begin{pmatrix} \text{take } \gamma = \frac{5}{3} \\ \text{(a)} 64p \\ \text{(c)} \frac{p}{64} \\ \text{(d)} 16p \\ \end{pmatrix}$$

Ans. (c)

For isothermal expansion process

$$pV = p' \times 2V \qquad [::V' = 2V]$$
$$p' = \frac{p}{2}$$

For adiabatic expansion, $pV^{\gamma} = \text{constant} \Rightarrow$ $p'V^{\gamma} = p''V'^{\gamma}$

$$\Rightarrow \frac{p}{2}(2V)^{5/3} = p''(16V)^{5/3}$$

$$\Rightarrow p'' = \frac{p}{2} \left[\frac{2V}{16V} \right]^{5/3} = \frac{p}{2} \left(\frac{1}{8} \right)^{5/3}$$
$$= \frac{p}{2} \left(\frac{1}{32} \right) = p/64$$

20 A thermodynamic system undergoes cyclic process *ABCDA* as shown in figure. The work done by the system in the cycle is [CBSE AIPMT 2014]



Ans.(d)

Work done in the cyclic process = Area bounded by the closed configuration

= Area of closed configuration = $\frac{1}{2} \times V_0 \times p_0 - \frac{1}{2} \times V_0 \times p_0 = 0$ (zero).

21 A gas is taken through the cycle $A \rightarrow B \rightarrow C \rightarrow A$, as shown. What is the net work done by the gas? [NEET 2013]



Ans. (b)

According to question, Net work done = Area enclosed in pVcurve i.e. ΔABC Area of ΔABC

$$= \frac{1}{2} \times 5 \times 10^{-3} \times 4 \times 10^{5} \text{ J}$$
$$= 10^{3} \text{ J} = 1000 \text{ J}$$

22 A thermodynamic system is taken through the cycle *ABCD* as shown in figure. Heat rejected by the gas during the cycle is

[CBSE AIPMT 2012]



(a)2*pV* (b)4*pV* (c) $\frac{1}{2}$ *pV* (d)*pV*

Ans. (a)

For given cyclic process, $\Delta U = 0 \implies Q = W$ Also, W = - area enclosed by the curve $= -AB \times AD$ = -(2p - p)(3V - V) $= -p \times 2V$

:. Heat rejected = 2pV

23 One mole of an ideal gas goes from an initial state *A* to final state *B via* two processes. It first undergoes isothermal expansion from volume *V* to 3*V* and then its volume is reduced from 3*V* to *V* at constant pressure. The correct *p*-*V* diagram representing the two processes is [CBSE AIPMT 2012]



Ans. (d)

According to question, firstly gas expands from volume V to 3V and after this volume is reduced from 3V to V at constant pressure. In isothermal expansion, *p*-V curve is

rectangular hyperbola.

- 24 In thermodynamic processes which of the following statements is not true ? [CBSE AIPMT 2009]
 - (a) In an adiabatic process the system is insulated from the surroundings

- (b) In an isochoric process pressure remains constant
- (c) In an isothermal process the temperature remains constant(d) In an adiabatic process
- $pV^{\gamma} = \text{constant}$

Ans. (b)

For an adiabatic process, there should not be any exchange of heat between the system and its surroundings. All walls of the container must be perfectly insulated. In adiabatic changes, gases obey Poisson's law, i.e., $pV^{\gamma} = \text{constant}$. In an isochoric process, volume remains constant and for isobaric process, pressure remains constant.

25 Which of the following processes is reversible? [CBSE AIPMT 2005]

- (a) Transfer of heat by radiation
- (b) Electrical heating of a nichrome wire
- (c) Transfer of heat by conduction
- (d) Isothermal compression

Ans. (d)

(a) It is the process by which heat is transmitted from one place to another without heating the intervening medium. Hence, it is not a reversible process.
(b) Nichrome wire is made of alloy and has high resistance. When current is passed through it, heat is produced. So, here electrical energy is converted into heat energy. Hence, it is not a reversible

process.

(c) It is the process by which heat is transmitted from one point to another through a substance in the direction of fall of temperature without the actual movement of the particles of the substances themselves. Hence, it cannot be reversible.

(d) Isothermal compression is reversible, for example-Carnot cycle, Heat engine.

Thus, choice (d) is correct.

26 One mole of an ideal gas at an initial temperature of *T* K does 6*R* joules of work adiabatically. If the ratio of specific heats of this gas at constant pressure and at constant volume is 5/3, the final temperature of gas will be

[CBSE AIPMT 2004]

(a) (T + 2.4) K (b) (T - 2.4) K (c) (T + 4) K (d) (T - 4) K

Ans. (d)

In an adiabatic process,

Q = 0

So, from 1st law of thermodynamics, $\Delta 0 = \Delta U + \Delta W$

As
$$\Delta Q = 0$$

so, $W = -\Delta U = -nC_v \Delta T$
 $= -n\left(\frac{R}{\gamma - 1}\right)(T_f - T_i)$
 $= \frac{nR}{\gamma - 1}(T_i - T_f)$...(i)

Given, work done, W = 6R J, n = 1 mol,

$$R = 8.31 \text{ J/mol-K}, \gamma = \frac{5}{3}, T_i = T \text{ K}$$

Substituting given values in Eq. (i), we get

$$\therefore \qquad 6R = \frac{R}{(5/3 - 1)}(T - T_r)$$

$$\Rightarrow \qquad 6R = \frac{3R}{2}(T - T_r) \Rightarrow T - T_r = 4$$

 $\begin{array}{ccc} & & T_{f} = (T-4) \ \mathrm{K} \\ & \mbox{Adiabatic expansions of mono, dia} \\ & \mbox{and polyatomic gases are shown below.} \\ & \mbox{1} \rightarrow \mbox{monoatomic} & \mbox{2} \rightarrow \mbox{diatomic} & \mbox{3} \\ & \mbox{} \rightarrow \mbox{polyatomic} \end{array}$



27 If the ratio of specific heat of a gas at constant pressure to that at constant volume is γ , the change in internal energy of a mass of gas when the volume changes from *V* to 2*V* at constant pressure *p* is

a)
$$\frac{R}{(\gamma-1)}$$
 (b) pV (c) $\frac{pV}{(\gamma-1)}$ (d) $\frac{\gamma pV}{(\gamma-1)}$

Ans. (c)

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Change in internal energy of a gas having atomicity $\boldsymbol{\gamma}$ is given by

$$\Delta U = \frac{1}{(\gamma - 1)} (p_2 V_2 - p_1 V_1)$$

Given, $V_1 = V, V_2 = 2V$
So, $\Delta U = \frac{1}{\gamma - 1} [p \times 2V - p \times V]$
 $= \frac{1}{\gamma - 1} \times pV = \frac{pV}{\gamma - 1}$

28 A sample of gas expands from volume V_1 to V_2 . The amount of work done by the gas is greatest when the expansion is

[CBSE AIPMT 1997]

(a) adiabatic (b) isobaric (c) isothermal (d) Equal in all above cases

Ans. (b)

The *p*-V diagram for isobaric, isothermal and adiabatic processes of an ideal das is shown in graph below



In thermodynamics, for some change in volume, the work done is maximum for the curve having maximum area enclosed with the volume axis. Area enclosed by the curve

∝(Slope of curve)

Note

$$\Rightarrow (Area)_{sobaric} > (Area)_{sothermal} > (Area)_{aciabatic}$$

Hence, work done is maximum in isobaric process

 $(\text{Slope})_{\text{adiabatic}} = -\gamma \left(\frac{p}{V}\right)$ and (Slope)_{isothermal} = $-\frac{p}{V}$

:.. $(Slope)_{adiabatic} = \gamma \times (slope)_{sothermal}$

Slope of adiabatic curve is always steeper than that of isothermal curve.

29 An ideal gas undergoing adiabatic change has the following pressure-temperature relationship [CBSE AIPMT 1996]

(a) $p^{\gamma-1}T^{\gamma} = \text{constant}$ (b) $p^{\gamma}T^{\gamma-1} = \text{constant}$

(c) $p^{\gamma}T^{1-\gamma}$ = constant

(d) $n^{1-\gamma T\gamma}$ - constant

$$(0)p$$
 $(1) = constant$

Ans. (d)

The adiabatic relation between pand V for a perfect gas is

$$pV^{\gamma} = k$$
 ...(i)

$$pV = RT$$
 or $V = \frac{RT}{p}$
Putting value of V in Eq. (i)
 $p\left(\frac{RT}{p}\right)^{\gamma} = k$
or $p^{1-\gamma}T^{\gamma} = \frac{k}{R^{\gamma}} =$ another constant

 $p^{1-\gamma}T^{\gamma} = \text{constant}$ i.e.,

30 An ideal gas A and a real gas B have their volumes increased from V to 2V under isothermal conditions. The increase in internal energy [CBSE AIPMT 1993]

(a) will be same in both A and B (b) will be zero in both the gases (c) of B will be more than that of A (d) of A will be more than that of B

Ans. (b)

A change in pressure and volume of a gas without any change in its temperature is called an isothermal change. In such a change, there is a free exchange of heat between the gas and its surroundings.

 \therefore T = constant, $\Delta T = 0$ So, internal energy (U) remains constant 0.

31 Which of the following is not thermodynamical function? [CBSE AIPMT 1993]

(b) Work done (a) Enthalpy (c) Gibb's energy

(d) Internal energy

Ans. (b)

The thermodynamic state of a homogeneous system may be represented by certain specific thermodynamic variables such as pressure p, volume V, temperature T and entropy S. Out of these four variables, any two are independent and when they are known the others may be determined. Thus, there are only two independent variables and the others may be considered their functions. For complete knowledge of the system certain relations are required and for this purpose we introduce some functions of variables p, V, T and S known as thermodynamic functions. There are four principal thermodynamic functions

(i) Internal energy (U)

- (ii) Helmholtz function (F)
- (iii) Enthalpy(H)
- (iv) Gibb's energy (G)

Hence, work done is not thermodynamic function.

32 A thermodynamic system is taken from state A to B along ACB and is brought back to A along BDA as shown in the p-V diagram. The net work done during the complete cycle is given by the area

[CBSE AIPMT 1992]





Ans. (c)

Work done during path ACB = area ACBB'A'A Work done during path BDA = area BDAA'B'B ... Work done during going from ACB and

then to BDA path is = area ACB B'A'A – area BDAA'B'B

= area ACBDA

Net work done in cyclic process is given by area under the cycle.

33 A thermodynamic process is shown in the figure. The pressure and volumes corresponding to some points in the figure are



 $p_A = 3 \times 10^4$ pa, $V_A = 2 \times 10^{-3}$ m³ $p_B = 8 \times 10^4$ pa, $V_B = 5 \times 10^{-3}$ m³ In process AB, 600 J of heat is added to the system and in process BC, 200 J of heat is added to the system. The change in internal energy of the system in process AC would be [CBSE AIPMT 1991]

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a)560 J	(b)800J	
c)600J	(d)640J	

Ans. (a)

For path AB, applying first law of thermodynamics dQ = dU + dW or $dQ = U_{B} - U_{A} + dW$ $600 = U_{B} - U_{A} + 0$ or (for isochoric process dV = 0 so, dW = 0) $U_{\rm P} - U_{\rm A} = 600$...(i) For path BC, dQ = dU + dW:. $200 = U_c - U_B + p_B (V_c - V_B)$ (as BC is isobaric process) $\therefore 200 = U_c - U_B + 8 \times 10^4$ $(5 \times 10^{-3} - 2 \times 10^{-3})$ or $200 = U_{c} - U_{B} + 240$ $\therefore U_{c} - U_{B} = -40 \text{ J}$...(ii) For AC, change in internal energy can be calculated by adding Eqs. (i) and (ii), $= U_c - U_A$

$$= U_c - U_B + U_B - U_z$$

= -40 + 600 = 560 J

TOPIC 3 Heat Engine, Second Law of Thermodynamics and Carnot Engine

33 The efficiency of a Carnot engine depends upon **[NEET (Oct.) 2020]**

- (a) the temperature of the sink only
- (b) the temperatures of the source and sink
- (c) the volume of the cylinder of the engine
- (d) the temperature of the source only

Ans. (b)

Efficiency of Carnot engine is given as

$$\eta = 1 - \frac{I_2}{T_1}$$

where, T_2 = temperature of sink and T_1 = temperature of source. Hence, η depends upon the temperature of source and sink both.

34 The efficiency of an ideal heat engine working between the freezing point and boiling point of water, is [NEET 2018]
(a) 6.25% (b) 20% (c) 26.8% (d) 12.5%

Ans. (c)

Efficiency of an ideal heat engine is given as

$$\eta = 1 - \frac{T_2}{T_1}$$

where,
$$T_1$$
 is the temperature of the
source and T_2 is the temperature of the
sink.

Here,
$$T_1 = 100 + 273 = 373 \text{ K}$$

 $T_2 = 0 + 273 = 273 \text{ K}$
 $\Rightarrow \eta = 1 - \frac{273}{373}$
 $= \frac{373 - 273}{373} = \frac{100}{373} = 0.268$
∴ $\eta \% = 0.268 \times 100 = 26.8 \%$

35 A Carnot engine having an efficiency of $\frac{1}{10}$ as heat engine, is used as a refrigerator. If the work done on the system is 10 J, the amount of energy absorbed from the reservoir at lower temperature

is [NEET 2017] (a)1J (b)90J (c)99J (d)100J

Ans. (b)

Consider schematic diagram for a Carnot engine as shown below.



In case of engine,

engine efficiency = $\frac{\text{work}}{\text{heat absorbed}} = \frac{W}{q_1}$

$$\therefore \qquad \frac{W}{q_1} = \frac{1}{10}$$

$$\Rightarrow \qquad \frac{10 \text{ J}}{q_1} = \frac{1}{10} \text{ or } q_1 = 100 \text{ J}$$

When this engine is reversed, it takes in work W and heat q_2 from cold reservoir and ejects 100 J of heat to hot reservoir.

$$\therefore \quad VV + q_2 = q_1$$

 \Rightarrow 10 + q₂ = 100 or q₂ = 90 J

36 A refrigerator works between 4°C and 30°C. It is required to remove 600 calories of heat every second in order to keep the temperature of the refrigerated space constant. The power required is

(Take, 1cal=4.2Joules)

[NEET 2016]

a) 23.65 W	(b)236.5 W
c) 2365 W	(d)2.365 W

Ans. (b)

Given, temperature of source,

 $T = 30^{\circ} C = 30 + 273 \implies T_1 = 303 \text{ K}$ Temperature of sink, $T_2 = 4^{\circ} C = 4 + 273$ $T_2 = 277 \text{ K}$ As, we know that $\frac{Q_1}{Q_2} = \frac{T_1}{T_2} \Longrightarrow \frac{Q_2 + W}{Q_2} = \frac{T_1}{T_2}$ $\{\because W = Q_1 - Q_2\}$

where Q_2 is the amount of heat drawn from the sink (at T_2), W is workdone on working substance, Q_1 is amount of heat rejected to source (at room temperature T_1).

$$\Rightarrow WT_{2} + T_{2}O_{2} = T_{1}O_{2}$$

$$\Rightarrow WT_{2} = T_{1}O_{2} - T_{2}O_{2}$$

$$\Rightarrow WT_{2} = O_{2}(T_{1} - T_{2})$$

$$\Rightarrow W = O_{2}\left(\frac{T_{1}}{T_{2}} - 1\right)$$

$$\Rightarrow W = 600 \times 4.2 \times \left(\frac{303}{277} - 1\right)$$

$$W = 600 \times 4.2 \times \left(\frac{26}{277}\right)$$

$$W = 236.5 \text{ Joules}$$

$$Power = \frac{Work \text{ done}}{Time} = \frac{W}{t} = \frac{236.5}{1}$$

37 The temperature inside a refrigerator is $t_2 \,^\circ C$ and the room temperature is $t_1 \,^\circ C$. The amount of heat delivered to the room for each joule of electrical energy consumed ideally will be **[NEET 2016]**

(a)
$$\frac{t_1}{t_1 - t_2}$$
 (b) $\frac{t_1 + 273}{t_1 - t_2}$
(c) $\frac{t_2 + 273}{t_1 - t_2}$ (d) $\frac{t_1 + t_2}{t_1 + 273}$

Ans.(b)

For a refrigerator, we know that

$$\frac{Q_1}{W} = \frac{Q_1}{Q_1 - Q_2} = \frac{T_1}{T_1 - T_2}$$

where,

 $Q_1 =$ amount of heat delivered to the room

W = electrical energy consumed

 $T_1 = \text{room temperature} = t_1 + 273$

 T_2 = temperature of sink = t_2 + 273

$$\begin{array}{ll} \therefore & \frac{Q_1}{1} = \frac{t_1 + 273}{t_1 + 273 - (t_2 + 273)} \\ \Rightarrow & Q_1 = \frac{t_1 + 273}{t_1 - t_2} \end{array}$$

38 A Carnot engine, having an efficiency of $\eta = \frac{1}{10}$ as heat engine, is used as a refrigerator. If the work done on the system is 10 J, the amount of energy absorbed from the reservoir at lower temperature [CBSE AIPMT 2015] is (a) 100 J (b) 99 J (c) 90 J (d) 1 J Ans. (c) $Q_1 + W = Q_2$ As, $\eta = \frac{1}{10}$ Given, Now, using $\eta = 1 - \frac{T_1}{\tau}$ $\frac{1}{10} = 1 - \frac{T_1}{T_2} \Longrightarrow \frac{T_1}{T_2} = \frac{9}{10}$ So, $\frac{Q_1}{Q_2} = \frac{T_1}{T_2} \Rightarrow \frac{Q_1}{Q_1 + W} = \frac{9}{10}$ Now $10Q_1 = 9Q_1 + 9W$ \Rightarrow $0_1 = 9W = 9 \times 10 = 90 \text{ J}$ \Rightarrow

39 The coefficient of performance of a refrigerator is 5. If the temperature inside freezer is -20° C, the temperature of the surroundings to which it rejects heat is [CBSE AIPMT 2015] (a) 31°C (b)41°C

(c)11°C	(d)21°C

Ans. (a)

Key Concept Coefficient of performance (β) of a refrigerator is defined as the ratio of quantity of heat removed per cycle (Q_2) to the work done on the working substance per cycle to remove this heat.

Given, coefficient of performance of a refrigerator, $\beta = 5$

Temperature of surface, i.e. inside freezer,

 $T_2 = -20^{\circ} \text{C} = -20 + 273 = 253 \text{K}$

Temperature of surrounding, i.e. heat rejected outside $T_1 = ?$

So,
$$\beta = \frac{I_2}{T_1 - T_2} \implies 5 = \frac{253}{T_1 - 253}$$

 \Rightarrow 5T, -1265 = 253

$$\Rightarrow 5 I_1 = 1518$$

$$T_1 = \frac{1518}{5} = 303.6 \text{ K}$$

$$T_1 = 303.6 - 273 = 31^{\circ} \text{ C}$$
40 An engine has an efficiency of $\frac{1}{6}$.
When the temperature of sink is
reduced by 62°C, its efficiency is
doubled. Temperature of the
source is **[CBSE AIPMT 2007]**
(a) 124°C (b) 37°C (c) 62°C (d) 99°C
Ans. (d)
Efficiency of engine is given by
 $\eta = 1 - \frac{T_2}{T_1}$
 $T_2 = \text{temperature of sink}$
 $T_1 = \text{temperature of source}$
 $\therefore \frac{T_2}{T_1} = 1 - \eta = 1 - \frac{1}{6} = \frac{5}{6}$...(i)
In other case,
 $\frac{T_2 - 62}{T_1} = 1 - \eta = 1 - \frac{2}{6} = \frac{2}{3}$...(ii)
or $T_2 - 62 = \frac{2}{3}T_1 = \frac{2}{3} \times \frac{6}{5}T_2$ [Using Eq. (i)]
or $\frac{1}{5}T_2 = 62$
 $\therefore T_2 = 310 \text{ K}$
 $= 310 - 273^{\circ}\text{C} = 37^{\circ}\text{C}$
Here, $T_1 = \frac{6}{5}T_2 = \frac{6}{5} \times 310$
 $= 372 \text{ K} = 372 - 273$

4 = 4 0

41 A Carnot engine whose sink is at 300 K has an efficiency of 40%. By how much should the temperature of source be increased so as to increase its efficiency by 50% of original efficiency?

[CBSE AIPMT 2006]

(a) 275 K (b) 325 K (c) 250 K (d) 380 K

Ans. (c)

The efficiency of Carnot engine is defined as the ratio of work done to the heat supplied i.e.

$$\eta = \frac{\text{Work done}}{\text{Heat supplied}} = \frac{W}{Q_1} = \frac{Q_1 - Q_2}{Q_1}$$
$$= 1 - \frac{Q_2}{Q_2} = 1 - \frac{T_2}{T_2}$$

Here, T_1 is the temperature of source, T_2 is the temperature of sink, Q_1 is heat absorbed and Q_2 heat rejected

As given, $\eta = 40\% = \frac{40}{100} = 0.4$ and $T_2 = 300 \, \text{K}$ So $0.4 = 1 - \frac{300}{T_1}$ $\Rightarrow T_1 = \frac{300}{1 - 0.4} = \frac{300}{0.6}$ $= 500 \, \text{K}$ Let temperature of the source be

increased by x K, then efficiency becomes

$$\eta' = 40\% + 50\% \text{ of } \eta$$

$$= \frac{40}{100} + \frac{50}{100} \times 0.4$$

$$= 0.4 + 0.5 \times 0.4 = 0.6$$
Hence, $0.6 = 1 - \frac{300}{500 + x}$

$$\Rightarrow \qquad \frac{300}{500 + x} = 0.4$$

$$\Rightarrow \qquad 500 + x = \frac{300}{0.4} = 750$$

$$\therefore \qquad x = 750 - 500 = 250 \text{ K}$$

All reversible heat engines working between same temperatures are equally efficient and no heat engine can be more efficient than Carnot engine (as it is ideal).

42 An ideal gas heat engine operates in Carnot cycle between 227°C and 127°C. It absorbs 6×10^4 cal of heat at higher temperature. Amount of heat converted to work is

[CBSE AIPMT 2005]

(a)2.4×10 ⁴ cal	(b) 6×10^4 cal
(c)1.2×10 ⁴ cal	(d)4.8 $\times 10^{4}$ cal

Ans. (c)

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According to the Carnot cycle in heat engine

 $\frac{Q_2}{Q_1} = \frac{T_2}{T_1}$

Given, heat absorbed, $Q_1 = 6 \times 10^4$ cal,

Temperature of source, $T_1 = 227 + 273 = 500 \text{ K}$

Temperature of sink,

$$T_2 = 127 + 273 = 400 \text{ K}$$

 $\therefore \quad \frac{Q_2}{6 \times 10^4} = \frac{400}{500}$

 \Rightarrow Heat rejected, $Q_2 = \frac{4}{5} \times 6 \times 10^4$ $=4.8 \times 10^{4}$ cal

Now, heat converted to work

$$W = Q_1 - Q_2 = 6.0 \times 10^4 - 4.8 \times 10^4$$
$$= 1.2 \times 10^4 \text{ cal}$$

43 An ideal gas heat engine operates in a Carnot cycle between 227°C and 127°C. It absorbs 6 kcal at the higher temperature. The amount of heat (in kcal) converted into work is equal to **[CBSE AIPMT 2003]** (a) 1.6 (b) 1.2 (c) 4.8 (d) 3.5

Ans.(b)

The efficiency of heat engine is $\eta = 1 - \frac{l_2}{T}$

or
$$\frac{W}{Q_1} = 1 - \frac{T_2}{T_1}$$

 $T_2 = \text{Temperature of sink}$

 $T_1 = \text{Temperature of source}$

Given, Q_1 = heat absorbed from the source

= 6 kcal $T_1 = 227 + 273 = 500 \text{ K}$ and $T_2 = 127 + 273 = 400 \text{ K}$ Hence, $\frac{W}{6} = 1 - \frac{400}{500}$ or $\frac{W}{6} = \frac{100}{500}$ or W = 1.2 kcal

Thus, amount of heat converted into work is 1.2 kcal.

44 The efficiency of Carnot engine is 50% and temperature of sink is 500 K. If the temperature of source is kept constant and its efficiency is to be raised to 60%, then the required temperature of the sink will be [CBSE AIPMT 2002]

(a) 600 K
(b) 500 K
(c) 400 K
(d) 100 K

Ans. (c)

Efficiency of the Carnot engine is given by

$$\eta = 1 - \frac{I_2}{T_1}$$

where, T_1 = temperature of source T_2 = temperature of sink

Given,
$$\eta = 50\% = 0.5$$
, $T_2 = 500$ K

Substituting in Eq. (i), we have

$$0.5 = 1 - \frac{500}{T_1}$$
 or $\frac{500}{T_1} = 0.5$

$$\therefore T_1 = \frac{500}{0.5} = 1000 \text{ K}$$

Now, the temperature of sink is changed to T_2' and the efficiency becomes 60% i.e., 0.6.

$$0.6 = 1 - \frac{r_2}{1000}$$

or $\frac{T'_2}{1000} = 1 - 0.6 = 0.4 \text{ or}$
 $T'_2 = 0.4 \times 100 = 400 \text{ K}$

τ,

Carnot engine is not a practical engine because many ideal situations have been assumed while designing this engine which cannot be obtained practically.

45 The temperatures of source and sink of a heat engine are 127°C and 27°C respectively. An inventor claims its efficiency to be 26%, then, [CBSE AIPMT 2001]
(a) it is impossible
(b) it is possible with high probability

(c) it is possible with low probability(d) Data is insufficient

Ans. (a)

Efficiency of heat engine is, $T_{r} = T_{r} - T_{r}$

 $\eta = 1 - \frac{T_2}{T_1}$ or $\eta = \frac{T_1 - T_2}{T_1}$ $T_2 = \text{temperature of sink}$ $T_1 = \text{temperature of source}$

Given,
$$T_1 = 273 + 127 = 400 \text{ K}$$

$$T_2 = 273 + 27 = 300 \text{ K}$$

$$\therefore \qquad \eta = \frac{400 - 300}{400} = \frac{100}{400} = 0.25 = 25\%$$

Hence, 26% efficiency is impossible for a given heat engine.

46 An engine takes heat from a reservoir and converts its 1/6 part into work. By decreasing temperature of sink by 62°C, its efficiency becomes double. The temperatures of source and sink must be [CBSE AIPMT 2000]

(a) 90°C, 37°C
(b) 99°C, 37°C
(c) 372°C, 37°C
(d) 206°C, 37°C

Ans. (b)

...(i)

If T_1 is temperature of source and T_2 the temperature of sink, the efficiency of engine

$$\eta = \frac{\text{Work done}(W)}{\text{Heat taken}(Q_1)} = 1 - \frac{T_2}{T_1}$$

$$\therefore \quad 1 - \frac{T_2}{T_1} = \frac{1}{6} \qquad \dots (i)$$

When temperature of sink is reduced by 62°C, then temperature of sink

$$T'_{2} = T_{2} - 62$$

$$\therefore \qquad \eta' = 1 - \frac{T'_{2}}{T_{1}}$$

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As according to question efficiency becomes double

o,
$$\eta' = 2\eta = \frac{2}{6} = \frac{1}{3}$$

 $\frac{1}{3} = 1 - \frac{T_2 - 62}{T_1}$...(ii)

From Eq. (i)
$$\frac{T_2}{T_1} = \frac{5}{6}$$
 ...(iii)

From Eq. (ii)
$$\frac{T_2 - 62}{T_1} = \frac{2}{3}$$
 ...(iv)

Dividing Eq. (iii) by Eq. (iv)

$$\frac{T_2}{T_2 - 62} = \frac{5}{4}$$

$$\Rightarrow \qquad 4T_2 = 5T_2 - 310$$

 \Rightarrow $T_2 = 310 \text{ K}$

and from Eq. (iii), we have

$$\frac{310}{T_1} = \frac{5}{6} \implies T_1 = 372 \text{ K}$$

Hence, $T_1 = 372 \text{ K} = 372 - 273 = 99^{\circ}\text{C}$ and $T_2 = 310 \text{ K} = 310 - 273 = 37^{\circ}\text{C}$ As kinetic energy of a gas depends on its atomicity.

47 The efficiency of a Carnot engine operating between temperatures of 100°C and – 23°C will be

	[CBSE AIPMT 1997]
(a) $\frac{100-23}{277}$	(b) $\frac{100+23}{373}$
273	373 100 – 23
$(c)\frac{100120}{100}$	$(d) \frac{100 - 20}{100}$

Ans. (b)

Efficiency of Carnot engine is given by

$$\eta = 1 - \frac{T_2}{T_1} = \frac{T_1 - T_2}{T_1} \qquad \dots (i)$$

Given, T_1 = temperature of reservoir = 100 + 273 = 373 K T_2 = temperature of sink = -23 + 273 = 250 K Substituting in Eq. (i), we get

$$\therefore \quad \eta = \frac{373 - 250}{373} = \frac{123}{373}$$
$$= \frac{100 + 23}{373}$$