Ordinary Thinking

Objective Questions

First Law of Thermodynamics ($\triangle Q = \triangle U + \triangle W$)

T	1	c	.1				1
First	law	ot	thermnod	lynamics	18	given	by

[CPMT 1977, 91]

(a)
$$dQ = dU + PdV$$

(b) $dO = dU \times PdV$

(c)
$$dQ = (dU + dV)P$$

(d) dQ = PdU + dV

2. The internal energy of an ideal gas depends upon

[RPMT 1997; MP PMT 1999; CPMT 2003]

- (a) Specific volume
- (b) Pressure
- (c) Temperature
- (d) Density
- **3.** In changing the state of thermodynamics from *A* to *B* state, the heat required is *Q* and the work done by the system is W. The change in its internal energy is

[MP PMT 1986; AMU (Med.) 2001]

- (a) Q + W
- (b) Q W

(c) Q

- (d) $\frac{Q-W}{2}$
- **4.** Heat given to a system is 35 joules and work done by the system is 15 *joules*. The change in the internal energy of the system will be
 - (a) -50 J
- (b) 20 J
- (c) 30 J
- (d) 50 J
- The temperature of an ideal gas is kept constant as it expands. The gas does external work. During this process, the internal energy of the gas [MP PMT 1990]
 - (a) Decreases
 - (b) Increases
 - (c) Remains constant
 - (d) Depends on the molecular motion
- The first law of thermodynamics is concerned with the conservation of [MP PMT 1987; CBSE PMT 1990, 92;

AFMC 1997; CPMT 1999; BHU 1999; DCE 2000; BCECE 2003]

- (a) Momentum
- (b) Energy
- (c) Mass
- (d) Temperature

P, V to P, 2 V. Then work done in the two cases is

[MP PMT 1990]

- (a) Zero, Zero
- (b) Zero, PV_1
- (c) PV_1 , Zero
- (d) PV_1, P_1V_1
- **8.** If the amount of heat given to a system be 35 *joules* and the amount of work done by the system be -15 *joules*, then the change in the internal energy of the system is

[MP PMT 1989]

- (a) −50 joules
- (b) 20 joules
- (c) 30 joules
- (d) 50 joules
- 9. A system is given 300 calories of heat and it does 600 joules of work. How much does the internal energy of the system change in this process

(J = 4.18 joules/cal)

[MP PET 1991]

- (a) 654 Joule
- (b) 156.5 *Joule*
- (c) 300 Joule
- (d) 528.2 *Joule*
- 10. Work done on or by a gas, in general depends upon the
 - (a) Initial state only
 - (b) Final state only
 - (c) Both initial and final states only
 - (d) Initial state, final state and the path
- 11. If R = universal gas constant, the amount of heat needed to raise the temperature of 2 mole of an ideal monoatomic gas from 273K to 373K when no work is done [MP PET 1990]
 - (a) 100 R
- (b) 150 R
- (c) 300 R
- (d) 500 R
- 12. Find the change in internal energy of the system when a system absorbs 2 *kilocalorie* of heat and at the same time does 500 *joule* of work [EAMCET 1984]
 - (a) 7900 J
- (b) 8200 J
- (c) 5600 J
- (d) 6400 J

(b) Becomes zero

(d) Decreases

(b) 150 J

(d) 40 J

If 150 J of heat is added to a system and the work done by the

system is $110\ \emph{J}$, then change in internal energy will be

(a) Remains constant

(c) Increases

(a) 260 J

(c) 110 J

22.

[CPMT 1981;] & KCET 2004]

[AMU (Engg.) 1999; BHU 2000]

14.	A container of volume $1m^3$ is divided into two equal compartmer by a partition. One of these compartments contains an ideal gas 300 K . The other compartment is vacuum. The whole system	at is	If ΔQ and ΔW represent the heat supplied to the system and the work done on the system respectively, then the first law of thermodynamics can be written as [Roorkee 2000]				
	thermally isolated from its surroundings. The partition is remove and the gas expands to occupy the whole volume of the contain-		(a) $\Delta Q = \Delta U + \Delta W$ (b) $\Delta Q = \Delta U - \Delta W$				
	Its temperature now would be [Manipal MEE 199	_	(c) $\Delta Q = \Delta W - \Delta U$ (d) $\Delta Q = -\Delta W - \Delta U$				
	(a) 300 K (b) 239 K		where ΔU is the internal energy				
	(c) 200 K (d) 100 K	24.	For free expansion of the gas which of the following is true				
15.	110 J of heat is added to a gaseous system, whose internal ener change is 40 J , then the amount of external work done is [CBSE P.		$\label{eq:amu (Med.) 2000} \mbox{[AMU (Med.) 2000]}$ IT (996, 4) AFWC=1999and $\Delta E_{\rm int}=0$				
	JIPMER 2000; MH CET 2000; Pb. PMT 200	03]	(b) $Q=0, W>0$ and $\Delta E_{\rm int}=-W$				
	(a) 150 <i>J</i> (b) 70 <i>J</i>		(c) $W=0, Q>0$, and $\Delta E_{\rm int}=Q$				
	(c) 110 J (d) 40 J						
16.	Which of the following is not thermodynamical function		(d) $W>0, Q<0$ and $\Delta E_{\mathrm{int}}=0$				
	[CBSE PMT 1993; CPMT 2001; DCE 1996; 20	oı] 25.	Which of the following can not determine the state of a				
	(a) Enthalpy (b) Work done		thermodynamic system [AFMC 2001] (a) Pressure and volume				
	(c) Gibb's energy (d) Internal energy		(b) Volume and temperature				
17.	When the amount of work done is 333 cal and change in interr	nal	(c) Temperature and pressure				
	energy is 167 cal, then the heat supplied is		(d) Any one of pressure, volume or temperature				
	[AFMC 199	^{)8]} 26.	Which of the following is not a thermodynamics co-ordinate				
	(a) 166 <i>cal</i> (b) 333 <i>cal</i>		[AIIMS 2001]				
	(c) 500 cal (d) 400 cal		(a) P (b) T				
18.	First law thermodynamics states that [KCET 1999]		(c) V (d) R				
	(a) System can do work	27.	In a given process for an ideal gas, $dW=0$ and $dQ<0$. Then for				
	(b) System has temperature		the gas [IIT-JEE (Screening) 2001]				
	(c) System has pressure		(a) The temperature will decrease				
	(d) Heat is a form of energy		(b) The volume will increase				
19.	A thermo-dynamical system is changed from state (P_1, V_1)	to	(c) The pressure will remain constant				
	(P_2, V_2) by two different process. The quantity which will remark		(d) The temperature will increase The specific heat of hydrogen gas at constant pressure is				
	same will be [RPET 199		The specific field of hydrogen gas at constant pressure is $C_P = 3.4 \times 10^3 cal/kg$ °C and at constant volume is				
	(a) ΔQ (b) ΔW	,	$C_P = 3.4 \times 10^3 cal/ kg^{\circ}$ C. If one kilogram hydrogen gas is heated				
	(c) $\Delta Q + \Delta W$ (d) $\Delta Q - \Delta W$		from $10^{\circ}C$ to $20^{\circ}C$ at constant pressure, the external work				
20.	In thermodynamic process, 200 <i>Joules</i> of heat is given to a gas at	nd	done on the gas to maintain it at constant pressure is				
20.	100 Joules of work is also done on it. The change in internal ener	gy	(a) 10^5 cal (b) 10^4 cal				
	of the gas is [AMU (Engg.) 199	99]	(c) 10^3 cal (d) 5×10^3 cal				
	(a) 100 <i>J</i> (b) 300 <i>J</i>	29.	· .				
21	(c) 419 <i>J</i> (d) 24 <i>J</i> A perfect gas contained in a cylinder is kept in vacuum. If the cylinder is kept in vacuum.	ha	thermodynamic state of matter [CPMT 2001; AIEEE 2003]				
21.	cylinder suddenly bursts, then the temperature of the gas	iie	(a) Volume (b) Temperature				

A system performs work ΔW when an amount of heat is ΔQ

added to the system, the corresponding change in the internal

energy is ΔU . A unique function of the initial and final states

(b) ΔW

(d) ΔU

(irrespective of the mode of change) is

13.

(a) ΔQ

(c) ΔU and ΔQ

	(c) Pressure (d) Work		energy when temperature of 2 moles of this gas is increased from
30.	In a thermodynamic system working substance is ideal gas, its internal energy is in the form of [MP PMT 2003]		340 K to 342 K [RPET 1997]
	(a) Kinetic energy only		(a) 27.80 cal (b) 19.84 cal (c) 13.90 cal (d) 9.92 cal
	(b) Kinetic and potential energy	40	(c) 13.90 <i>cal</i> (d) 9.92 <i>cal</i> Temperature is a measurement of coldness or hotness of an object.
	(c) Potential energy	40.	This definition is based on [RPET 2003]
	(d) None of these		(a) Zeroth law of thermodynamics
31.	Which of the following statements is correct for any thermodynamic		(b) First law of thermodynamics
	system [AIEEE 2004]		(c) Second law of thermodynamics
	(a) The internal energy changes in all processes		(d) Newton's law of cooling
	(b) Internal energy and entropy are state functions	41.	When heat energy of 1500 Joules, is supplied to a gas at constant
	(c) The change in entropy can never be zero		pressure $2.1 \times 10^5 N/m^2$, there was an increase in its volume
	(d) The work done in an adiabatic process is always zero		equal to $2.5 \times 10^{-3} m^3$. The increase in internal energy of the gas in Joules is [EAMCET (Engg.) 1999]
32.	A system is provided with 200 cal of heat and the work done by the system on the surrounding is 40 <i>J</i> . Then its internal energy		(a) 45[Orissa PMT 2004] (b) 525
	(a) Increases by 600 J (b) Decreases by 800 J		(c) 975 (d) 2025
	(c) Increases by 800 J (d) Decreases by 50 J	42.	If heat given to a system is 6 kcal and work done is 6 kJ. Then
33.	In a thermodynamic process, pressure of a fixed mass of a gas is		change in internal energy is [BHU Med. 2000]
	changed in such a manner that the gas molecules gives out 20 <i>J</i> of heat and 10 <i>J</i> of work is done on the gas. If the initial internal energy		(a) 19.1 <i>kJ</i> (b) 12.5 <i>kJ</i>
	of the gas was 40 <i>J</i> , then the final internal energy will be		(c) 25[DPMT 2004] (d) Zero
	(a) 30 <i>J</i> (b) 20 <i>J</i>	43.	In a thermodynamics process, pressure of a fixed mass of a gas is
	(c) 60 J (d) 40 J		changed in such a manner that the gas releases 20 <i>J</i> of heat and 8, of work is done on the gas. If the initial internal energy of the gas
34.	Heat is not being exchanged in a body. If its internal energy is		was 30/. The final internal energy will be
	increased, then [RPMT 2002]		(a) 18/ (b) 9/
	(a) Its temperature will increase		(c) 4.5/ (d) 36/
	(b) Its temperature will decrease	44.	A monoatomic gas of n -moles is heated from temperature T to T
	(c) Its temperature will remain constant	• •	under two different conditions (i) at constant volume and (ii) at
~-	(d) None of these		constant pressure. The change in internal energy of the gas is
35.	Out of the following which quantity does not depend on path		(a) Mare To 2002
	(a) Temperature (b) Energy (c) Work (d) None of these		(b) More for (ii)
26			(c) Same in both cases
36.	First law of thermodynamics is a special case of [CPMT 1985; RPET 2000; DCE 2000;		(d) Independent of number of moles
	CBSE PMT 2000; AIEEE 2002; AFMC 2002]	45.	The state of a thermodynamic system is represented by
	(a) Newton's law		[MH CET 2004]
	(b) Law of conservation of energy		(a) Pressure only
	(c) Charle's law		(b) Volume only
	(d) Law of heat exchange		(c) Pressure, volume and temperature (d) Number of moles
37.	One mole of an ideal monoatomic gas is heated at a constant	46.	A perfect gas goes from state <i>A</i> to another state <i>B</i> by absorbing
	pressure of one atmosphere from $0^{o}C$ to $100^{o}C$. Then the change in the internal energy is [Pb. PMT 2001]	40.	$8 \times 10^5 J$ of heat and doing $6.5 \times 10^5 J$ of external work. It is
	(a) 6.56 <i>joules</i> (b) 8.32×10^2 <i>joules</i>		now transferred between the same two states in another process in which it absorbs $10^5 J$ of heat. Then in the second process
	(c) 12.48×10^2 joules (d) 20.80 joules		·
38.	If the ratio of specific heat of a gas at constant pressure to that at		(a) Work done on the gas is $0.5 \times 10^5 J$
	constant volume is γ , the change in internal energy of a mass of		(b) Work done by gas is $0.5 \times 10^5 J$
	gas, when the volume changes from V to $2V$ constant pressure p , is		[CBSE PMT 1998] (c) Work done on gas is $10^5 J$
	(a) $R/(\gamma-1)$ (b) pV		(d) Work done by gas is $10^5 J$
	(c) $pV/(\gamma-1)$ (d) $\gamma pV/(\gamma-1)$	47.	If a system undergoes contraction of volume then the work done by
39.	If $C_V = 4.96 cal/mole$ K, then increase in internal	***	the system will be [BHU 1999]
			(a) Zero (b) Negligible

- (c) Negative
- (d) Positive
- 48. Which of the following is incorrect regarding the first law of thermodynamics [AIEEE 2005]
 - (a) It introduces the concept of the internal energy
 - (b) It introduces the concept of the entropy
 - It is not applicable to any cyclic process
 - (d) None of the above

Isothermal Process

For an ideal gas, in an isothermal process 1.

- [BHU 1998]
- (a) Heat content remains constant
- (b) Heat content and temperature remain constant
- (c) Temperature remains constant
- (d) None of the above
- 2. Can two isothermal curves cut each other
 - (a) Never
 - (b) Yes
 - They will cut when temperature is $0^{\circ}C$
 - (d) Yes, when the pressure is critical pressure

In an isothermal expansion 3.

[KCET 2000; AFMC 2001]

- (a) Internal energy of the gas increases
- (b) Internal energy of the gas decreases
- (c) Internal energy remains unchanged
- (d) Average kinetic energy of gas molecule decreases
- In an isothermal reversible expansion, if the volume of 96 gm of oxygen at 27°C is increased from 70 litres to 140 litres, then the work done by the gas will be
 - $300 R \log_{10} 2$ (a)
- (b) $81 R \log_{e} 2$
- $900 R \log_{10} 2$
- (d) $2.3 \times 900 R \log_{10} 2$
- A vessel containing 5 litres of a gas at 0.8 m pressure is connected 5. to an evacuated vessel of volume 3 litres. The resultant pressure inside will be (assuming whole system to be isolated)
 - (a) 4/3 m
- (b) 0.5 m
- (c) 2.0 m
- (d) 3/4 m
- For an isothermal expansion of a perfect gas, the value of $\frac{\Delta P}{P}$ is 6.

[CPMT 1980]

- (a) $-\gamma^{1/2} \frac{\Delta V}{V}$

- (c) $-\gamma \frac{\Delta V}{V}$ (d) $-\gamma^2 \frac{\Delta V}{V}$
- The gas law $\frac{PV}{T}$ = constant is true for 7.

[MNR 1974; MP PMT 1984; BHU 1995, 98, 2000]

- (a) Isothermal changes only
- (b) Adiabatic changes only
- Both isothermal and adiabatic changes
- Neither isothermal nor adiabatic changes

- 8. One mole of O_2 gas having a volume equal to 22.4 litres at $0^{\circ}C$ and 1 atmospheric pressure in compressed isothermally so that its volume reduces to 11.2 litres. The work done in this process is
 - (a) 1672.5 J
- (b) 1728 /
- (c) -1728J
- (d) -1572.5 J
- If a gas is heated at constant pressure, its isothermal compressibility 9.
 - (a) Remains constant
 - (b) Increases linearly with temperature
 - (c) Decreases linearly with temperature
 - (d) Decreases inversely with temperature
- 10. Work done per mol in an isothermal change is

[RPMT 2004; BCECE 2005]

- (a) $RT \log_{10} \frac{V_2}{V_2}$
- (b) $RT \log_{10} \frac{V_1}{V_2}$
- (c) $RT \log_e \frac{V_2}{V_1}$
- (d) $RT \log_e \frac{V_1}{V_2}$
- 11. The isothermal Bulk modulus of an ideal gas at pressure P is

[CPMT 1974, 81; UPSEAT 1998; IIT 1998]

(a) P

- (b) γP
- (c) P/2
- (d) P/γ
- 12. In isothermal expansion, the pressure is determined by

[AFMC 1995]

- (a) Temperature only
- (b) Compressibility only
- Both temperature and compressibility
- (d) None of these
- The isothermal bulk modulus of a perfect gas at normal pressure is 13.
 - $1.013 \times 10^5 \, N \, / \, m^2$
- (b) $1.013 \times 10^6 N/m^2$
- (c) $1.,013 \times 10^{-11} N/m^2$
- (d) $1.013 \times 10^{11} N/m^2$
- In an isothermal change, an ideal gas obeys 14.

[MP PMT 1993]

[EAMCET 1994; CPMT 1999]

- (a) Boyle's law
- (b) Charle's law
- (c) Gaylussac law

- (d) None of the above
- In isothermic process, which statement is wrong 15.

[RPMT 1997]

- (a) Temperature is constant
- (b) Internal energy is constant
- (c) No exchange of energy
- (d) (a) and (b) are correct
- 16. An ideal gas A and a real gas B have their volumes increased from Vto 2 V under isothermal conditions. The increase in internal energy

[CBSE PMT 1993; JIPMER 2001, 02]

- (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
- The specific heat of a gas in an isothermal process is 17.

[AFMC 1998]

- (a) Infinite
- (b) Zero
- (c) Negative
- (d) Remains constant

- 18. A thermally insulated container is divided into two parts by a screen. In one part the pressure and temperature are P and T for an ideal gas filled. In the second part it is vacuum. If now a small hole is created in the screen, then the temperature of the gas will Decrease Remain same (d) None of the above (c) A container that suits the occurrence of an isothermal process
- 19. should be made of [Pb. PMT 2000]
 - (a) Copper
- (b) Glass
- (c) Wood
- (d) Cloth
- In an isothermal process the volume of an ideal gas is halved. One 20. [MP PMT 2004]
 - (a) Internal energy of the system decreases
 - (b) Work done by the gas is positive
 - (c) Work done by the gas is negative
 - (d) Internal energy of the system increases
- A thermodynamic process in which temperature T of the system 21. remains constant though other variable P and V may change, is called [Pb. PMT 2004]
 - (a) Isochoric process
- (b) Isothermal process
- (c) Isobaric process
- (d) None of these
- 22. If an ideal gas is compressed isothermally then

[RPMT 2003]

- (a) No work is done against gas
- (b) Heat is relased by the gas
- The internal energy of gas will increase
- (d) Pressure does not change
- When an ideal gas in a cylinder was compressed isothermally by a 23 piston, the work done on the gas was found to 1.5×10^4 joules. During this process about

[MP PMT 1987]

- 3.6×10^3 cal of heat flowed out from the gas
- 3.6×10^3 cal of heat flowed into the gas
- 1.5×10^4 cal of heat flowed into the gas
- 1.5×10^4 cal of heat flowed out from the gas
- When heat is given to a gas in an isothermal change, the result will 24. [MP PET 1995; RPMT 1997]
 - External work done
 - Rise in temperature
 - (c) Increase in internal energy
 - External work done and also rise in temp.
- When 1 gm of water at $0^{\circ}C$ and $1\times10^{5} N/m^{2}$ pressure is 25. converted into ice of volume $1.091 cm^2$, the external work done will be
 - (a) 0.0091 joule
- (b) 0.0182 joule
- (c) 0.0091 joule
- (d) 0.0182 joule
- 26. The latent heat of vaporisation of water is 2240 J/gm. If the work done in the process of expansion of 1 g is 168 J, then increase in internal energy is [Pb. PET 1998; CPMT 2000]
 - (a) 2408 J
- (b) 2240 J
- (c) 2072 I
- (d) 1904 /

- 27. 540 calories of heat convert 1 cubic centimeter of water at $100^{\circ} C$ into 1671 cubic centimeter of steam at $100^{\circ} C$ at a pressure of one atmosphere Then the work done against the atmospheric pressure is nearly
 - (a) 540 cal
- (b) 40 cal
- (c) Zero cal
- (d) 500 cal
- 28. One mole of an ideal gas expands at a constant temperature of 300 K from an initial volume of 10 litres to a final volume of 20 litres. The work done in expanding the gas is

 $(R = 8.31 \ || mole-K)$

[MP PMT 1995; UPSEAT 2000]

- (a) 750 joules
- (b) 1728 joules
- (c) 1500 joules
- (d) 3456 joules
- 29. A cylinder fitted with a piston contains 0.2 moles of air at temperature 27°C. The piston is pushed so slowly that the air within the cylinder remains in thermal equilibrium with the surroundings. Find the approximate work done by the system if the final volume is twice the initial volume

[BHU (Med.) 2000]

- (a) 543 J
- (b) 345 J
- (c) 453 /
- (d) 600 /
- The volume of an ideal gas is 1 litre and its pressure is equal to 30. 72cm of mercury column. The volume of gas is made 900 cm by compressing it isothermally. The stress of the gas will be
 - (a) 8 cm (mercury)
- (b) 7 *cm* (mercury)
- (c) 6 cm (mercury)
- (d) 4 cm (mercury)
- 31. During an isothermal expansion of an ideal gas

[UPSEAT 2005]

- (a) Its internal energy decreases
- (b) Its internal energy does not change
- The work done by the gas is equal to the quantity of heat absorbed by it
- (d) Both (b) and (c) are correct

Adiabatic Process

- If a cylinder containing a gas at high pressure explodes, the gas undergoes [MP PET/PMT 1988]
 - (a) Reversible adiabatic change and fall of temperature
 - (b) Reversible adiabatic change and rise of temperature
 - (c) Irreversible adiabatic change and fall of temperature
 - (d) Irreversible adiabatic change and rise of temperature
- The work done in an adiabatic change in a gas depends only on [CPMT 1971; MF 2.
 - (a) Change is pressure
- (b) Change is volume
- (c) Change in temperature
- (d) None of the above
- In adiabatic expansion
- [DPMT 1999]
- (a) $\Delta U = 0$
- $\Delta U = \text{negative}$
- (c) $\Delta U = \text{positive}$
- $\Delta W = zero$ (d)
- The pressure in the tyre of a car is four times the atmospheric pressure at 300 K. If this tyre suddenly bursts, its new temperature will be $(\gamma = 1.4)$

[RPMT 1996; MP PMT 1990]

- $300(4)^{1.4/0.4}$
- $300(2)^{-0.4/1.4}$
- (d) $300(4)^{-0.4/1.4}$

- A gas at NTP is suddenly compressed to one-fourth of its original 5. volume. If γ is supposed to be $\frac{3}{2}$, then the final pressure is
 - (a) 4 atmosphere
- (b) $\frac{3}{2}$ atmosphere
- 8 atmosphere
- (d) $\frac{1}{4}$ atmosphere
- A monoatomic gas $(\gamma = 5/3)$ is suddenly compressed to $\frac{1}{8}$ of its 6. original volume adiabatically, then the pressure of the gas will [CPMT 1976, 83; MP PMT 1994;

DPMT 1996; Roorkee 2000; KCET 2000; Pb. PMT 1999, 2001]

- (b) 8
- (d) 32 times its initial pressure
- The pressure and density of a diatomic gas $(\gamma = 7/5)$ change 7. adiabatically from (P, d) to (P, d). If $\frac{d'}{d} = 32$, then $\frac{P'}{P}$ should be [CPMT 1982; EAWCET 2001]

- (d) None of the above
- An ideal gas at $27^{\circ}C$ is compressed adiabatically to $\frac{8}{27}$ of its 8.

original volume. If $\gamma = \frac{5}{3}$, then the rise in temperature is [CPMT 1984; CBSE PMT 1999; DPMT 2000;

BHU 2001; Pb. PET 2001; UPSEAT 2002, 03; KCET 2003;]

- (a) 450 K
- (b) 375 K
- (c) 225 K
- (d) 405 K
- Two identical samples of a gas are allowed to expand (i) isothermally
 - (ii) adiabatically. Work done is [MNR 1998]
 - More in the isothermal process
 - More in the adiabatic process
 - Neither of them
 - (d) Equal in both processes
- Which is the correct statement [MP PMT 1993] 10.
 - For an isothermal change PV = constant
 - In an isothermal process the change in internal energy must be equal to the work done
 - (c) For an adiabatic change $\frac{P_2}{P_1} = \left(\frac{V_2}{V_1}\right)^{\gamma}$, where γ is the ratio of
 - (d) In an adiabatic process work done must be equal to the heat entering the system
- The slopes of isothermal and adiabatic curves are related as 11.

[CPMT 1971; BHU 1996; MH CET 1999; UPSEAT 2000; RPET 2003]

- Isothermal curve slope = adiabatic curve slope
- Isothermal curve slope = $\gamma \times$ adiabatic curve slope
- Adiabatic curve slope = $\gamma \times$ isothermal curve slope

- (d) Adiabatic curve slope = $\frac{1}{2}$ × isothermal curve slope
- Pressure-temperature relationship for an ideal gas undergoing adiabatic change is $(\gamma = C_p / C_v)$

[CPMT 1992; MP PMT 1986, 87, 94, 97; Pb. PET 1998; DCE 2001; MP PET 2001; UPSEAT 1999, 2001; AFMC 2002]

- (a) $PT^{\gamma} = \text{constant}$
- (b) $PT^{-1+\gamma} = \text{constant}$
- (c) $P^{\gamma-1}T^{\gamma} = \text{constant}$
- (d) $P^{1-\gamma}T^{\gamma} = \text{constant}$
- The amount of work done in an adiabatic expansion from temperature T to T_1 is MP PMT 1989
 - (a) $R(T-T_1)$
- (b) $\frac{R}{\gamma 1} (T T_1)$
- (d) $R(T-T_1)(\gamma-1)$
- During the adiabatic expansion of 2 moles of a gas, the internal 14. energy of the gas is found to decrease by 2 joules, the work done during the process on the gas will be equal to

[CPMT 1988]

- (b) -1 J
- (d) -2J
- The adiabatic elasticity of hydrogen gas ($\gamma = 1.4$) at NTP is

[MP PMT 1990]

- (a) $1 \times 10^5 \ N/m^2$
- (b) $1 \times 10^{-8} \ N / m^2$
- - (c) 1.4 N/m
- (d) $1.4 \times 10^5 N / m^2$
- If γ denotes the ratio of two specific heats of a gas, the ratio of slopes of adiabatic and isothermal PV curves at their point of intersection is

[NCERT 1990; MH CET 1999; MP PMT 2000]

- (a) $1/\gamma$
- (c) $\gamma 1$
- (d) $\gamma + 1$
- Air in a cylinder is suddenly compressed by a piston, which is then 17. maintained at the same position. With the passage of time [NCERT 1971; DPMT 1

KCET 2000; AllMS 2000; MH CET 2001]

- The pressure decreases
- (b) The pressure increases
- The pressure remains the same
- The pressure may increase or decrease depending upon the nature of the gas
- When a gas expands adiabatically [CPMT 1990]
 - (a) No energy is required for expansion
 - Energy is required and it comes from the wall of the container
 - (c) Internal energy of the gas is used in doing work
 - (d) Law of conservation of energy does not hold
- One gm mol of a diatomic gas $(\gamma = 1.4)$ is compressed adiabatically so that its temperature rises from $27^{\circ}C$ to $127^{\circ}C$. The work done will be
 - (a) 2077.5 joules
- (b) 207.5 joules
- (c) 207.5 ergs
- (d) None of the above

- Compressed air in the tube of a wheel of a cycle at normal 20. temperature suddenly starts coming out from a puncture. The air inside [NCERT 1970]
 - (a) Starts becoming hotter
 - (b) Remains at the same temperature
 - (c) Starts becoming cooler
 - May become hotter or cooler depending upon the amount of water vapour present
 - The adiabatic Bulk modulus of a perfect gas at pressure is given by [CPMT 1982; MH CET 200] o o

2P

- (a) P
- γP (c) P/2(d)
- 22. An adiabatic process occurs at constant

[MNR 1985; AFMC 1996; AIIMS 1999; UPSEAT 1999, 2000; Pb. PET 2004]

- (a) Temperature
- (b) Pressure
- (c) Heat

21.

- Temperature and pressure
- A polyatomic gas $\left(\gamma = \frac{4}{3}\right)$ is compressed to $\frac{1}{8}$ of its volume 23.

adiabatically. If its initial pressure is P_o , its new pressure will be [MP PET 1994; BHU c 1995] $(C_P + C_V)(T_i - T_f)$ (d) $(C_P - C_V)(T_i - T_f)$

- (a) $8P_a$
- (b) $16P_0$
- (c) $6P_a$
- (d) $2P_{o}$
- For adiabatic processes $\left(\gamma = \frac{C_p}{C_v} \right)$ 24.

[KCET 1999; MP PET 1995; CPMT 2003]

- (a) $P^{\gamma}V = \text{constant}$
- (b) $T^{\gamma}V = \text{constant}$
- (c) $TV^{\gamma-1}$ =constant
- (d) $TV^{\gamma} = \text{constant}$
- An ideal gas is expanded adiabatically at an initial temperature of 25. 300 K so that its volume is doubled. The final temperature of the hydrogen gas is $(\gamma = 1.40)$

[MP PMT 1995; DPMT 1999]

- (a) 227.36 K
- (b) 500.30 K
- (c) 454.76 K
- (d) $-47^{\circ} C$
- A given system undergoes a change in which the work done by the 26. system equals the decrease in its internal energy. The system must have undergone an

[Haryana CEE 1996; UPSEAT 2003]

- (a) Isothermal change
- (b) Adiabatic change
- (c) Isobaric change
- (d) Isochoric change
- During the adiabatic expansion of 2 moles of a gas, the internal 27. energy was found to have decreased by 100 J. The work done by the gas in this process is [MP PET 1996, 97]
 - (a) Zero
- (b) -100 /
- (c) 200 J
- (d) 100 J
- 28. In an adiabatic expansion of a gas initial and final temperatures are T_1 and T_2 respectively, then the change in internal energy of the gas is [MP PET 1997]

 - (a) $\frac{R}{v-1}(T_2-T_1)$ (b) $\frac{R}{v-1}(T_1-T_2)$

- (c) $R(T_1 T_2)$
- (d) Zero
- Helium at $27^{\circ}C$ has a volume of 8 *litres*. It is suddenly 29. compressed to a volume of 1 litre. The temperature of the gas will be $[\gamma = 5 / 3]$

[CBSE PMT 1993; MP PMT 1999; Pb. PMT 2002]

- (a) $108^{\circ} C$
- 9327°C
- - 927° C
- A cycle tyre bursts suddenly. This represents an

[SCRA 1994]

- (a) Isothermal process
- (b) Isobaric process
- (c) Isochoric process
- (d) Adiabatic process
- One mole of helium is adiabatically expanded from its initial state 31. (P_i, V_i, T_i) to its final state (P_f, V_f, T_f) . The decrease in the internal energy associated with this expansion is equal to

SCRA 1994: BHU 20021

[RPMT 1997]

- (a) $C_V(T_i T_f)$
- (b) $C_P(T_i T_f)$

- 32. At N.T.P. one mole of diatomic gas is compressed adiabatically to half of its volume $\gamma = 1.41$. The work done on gas will be
 - (a) 1280 J
- (b) 1610 J
- (c) 1815 /
- (d) 2025 /
- For adiabatic process, wrong statement is 33.

 - (a) dQ = 0
- (b) dU = -dW
- (c) Q = constant
- (d) Entropy is not constant
- A diatomic gas initially at 18°C is compressed adiabatically to oneeighth of its original volume. The temperature after compression will be

[Pb. PET 1995; CBSE PMT 1996; CPMT 1999]

- (a) $10^{\circ} C$
- (b) $887^{\circ} C$
- (c) 668K
- (d) 144° C
- A gas is being compressed adiabatically. The specific heat of the gas 35. during compression is [SCRA 1996]
 - (a) Zero
- (b) Infinite
- (c) Finite but non-zero
- (d) Undefined
- 36. The process in which no heat enters or leaves the system is termed [Pb. PET 1996; BHU 1998; BCECE 2003]
 - (a) Isochoric
- (b) Isobaric
- (c) Isothermal
- (d) Adiabatic
- 37. Two moles of an ideal monoatomic gas at 27° C occupies a volume of V. If the gas is expanded adiabatically to the volume 2V, the work done by the gas will be $[\gamma=5/3,R=8.31J/mol\,K]$ [RPET 1999]
 - -2767.23J
- (b) 2767.23J
- 2500J
- (d) -2500J

38.	At $27^{\circ}C$ a gas is suddenly compressed such that its pressure
	becomes $\frac{1}{8}th$ of original pressure. Temperature of the gas will be

 $(\gamma = 5/3)$

[BHU 2000]

- (a) 420K
- (b) $327^{\circ} C$
- (c) 300K
- (d) $-142^{\circ} C$

 $\Delta U + \Delta W = 0$ is valid for 39.

[RPMT 2000]

- (a) Adiabatic process
- (b) Isothermal process

- (c) Isobaric process
- (d) Isochoric process
- An ideal gas at a pressures of 1 atmosphere and temperature of 40. 27°C is compressed adiabatically until its pressure becomes 8 times the initial pressure, then the final temperature is $(\gamma = 3/2)$ [EAMCET (Engs.)2000]
 - (a) $627^{\circ} C$
- (b) 527° C
- (c) $427^{\circ} C$
- (d) $327^{\circ} C$
- Air is filled in a motor tube at $27^{\circ}C$ and at a pressure of 8 41. atmospheres. The tube suddenly bursts, then temperature of air is [Given γ of air = 1.5] [MP PMT 2002]
 - (a) 27.5° C
- (b) 75° K
- (c) 150 K
- (d) $150^{\circ} C$
- If $\gamma = 2.5$ and volume is equal to $\frac{1}{8}$ times to the initial volume then pressure P' is equal to (Initial pressure = P)

[RPET 2003]

- (a) P' = P
- (c) $P' = P \times (2)^{15/2}$
- (d) P' = 7P
- In an adiabatic process, the state of a gas is changed from 43. P_1, V_1, T_1 , to P_2, V_2, T_2 . Which of the following relation is [Orissa JEE 2003]

 - (a) $T_1 V_1^{\gamma 1} = T_2 V_2^{\gamma 1}$ (b) $P_1 V_1^{\gamma 1} = P_2 V_2^{\gamma 1}$
 - (c) $T_1 P_1^{\ \gamma} = T_2 P_2^{\ \gamma}$
- (d) $T_1 V_1^{\gamma} = T_2 V_2^{\gamma}$
- During an adiabatic process, the pressure of a gas is found to be 44. proportional to the cube of its absolute temperature. The ratio C_n / C_v for the gas is [AIEEE 2003]

(c) 2

- 45. In adiabatic expansion of a gas

[BCECE 2001; MP PET 2003]

- (a) Its pressure increases
- (b) Its temperature falls
- (c) Its density increases
- (d) Its thermal energy increases
- One mole of an ideal gas at an initial temperature of TK does 6 R 46. 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 PMT 2004]

- (a) (T+2.4)K
- (b) (T-2.4)K

(c)
$$(T+4)K$$

- (d) (T-4)K
- A gas is suddenly compressed to 1/4 th of its original volume at 47. normal temperature. The increase in its temperature is $(\gamma = 1.5)$
 - (a) 273 K
- (b) 573 K (d) 473 K
- (c) 373 K
- A gas ($\gamma = 1.3$) is enclosed in an insulated vessel fitted with insulating piston at a pressure of $10^5 N/m^2$. On suddenly pressing the piston the volume is reduced to half the initial volume.

The final pressure of the gas is [RPET 2002]

- (a) $2^{0.7} \times 10^5$
- (b) $2^{1.3} \times 10^5$
- (c) $2^{1.4} \times 10^5$
- (d) None of these

[MP PMT 1989; RPMT 2001]

- (a) Adiabatic expansion
- (b) Adiabatic compression
- (c) Isothermal expansion
- (d) Isothermal compression

50. 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 PMT 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
- A gas is suddenly compressed to one fourth of its original volume. What will be its final pressure, if its initial pressure is P

[Pb. PET 2002]

- (a) Lesss than P
- (b) More than P

- (d) Either (a) or (c)
- A gas for which $\gamma = 1.5$ is suddenly compressed to $\frac{1}{4}$ th of the

initial volume. Then the ratio of the final to the initial pressure is [EAMCET 200

- (a) 1:16
- (b) 1:8
- (c) 1:4
- (d) 8:1
- One mole of an ideal gas with $\gamma = 1.4$, is adiabatically compressed 53. so that its temperature rises from 27°C to 35°C. The change in the internal energy of the gas is (R = 8.3 J/mol.K)[EAMCET 2001]
 - (a) -166 /
- (b) 166 /
- (c) -168 J
- (d) 168 J
- The volume of a gas is reduced adiabatically to $\frac{1}{4}$ of its volume at 54.

 27° C, if the value of $\gamma = 1.4$, then the new temperature will be

- (a) $350 \times 4^{0.4} K$
- (b) $300 \times 4^{0.4} K$
- (c) $150 \times 4^{0.4} K$
- (d) None of these
- During an adiabatic expansion of 2 moles of a gas, the change in 55. internal energy was found -50/. The work done during the process [Pb. PET 1996]
 - (a) Zero
- (b) 100/
- (d) 50/
- Adiabatic modulus of elasticity of a gas is $2.1 \times 10^5 \, N/m^2$. What 56.

will be its isothermal modulus of elasticity $\left(\frac{C_p}{C} = 1.4\right)$

[UPSEAT 1999]

- (a) $1.8 \times 10^5 \, N / m^2$
- (b) $1.5 \times 10^5 N/m^2$
- (c) $1.4 \times 10^5 \, N / m^2$
- (d) $1.2 \times 10^5 N/m^2$
- For an adiabatic expansion of a perfect gas, the value of $\frac{\Delta P}{P}$ is 57.

[CPMT 1983; MP PMT 1990]

(a)
$$-\sqrt{\gamma} \frac{\Delta V}{V}$$

(b)
$$-\frac{\Delta V}{V}$$

(c)
$$-\gamma \frac{\Delta V}{V}$$

(d)
$$-\gamma^2 \frac{\Delta V}{V}$$

Isobaric and Isochoric Processes

A gas expands under constant pressure P from volume V_1 to V_2 . The work done by the gas is

[CBSE PMT 1990; RPMT 2003]

- (a) $P(V_2 V_1)$
- (b) $P(V_1 V_2)$
- (c) $P(V_1^{\gamma} V_2^{\gamma})$
- (d) $P \frac{V_1 V_2}{V_2 V_1}$
- 2. When heat in given to a gas in an isobaric process, then

[DPMT 2001]

- (a) The work is done by the gas
- (b) Internal energy of the gas increases
- (c) Both (a) and (b)
- (d) None from (a) and (b)
- One mole of a perfect gas in a cylinder fitted with a piston has a 3. pressure P_i volume V and temperature T_i . If the temperature is increased by 1 K keeping pressure constant, the increase in volume

- (d) V
- A gas is compressed at a constant pressure of $50N/m^2$ from a 4 volume of $10m^3$ to a volume of $4m^3$. Energy of 100 / then added to the gas by heating. Its internal energy is

[MNR 1994]

- (a) Increased by 400 /
- (b) Increased by 200 /
- (c) Increased by 100 /
- (d) Decreased by 200 /
- Work done by air when it expands from 50 litres to 150 litres at a 5. constant pressure of 2 atmosphere is
 - (a) 2×10^4 joules
- (b) 2×100 joules
- (c) $2 \times 10^5 \times 100$ joules
- (d) $2 \times 10^{-5} \times 100$ joules
- Work done by 0.1 mole of a gas at $27^{\circ}C$ to double its volume at 6. constant pressure is $(R = 2 \text{ cal mol}^{\perp}C)$

[EAMCET 1994]

- (a) 54 cal
- (b) 600 cal
- (c) 60 cal
- (d) 546 cal
- Unit mass of a liquid with volume V_1 is completely changed into a 7. gas of volume V_2 at a constant external pressure \emph{P} and temperature T. If the latent heat of evaporation for the given mass is L, then the increase in the internal energy of the system is

- (a) Zero
- (b) $P(V_2 V_1)$
- (c) $L P(V_2 V_1)$
- (d) L
- A gas expands $0.25m^3$ at constant pressure $10^3 N/m^2$, the work done is

[CPMT 1997; UPSEAT 1999; JIPMER 2001, 02]

- (a) 2.5 ergs
- (b) 250 J
- (c) 250 W
- (d) 250 N
- Two kg of water is converted into steam by boiling at atmospheric 9. pressure. The volume changes from $2 \times 10^{-3} m^3$ to $3.34 m^3$. The work done by the system is about
 - (a) $-340 \, kJ$
- (b) $-170 \, kJ$
- (c) 170 kJ
- (d) 340 kJ
- 10. An ideal gas has volume V_0 at 27° C. It is heated at constant pressure so that its volume becomes $2V_0$. The final temperature is [BCECE 2003]
 - (a) $54^{\circ} C$
- (b) 32.6° C
- (c) 327 °C
- (d) 150 K
- If 300 ml of a gas at $27^{\circ}C$ is cooled to $7^{\circ}C$ at constant 11. pressure, then its final volume will be

[Pb. PET 1999; BHU 2003; CPMT 2004]

- (a) 540 ml
- (b) 350 ml
- (c) 280 ml
- (d) 135 ml
- Which of the following is correct in terms of increasing work done for the same initial and final state [RPMT 1996]
 - (a) Adiabatic < Isothermal < Isobaric
 - (b) Isobaric < Adiabatic < Isothermal
 - (c) Adiabatic < Isobaric < Isothermal
 - (d) None of these
- A sample of gas expands from volume V_1 to V_2 . The amount of work 13. done by the gas is greatest when the expansion is

[CBSE PMT 1997; AlIMS 1998; JIPMER 2000]

- (a) Isothermal
- (b) Isobaric
- (c) Adiabatic
- (d) Equal in all cases
- Which of the following is a slow process
- [] & K CET 2000]

- (a) Isothermal
- (b) Adiabatic
- (c) Isobaric
- (d) None of these
- How much work to be done in decreasing the volume of and ideal 15. gas by an amount of $2.4 \times 10^{-4} m^3$ at normal temperature and constant normal pressure of $1 \times 10^5 N/m^2$

[UPSEAT 1999]

- (a) 28 joule
- (b) 27 joule
- (c) 25 joule
- (d) 24 joule
- 16. A Container having 1 mole of a gas at a temperature 27°C has a movable piston which maintains at constant pressure in container of 1 atm. The gas is compressed until temperature becomes 127° C. The work done is (C for gas is 7.03 cal|molK)
 - (a) 703 J
- (b) 814 1
- (c) 121 J
- (d) 2035 /
- In a reversible isochoric change 17.
- [NCERT 1990]

[Roorkee 1999]

18. Entropy of a thermodynamic system does not change when this system is used for [AIIMS 1995] Irreversible process is 5. (a) Conduction of heat from a hot reservoir to a cold reservoir (a) Adiabatic process (b) Conversion of heat into work isobarically (b) Joule-Thomson expansion (c) Conversion of heat into internal energy isochorically (c) Ideal isothermal process (d) Conversion of work into heat isochorically The work done in which of the following processes is zero (d) None of the above 19. [UPSEAT 2003] For a reversible process, necessary condition is (a) Isothermal process (b) Adiabatic process energy should be zero (c) Isochoric process (d) None of these That the process should be too fast In which thermodynamic process, volume remains same 20. That the process should be slow so that the working substance [Orissa PMT 2004] (b) Isothermal (a) Isobaric surroundings (c) Adiabatic (d) Isochoric In an isochoric process if $T_1 = 27^{\circ} C$ and $T_2 = 127^{\circ} C$, then In a cyclic process, work done by the system is 21. P_1 / P_2 will be equal to (b) Equal to heat given to the system (a) 9 / 59 (b) 2/3 (c) More than the heat given to system (c) 3/4 (d) None of these (d) Independent of heat given to the system Which is incorrect [DCE 2001] 22. (a) In an isobaric process, $\Delta p = 0$

[RPMT 2001; BCECE 2003]

(b) $\Delta Q = 0$

(d) $\Delta U = 0$

(b) In an isochoric process, $\Delta W = 0$

(c) In an isothermal process, $\Delta T = 0$

(d) In an isothermal process, $\Delta Q = 0$

Which relation is correct for isometric process

(a) $\Delta Q = \Delta U$

23.

(a) $\Delta W = 0$

(c) $\Delta T = 0$

(b) $\Delta W = \Delta U$

(c) $\Delta Q = \Delta W$

(d) None of these

Heat Engine, Refrigerator and Second Law of Thermodynamics

A Carnot engine working between 300 K and 600 K has work output of 800 / per cycle. What is amount of heat energy supplied to the engine from source per cycle

[DPMT 1999; Pb. PMT 2002, 05; Kerala PMT 2004]

(a) 1800 //cycle

(b) 1000 //cycle

(c) 2000 //cycle

- (d) 1600 //cycle
- The coefficient of performance of a Carnot refrigerator working between $30^{\circ} C$ and $0^{\circ} C$ is [UPSEAT 2002]

(a) 10

(c) 9

- (b) 1 (d) 0
- If the door of a refrigerator is kept open, then which of the [DPMT 2001; BHU 2001; following is true

IIPMER 2002; AIEEE 2002; CPMT 2003]

- (a) Room is cooled
- Room is heated
- Room is either cooled or heated
- Room is neither cooled nor heated

- In a cyclic process, the internal energy of the gas
 - (a) Increases
- (b) Decreases
- (c) Remains constant
- (d) Becomes zero

- In the whole cycle of the system, the loss of any type of heat
- should remain in thermal and mechanical equilibrium with the
- The loss of energy should be zero and it should be quasistatic
- [BHU 2002]
- 8. An ideal gas heat engine operates in a Carnot's cycle between $227^{\,o}\,C\,$ and $\,127^{\,o}\,C\,.$ It absorbs 6 \times 10 $^{\circ}\textit{J}$ at high temperature. The amount of heat converted into work is

[KCET 2004]

(a) $4.8 \times 10^4 J$

(b) $3.5 \times 10^4 J$

(c) $1.6 \times 10^4 J$

(d) $1.2 \times 10^4 J$

- An ideal heat engine exhausting heat at $77^{\circ}C$ is to have a 30% 9. efficiency. It must take heat at [BCECE 2004]
 - (a) $127^{\circ} C$
- (b) 227° C
- (c) $327^{\circ} C$
- $673^{\circ} C$ (d)
- Efficiency of Carnot engine is 100% if [Pb. PET 2000]
 - (a) $T_2 = 273 \ K$
- (b) $T_2 = 0 K$
- (c) $T_1 = 273 K$
- (d) $T_1 = 0 \ K$
- A Carnot's engine used first an ideal monoatomic gas then an ideal diatomic gas. If the source and sink temperature are $411^{\circ} C$ and $69^{o}\,C$ respectively and the engine extracts 1000 J of heat in each cycle, then area enclosed by the PV diagram is
 - (a) 100 J
- (b) 300 J
- (c) 500 J
- (d) 700 J
- 12. A Carnot engine absorbs an amount Q of heat from a reservoir at an abosolute temperature T and rejects heat to a sink at a temperature of T/3. The amount of heat rejected is

[UPSEAT 2004]

- (a) Q/4
- (b) Q/3
- (c) Q/2
- (d) 2Q/3
- 13. The temperature of sink of Carnot engine is 27° C. Efficiency of engine is 25%. Then temperature of source is

(a) $227^{\circ} C$

(c) 127° C

(a) 300 K

(c) 500 K

	(c) 300 A	(d) 700 K	25	A Compet angine takes	$3 \times 10^6 \ cal$. of heat from a reservoir	-+
15.	In a Carnot engine, when	$T_2 = 0^o C$ and $T_1 = 200^o C$, its	25.		sink at 27° <i>C</i> . The work done by the engi	
	efficiency is η_1 and when T_1	$=0~^{o}C~$ and $T_{2}=-200~^{o}C$, Its		is [AIEEE 2003]	20 2, eee 20e e. g .	
	efficiency is η_2 , then what is η_2	η_1/η_2 [DCE 2004]		(a) $4.2 \times 10^6 J$	(b) $8.4 \times 10^6 J$	
	(a) 0.577	(b) 0.733		(c) $16.8 \times 10^6 J$	(d) Zero	
	(c) 0.638	(d) Can not be calculated	26.	The first operation involv	ed in a Carnot cycle is	
16.	The efficiency of Carnot's en	ngine operating between reservoirs,			[AFMC 199	8]
	maintained at temperatures 27	$7^{\circ} C$ and $-123^{\circ} C$, is	[1	OPMT 2002; 03; 18V 2004] sion	n (b) Adiabatic expansion	
	(a) 50%	(b) 24%		(c) Isothermal compress	sion (d) Adiabatic compression	
	(c) 0.75%	(d) 0.4%	27.	For which combination Carnot's engine is highest	of working temperatures the efficiency (KCET 200	
17.	- ,	where $227^{\circ} C$ and $27^{\circ} C$. Efficiency		(a) 80 K, 60 K	(b) 100 K, 80 K	
	of the engine will be	[DCE 1999; BHU 2004]		(c) 60 K, 40 K	(d) 40 K, 20 K	
	(a) $\frac{1}{3}$	(b) $\frac{2}{5}$	28.	The efficiency of Carnot sink temperature is <i>T</i> wil	engine when source temperature is T at I be [DCE 2000]	nd
	(c) $\frac{3}{4}$	(d) $\frac{3}{5}$		(a) $\frac{T_1 - T_2}{T_1}$	(b) $\frac{T_2 - T_1}{T_2}$	
18.	A measure of the degree of disc	order of a system is known as			2	
	3	Pb. PET 1997; MH CET 1999]		(c) $\frac{T_1 - T_2}{T_2}$	(d) $\frac{T_1}{T_2}$	
	(a) Isobaric	(b) Isotropy		2	2	
	(c) Enthalpy	(d) Entropy	29.	· ·	king between temperature <i>T</i> and <i>T</i> has	
19.	A carnot engine has the same and $x K$ to 600 K . The value of	efficiency between 800 <i>K</i> to 500 <i>K x</i> is		efficiency η , the new temperature are doubled,	efficiency if both the source and si will be [DPMT 2000]	nk
		[Pb. PMT 1996; CPMT 1996]		(a) $\frac{\eta}{2}$	(b) η	
	(a) 1000 K	(b) 960 K		2	(6) 1	
	(c) 846 K	(d) 754 K		(c) 2η	(d) 3η	
20.		ncy of his heat engine which operates and sink temperature $27^{\circ}C$ is 26% ,	30.		a freezer at a temperature of $-13^{\circ}C$. The of the engine is 5. The temperature of the temperature of the temperature of the content of th	
	(a) It is impossible				[BHU 2000; CPMT 200)2]
	(b) It is possible but less prob	pable		(a) 325° C	(b) 325 <i>K</i>	
	(c) It is quite probable			(c) 39° <i>C</i>	(d) 320° <i>C</i>	
	(d) Data are incomplete		31.		rator, the low temperature coils are at nd the compressed gas in the condenser h	
21.		ork between 200° C and 0° C first and The ratio of efficiencies of the engine			he theoretical coefficient of performance [UPSEAT 2001]	
	in the two cases is	[KCET 2002]		(a) 5	(b) 8	
	(a) 1.73:1	(b) 1:1.73		(c) 6	(d) 6.5	
	(c) 1:1	(d) 1:2	32.		to operate between two reservoirs 227° C. The maximum possible efficiency	
22.		s 50% when temperature of outlet is		such an engine is	[UPSEAT 2005]	
		se efficiency up to 60% keeping	m1	(a) 1/2	(b) 1/4	
	()	what is temperature of outlet [CBSE PM'	1 2002]	(c) 3/4	(d) 1	
	(a) 200 <i>K</i> (c) 600 <i>K</i>	(b) 400 <i>K</i> (d) 800 <i>K</i>	33.	-	e operates in Carnot cycle between 227	
23.	· /	2 100% efficiency because we cannot[AIEE	E 20021	and 127° C. It absorbs 6 Amount of heat converted	$ imes 10^4$ cals of heat at higher temperatured to work is	re.
	(a) Prevent radiation			I made converted	[CBSE PMT 200)5]
	(b) Find ideal sources			(a) $2.4 \times 10^4 \ cal$	(b) 6×10^4 cal	- 4
	,			\-/	(-)	

[DCE 2002; CPMT 2002]

(b) 327° C

(d) 27° C

(b) 400 K

(d) 700 K

The temperature of reservoir of Carnot's engine operating with an

efficiency of 70% is $1000\,\mbox{\it K}$. The temperature of its sink is

(c) Reach absolute zero temperature

(a) Second law of thermodynamics

(b) Concevation of momentum

(d) First law of thermodynamics

(c) Conservation of mass

"Heat cannot by itself flow from a body at lower temperature to a

body at higher temperature" is a statement or consequence of [AIEEE 2003, EAA

(d) Eliminate friction

24.

- 1.2×10^4 cal
- 34 Which of the following processes is reversible

[CBSE PMT 2005]

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

Critical Thinking Objective Questions

When an ideal diatomic gas is heated at constant pressure, the fraction of the heat energy supplied which increases the internal energy of the gas, is

[IIT 1990: UPSEAT 1998: RPET 2000]

- $1cm^3$ of water at its boiling point absorbs 540 calories of heat to 2. become steam with a volume of 1671cm3.If the atmospheric pressure = $1.013x10^5 N/m^2$ and the mechanical equivalent of heat = 4.19 J/calorie, the energy spent in this process in overcoming intermolecular forces is

[MP PET 1999, 2001; Orissa JEE 2002]

- (a) 540 cal
- (b) 40 cal
- (c) 500 cal
- (d) Zero
- 3. During the melting of a slab of ice at 273 K at atmospheric pressure
 - (a) Positive work is done by ice-water system on the atmosphere
 - Positive work is done on the ice-water system by the atmosphere
 - The internal energy of the ice-water system increases
 - (d) The internal energy of the ice-water system decreases
- Two identical containers A and B with frictionless pistons contain the same ideal gas at the same temperature and the same volume V. The mass of the gas in A is m_A and that in B is m_B . The gas in each cylinder is now allowed to expand isothermally to the same final volume 2V. The changes in the pressure in A and B are found to be ΔP and 1.5 ΔP respectively. Then
 - (a) $4m_A = 9m_B$
- (b) $2m_A = 3m_B$
- (c) $3m_A = 2m_B$
- (d) $9m_A = 3m_B$
- A monoatomic ideal gas, initially at temperature T_1 , is enclosed in a cylinder fitted with a frictionless piston. The gas is allowed to expand adiabatically to a temperature. T_2 by releasing the piston suddenly. If L_1 and L_2 are the lengths of the gas column before and after expansion respectively, then $\,T_1\,/\,T_2\,$ is given by

[IIT-JEE (Screening) 2000]

(a)
$$\left(\frac{L_1}{L_2}\right)^{2/3}$$

(c)
$$\frac{L_2}{L_1}$$

(d)
$$\left(\frac{L_2}{L_1}\right)^{2/2}$$

A closed hollow insulated cylinder is filled with gas at 0^{o} C and 6. also contains an insulated piston of negligible weight and negligible thickness at the middle point. The gas on one side of the piston is heated to 100° C. If the piston moves $5 \, cm$, the length of the hollow cylinder is [EAMCET 2001]

- (a) 13.65 cm
- (b) 27.3 cm
- (c) 38.6 cm
- (d) 64.6 cm
- A mono atomic gas is supplied the heat Q very slowly keeping the 7. pressure constant. The work done by the gas will be

- 8. A gas mixture consists of 2 moles of oxygen and 4 moles argon at temperature T. Neglecting all vibrational modes, the total internal energy of the system is

[IIT 1999; UPSEAT 2003]

- (a) 4 RT
- (b) 15 RT
- (c) 9 RT
- (d) 11 RT
- An ideal gas expands isothermally from a volume V_1 to V_2 and then compressed to original volume V_1 adiabatically. Initial pressure is P_1 and final pressure is P_3 . The total work done is $\it W$. Then
 - (a) $P_3 > P_1, W > 0$
- (b) $P_3 < P_1, W < 0$
- (c) $P_{[1]} = P_{[2]} = W < 0$
- (d) $P_3 = P_1, W = 0$
- 10. Work done by a system under isothermal change from a volume V_1 V_2 for a gas which obeys Vander Waal's equation

$$(V - \beta n) \left(P + \frac{\alpha n^2}{V} \right) = nRT$$

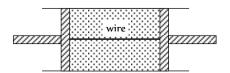
(a)
$$nRT \log_e \left(\frac{V_2 - n\beta}{V_1 - n\beta} \right) + \alpha n^2 \left(\frac{V_1 - V_2}{V_1 V_2} \right)$$

$$\text{(b)} \quad nRT \log_{10} \left(\frac{V_2 - \alpha\beta}{V_1 - \alpha\beta} \right) + \alpha \, n^2 \left(\frac{V_1 - V_2}{V_1 V_2} \right)$$

(c)
$$nRT \log_e \left(\frac{V_2 - n\alpha}{V_1 - n\alpha} \right) + \beta n^2 \left(\frac{V_1 - V_2}{V_1 V_2} \right)$$

(d)
$$nRT \log_e \left(\frac{V_1 - n\beta}{V_2 - n\beta} \right) + \alpha n^2 \left(\frac{V_1 V_2}{V_1 - V_2} \right)$$

A cylindrical tube of uniform cross-sectional area A is fitted with 11. two air tight frictionless pistons. The pistons are connected to each other by a metallic wire. Initially the pressure of the gas is P and temperature is T, atmospheric pressure is also P. Now the temperature of the gas is increased to 2T, the tension in the wire



work of $\frac{Q}{4}$ when a heat of Q is supplied to it is

The molar heat capacity in a process of a diatomic gas if it does a

(b) $\frac{5}{2}R$

(d) $\frac{6}{7}R$

An insulator container contains 4 moles of an ideal diatomic gas at temperature T. Heat Q is supplied to this gas, due to which 2 moles

of the gas are dissociated into atoms but temperature of the gas

then what percentage of given heat energy will be utilised in doing

(b) 30 %

(a) $2P_0A$

(c) $\frac{10}{3}R$

remains constant. Then

12.

13.

14.

15.

16.

17.

18.

external work

(a) 40 %

(c) $\frac{P_0 A}{2}$

(a) $Q = 2RT$	(b) $Q = RT$		(a) 1 <i>P</i>	(b)	2 <i>P</i>		
(c) $Q = 3RT$	(d) $Q = 4RT$		(c) 4 <i>P</i>	` '	8 <i>P</i>		
	s by 5% in its adiabatic expansion. The	22.	A thermally insulated rigid container contains an ideal gas heated b a filament of resistance 100 Ω through a current of 1 A for 5 mi then change in internal energy is				
(a) 5%	(b) 6%				[IIT-JEE (Screening	g) 2005]	
(c) 7%	(d) 8%		(a) 0 <i>kJ</i>	(b)	10 <i>kJ</i>		
The temperature of a hypot	thetical gas increases to $\sqrt{2}$ times when		(c) 20 <i>kJ</i>	(d)	30 <i>kJ</i>		
	half the volume. Its equation can be	23.	A reversible engine conv When the temperature o of the engine is doubled	of the sink is	reduced by $62^{\circ}C$, the eff	ficiency	
(a) $PV^{3/2} = \text{constant}$	(b) $PV^{5/2} = constant$		are		[CBSE PMT 2000]		
(c) $PV^{7/3} = \text{constant}$	(d) $PV^{4/3} = \text{constant}$		(a) 80°C, 37°C	(b)	95° <i>C</i> , 28° <i>C</i>		
` ,	<i>B</i> are operated in succession. The first		(c) 90°C, 37°C	(d)	99° <i>C</i> , 37° <i>C</i>		
one, A receives heat from	a source at $T_1 = 800K$ and rejects to ngine B receives heat rejected by the first	24.	An engineer claims to ha	f 1 <i>g/sec</i> . The		•	
-	ther sink at $T_3 = 300 K$. If the work		<i>kcal/g</i> . Is the claim of the	engineer			
	equal, then the value of T_2 is		()		[] & K CE	Γ 2000]	
			(a) Valid				
(a) 100 <i>K</i>	(b) 300 <i>K</i>		(b) Invalid				
(c) 550 <i>K</i>	(d) 700 <i>K</i>		(c) Depends on engine	design			
	ic gas is heated at constant pressure, plied which increases the internal energy		(d) Depends of the load	ł			
of gas, is	[AliMS 1995]	25.	Find the change in the e		٠.	•	
. 2	3		ice at 0°C melts when (Assume temperature of				
(a) $\frac{}{5}$	(b) $\frac{3}{5}$		(a) - 4.5 <i>cal/K</i>		+ 4.5 <i>cal/K</i>	, -,	
(c) $\frac{3}{7}$	(d) $\frac{3}{4}$		(c) +5.4 <i>cal/K</i>	(d)	– 5.4 <i>cal/K</i>		
(c) $\frac{3}{7}$	$\frac{(u)}{4}$	26.	An ideal gas expands in	such a manno	er that its pressure and	volume	
When an ideal gas ($\gamma = 5$	/3) is heated under constant pressure,		can be related by equati				

(c) 60 %

19.

20.

21.

the gas is

(a) Heated

(b) Cooled

(d) 20 %

[SCRA 1998]

[UPSEAT 2002]

[MP PET 1996, 99; MP PMT 1997, 99]

Which one of the following gases possesses the largest internal

(d) 6×10^{26} molecules of argon occupying $40m^3$ at 900 K

Two samples A and B of a gas initially at the same pressure and

temperature are compressed from volume V to V/2 (A isothermally

Initial pressure and volume of a gas are P and V respectively. First it

is expanded isothermally to volume 4V and then compressed

adiabatically to volume V. The final pressure of gas will be [CBSE PMT 1999]

(a) 2 moles of helium occupying $1m^3$ at 300 K

(b) 56 kg of nitrogen at $107Nm^{-2}$ and 300 K

(c) 8 grams of oxygen at 8 atm and 300 K

and adiabatically). The final pressure of A is

(a) Greater than the final pressure of B

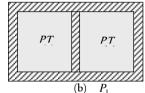
(b) Equal to the final pressure of B (c) Less than the final pressure of B (d) Twice the final pressure of B

- (c) Neither heated nor cooled
- (d) First heated and then cooled
- A Carnot engine whose low temperature reservoir is at $7^{\circ}C$ has an 27. efficiency of 50%. It is desired to increase the efficiency to 70%. By how many degrees should the temperature of the high temperature reservoir be increased

[UPSEAT 2005]

- (a) 840 K
- (b) 280 K
- (c) 560 K
- (d) 380 K
- 28. P-V diagram of a diatomic gas is a straight line passing through origin. The molar heat capacity of the gas in the process will be
 - (a) 4 R
- (b) 2.5 R

- (c) 3 R
- Following figure shows on adiabatic cylindrical container of volume 29. V_0 divided by an adiabatic smooth piston (area of cross-section =
 - A) in two equal parts. An ideal gas $(C_P / C_V = \gamma)$ is at pressure Pand temperature T in left part and gas at pressure P and temperature T in right part. The piston is slowly displaced and released at a position where it can stay in equilibrium. The final pressure of the two parts will be (Suppose x = displacement of the piston)



- (a) P_2

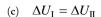
- 30. Two cylinders A and B fitted with pistons contain equal amounts of an ideal diatomic gas at 300 K. The piston of A is free to move while that of B is held fixed. The same amount of heat is given to the gas in each cylinder. If the rise in temperature of the gas in A is 30 K, then the rise in temperature of the gas in B is
 - (a) 30 K
- (b) 18 K
- 50 K (c)
- (d) 42 K

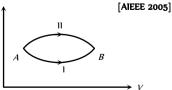
Graphical Questions

A system goes from A to B via two processes I and II as shown in figure. If ΔU_1 and ΔU_2 are the changes in internal energies in the processes I and II respectively, then

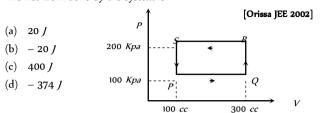


 $\begin{array}{ll} \mbox{(a)} & \Delta U_{\rm II} > \Delta U_{\rm I} \\ \mbox{(b)} & \Delta U_{\rm II} < \Delta U_{\rm I} \end{array}$





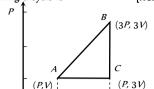
- (d) Relation between $\Delta U_{
 m I}$ and $\Delta U_{
 m II}$ can not be determined
- A thermodynamic system is taken through the cycle PQRSP process. 2. The net work done by the system is



[IIT 1998]

An ideal gas is taken around ABCA as shown in the above P-V 3. [KCET 2001] diagram. The work done during a cycle is

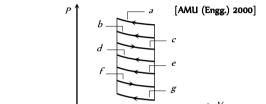
2PV



1/2PV

- (d) Zero
- The P-V diagram shows seven curved paths (connected by vertical paths) that can be followed by a gas. Which two of them should be parts of a closed cycle if the net work done by the gas is to be at its maximum value





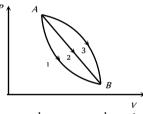
- An ideal gas of mass m in a state A goes to another state B via three 5. different processes as shown in figure. If $\,Q_1,Q_2\,$ and $\,Q_3\,$ denote the heat absorbed by the gas along the three paths, then

(a)
$$Q_1 < Q_2 < Q_3$$

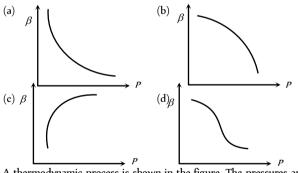
(b)
$$Q_1 < Q_2 = Q_3$$

(c)
$$Q_1 = Q_2 > Q_3$$

(d)
$$Q_1 > Q_2 > Q_3$$



6. Which of the following graphs correctly represents the variation of $\beta = -(dV/dP)/V$ with P for an ideal gas at constant temperature [IIT-JEE (Screening) 2002]

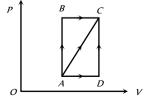


A thermodynamic process is shown in the figure. The pressures and 7. volumes corresponding to some points in the figure are :

$$P_A = 3 \times 10^4 Pa$$
, $P_B = 8 \times 10^4 Pa$ and $V_A = 2 \times 10^{-3} m^3$, $V_D = 5 \times 10^{-3} m^3$

In process AB, 600 / 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





P-V plots for two gases during adiabatic process are shown in the figure. Plots 1 and 2 should correspond respectively to

[IIT-JEE (Screening) 2001]

(a)
$$He$$
 and O_2

(b)
$$O_2$$
 and He

c)
$$He$$
 and Ar

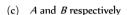
(d)
$$O_2$$
 and N_2



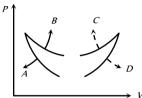
given amount of gas. The curves which represent adiabatic and isothermal changes are

[CPMT 1986; UPSEAT 1999]



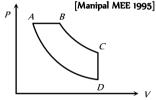




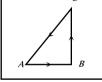


- In pressure-volume diagram given below, the isochoric, isothermal, 10 and isobaric parts respectively, are
 - BA, A [MP PET 1992]





The P-V diagram of a system undergoing thermodynamic 11. transformation is shown in figure. The work done on the system in going from $A \rightarrow B \rightarrow C$ is 50 J and 20 cal heat is given to the system. The change in internal energy between A and C is

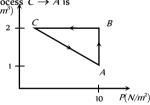


An ideal gas is taken through the cycle $A \rightarrow B \rightarrow C \rightarrow A$, as shown in the figure. If the net heat supplied to the gas in the cycle is 5 J, the work done by the gas in the process



(c)
$$-15 J$$



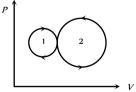


In the following indicator diagram, the net amount of work done 13. will be

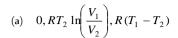




[CBSE PMT 1992]



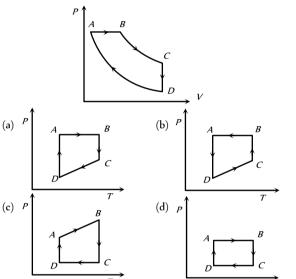
- (d) Infinity
- A cyclic process for 1 mole of an ideal gas is shown in figure in the V-T, diagram. The work done in AB, BC and CA respectively



- (b) $R(T_1 T_2), 0, RT_1 \ln \frac{V_1}{V_2}$
- V_2 V_2 V_3 V_4 V_5 V_6 V_7 V_8 V_8 V_9 V_9
- (d) $0, RT_2 \ln \left(\frac{V_2}{V_1}\right), R(T_2 T_1)$

(c) $0, RT_2 \ln \left(\frac{V_2}{V_1}\right), R(T_1 - T_2)$

15. A cyclic process *ABCD* is shown in the figure *P-V* diagram. Which of the following curves represent the same process



16. Carnot cycle (reversible) of a gas represented by a Pressufre-Volume curve is shown in the diagram

Consider the following statements

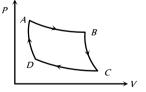
- 1. Area *ABCD* = Work done on the gas
- II. Area *ABCD* = Net heat absorbed
- III. Change in the internal energy in cycle = 0

Which of these are correct

[AMU (Med.) 2001]

[AIEEE 2005]

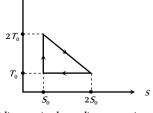
- (a) 1 only
- (b) Il only
- (c) II and III
- (d) 1, 11 and 111



17. The temperature-entropy diagram of a reversible engine cycle is given in the figure. Its efficiency is



- (b) 2/3
- (c) 1/2
- (d) 1/4

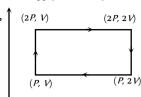


18. Work done in the given *P-V* diagram in the cyclic process is

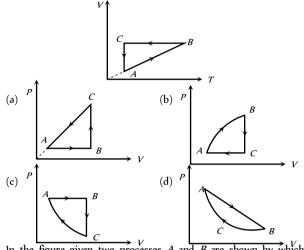
[UPSEAT 1998; RPET 2000; Kerala PMT 2002]

- (a) *PV*
- (b) 2*PV*
- (c) PV/2
- (d) 3*PV*

19.



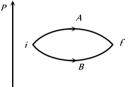
A cyclic process *ABCA* is shown in the *V-T* diagram. Process on the P-V diagram is



20. In the figure given two processes A and B are shown by which a thermo-dynamical system goes from initial to final state F. If ΔQ_A and ΔQ_B are respectively the heats supplied to the systems then

(a)
$$\Delta Q_A = \Delta Q_B$$

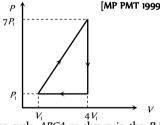
- (b) $\Delta Q_A \ge \Delta Q_B$
- (c) $\Delta Q_A < \Delta Q_B$
- (d) $\Delta Q_A > \Delta Q_B$



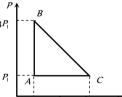
21. In the cyclic process shown in the figure, the work done by the gas in one cycle is $P \uparrow$ [MP PMT 1999]



- (b) $14 P_1 V_1$
- (c) $18 P_1 V_1$
- (d) $9 P_1 V_1$

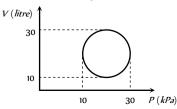


- 22. An ideal gas is taken around the cycle $\stackrel{V_1}{ABCA}$ as shown in the P-V diagram. The net work done by the gas during the cycle is equal to
 - (a) $12 P_1 V_1$
 - (b) $6 P_1 V_1$
 - (c) $3 P_1 V_1$
 - (1) 2017
 - (d) $2P_1V_1$

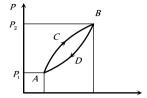


23. Heat energy absorbed by a system in going through, a cyclic process shown in figure is [AIIMS 1995; BHU 2002]

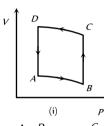


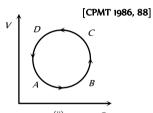


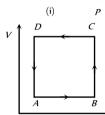
- (c) 10·π J
- (d) $10^{-3} \pi J$
- 24. A thermodynamic system is taken from state A to B along ACB and is brought back to A along BDA as shown in the PV diagram. The net work done during the complete cycle is given by the area
 - PACBPP
 - ACBB'A'A
 - **ACBDA**
 - ADBB'A'A

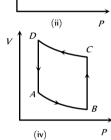


In the diagrams (i) to (iv) of variation of volume with changing 25. pressure is shown. A gas is taken along the path ABCD. The change in internal energy of the gas will be



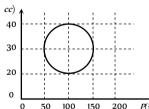






- (a) Positive in all cases (i) to (iv)
- Positive in cases (i), (ii) and (iii) but zero in (iv) case
- Negative in cases (i), (ii) and (iii) but zero in (iv) case
- Zero in all four cases
- 26. A system is taken through a cyclic process represented by a circle as shown. The heat absorbed by the system is



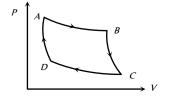


- $4\pi \times 10^2 J$
- 100
- 27. A thermodynamic system undergoes cyclic process ABCDA as shown in figure. The work done by the system is
 - (a) P_0V_0

 $2P_{0}V_{0}$

- $3P_0$

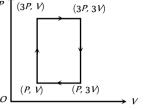
- 28. The P-V graph of an ideal gas cycle is shown here as below. The adiabatic process is described by
 - [CPMT 1985; UPSEAT 2003]



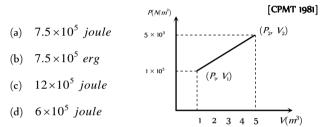
- AB and BC
- (b) AB and CD
- (c) BC and DA
- (d) BC and CD
- An ideal mensus town ideal mensus taken round the cycle ABCDA as shown 29. in following P-V diagram. The work done during the cycle is [IIT 1983; CPMT 19



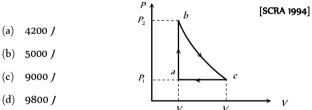
- (b) 2 PV
- (c) 4 PV
- (d) Zero



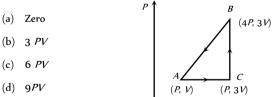
A system changes from the state (P_1, V_1) to (P_2V_2) as shown in 30. the figure. What is the work done by the system



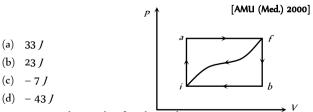
Carbon monoxide is carried around a closed cycle abc in which bc is 31. an isothermal process as shown in the figure. The gas absorbs 7000 J of heat as its temperature increases from 300 K to 1000 K in going from a to b. The quantity of heat rejected by the gas during the process ca is



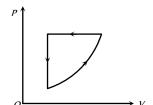
A sample of ideal monoatomic gas is taken round the cycle ABCA as 32. shown in the figure. The work done during the cycle is



When a system is taken from state i to a state f along path iaf, 33. Q = 50 J and W = 20 J. Along path ibf, Q = 35 J. If W = -13J for the curved return path fi, Q for this path is



For one complete cycle of a thermodynamic process on a gas as 34. shown in the P-V diagram, Which of following is correct



(a)
$$\Delta E_{\rm int} = 0, Q < O$$

(b)
$$\Delta E_{\text{int}} = 0, Q > 0$$

(c)
$$\Delta E_{\text{int}} > 0, Q < 0$$

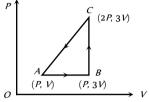
(d)
$$\Delta E_{\rm int} < 0, Q > 0$$

An ideal gas is taken around ABCA as shown in the above P-V 35. diagram. The work done during a cycle is

[UPSEAT 2001]

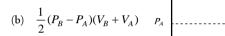
(a) Zero





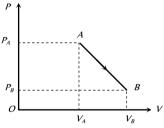
An ideal gas is taken from point A to the point B, as shown in the 36. P-V diagram, keeping the temperature constant. The work done in the process is [UPSEAT 2005]

(a)
$$(P_A - P_B)(V_B - V_A)$$



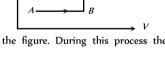
(c)
$$\frac{1}{2}(P_B - P_A)(V_B - V_A)$$
 P_B

(d)
$$\frac{1}{2}(P_B + P_A)(V_B - V_A)$$



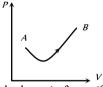
The P-V diagram of a system undergoing thermodynamic 37. transformation is shown in figure. The work done by the system in going from $A \rightarrow B \rightarrow C$ is 30/ and 40/ heat is given to the system. The change in internal energy between A and C is

- (a) 10 /
- 70 *l*
- 84 /
- (d) 134 J



Consider a process shown in the figure. During this process the 38. work done by the system

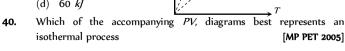
- (a) Continuously increases
 - Continuously decreases
 - (c) First increases, then decreases
 - (d) First decreases, then increases

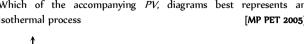


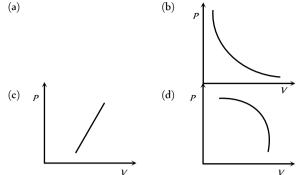
Six moles of an ideal gas performs a cycle shown in figure. If the 39. temperature are $T = 600 \, \text{K}$, $T = 800 \, \text{K}$, $T = 2200 \, \text{K}$ and $T = 1200 \, \text{K}$, the work done per cycle is



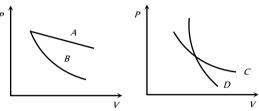
- 20 kl
- (b) 30 kJ
- 40 *kJ*
- (d) 60 kl







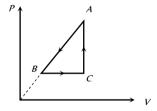
In the following figure, four curves A, B, C and D are shown. The [DCE 2003]



- Isothermal for A and D while adiabatic for B and C
- Adiabatic for A and C while isothermal for B and D
- Isothermal for A and B while adiabatic for C and D
- (d) Isothermal for A and C while adiabatic for B and D

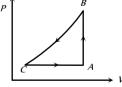
P-V diagram of a cyclic process ABCA is as shown in figure. Choose the correct statement

- (a) $\Delta Q_{A \to B} = \text{negative}$
- $\Delta U_{B \to C}$ = positive
- [BCECE 2005] ΔW_{CAB} = negative
- (d) All of these

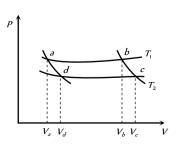


43. A sample of an ideal gas is taken through a cycle a shown in figure. It absorbs 50/ of energy during the process AB, no heat during BC, rejects 70/ during CA. 40/ of work is done on the gas during BC. Internal energy of gas at A is 1500), the internal energy at C would

- 1590 / (a)
- 1620 /
- 1540 *J*
- (d) 1570 J



In the following P-V diagram two adiabatics cut two isothermals at 44. temperatures T and T (fig.). The value of $\frac{V_a}{V_c}$ will be



				Thermodynemics 677
	(a) $\frac{V_b}{V_a}$ (b) $\frac{1}{V_b}$	V _c	Reason	: When a system changes from one thermal equilibrium to another, some heat is absorbed by it.
	V_d	12.	Assertion	: A room can be cooled by opening the door of a refrigerator in a closed room.
	(c) $\frac{\sqrt[4]{d}}{V_a}$ (d) V	$V_b V_c$	Reason	: Heat flows from lower temperature (refrigerator) to higher temperature (room).
	Assertion & F		. Assertion	: It is not possible for a system, unaided by an external agency to transfer heat from a body at lower temperature to another body at higher temperature.
	For the assertion and reason carefully to mark obtions given below: If both assertion and reason are true are		Reason	: According to Clausius statement, "No process is possible whose sole result is the transfer of heat from a cooled object to a hotter object.
(b)	explanation of the assertion. If both assertion and reason are true by	14.	. Assertion	: If an electric fan be switched on in a closed room, the air of the room will be cooled.
(c)	explanation of the assertion. If assertion is true but reason is false.		Reason	: Fan air decrease the temperature of the room.
(d) (e)	If the assertion and reason both are false If assertion is false but reason is true.	2. 15.	. Assertion	: The internal energy of an isothermal process does not change.
1.	Assertion : Reversible systems are d Reason : Most processes are dissip	ifficult to find in real world. native in nature	Reason	: The internal energy of a system depends only on pressure of the system.
2.		s adiabatic expansion.[AIIMS 2005]	. Assertion	: In an adiabatic process, change in internal energy of a gas is equal to work done on or by the gas in the process.
3.	Assertion : Thermodynamic process Reason : Dissipative effects can no		Reason	: Temperature of gas remains constant in a adiabatic process.
4.		arbonated drink is opened, a 17.	Assertion	: An adiabatic process is an isoentropic process.
		ne opening. the gas causes lowering of sation of water vapours.[A11MS 2003]	Reason 1	: Change in entropy is zero in case of adiabatic process.
5.	Assertion : The isothermal curves certain point.	intersect each other at a 18. takes place slowly, so the	•	: Work done by a gas in isothermal expansion is more than the work done by the gas in the same expansion, adiabatically.
	isothermal curves have v		Reason	: Temperature remains constant in isothermal expansion and not in adiabatic expansion.
6.	temperature of the syste		. Assertion	: First law of thermodynamics is a restatement of the principle of conservation
	Reason : The adiabatic compression	[AllMS 2001]	Reason	: Energy is fundamental quantity.
7.	Assertion : In isothermal process w supplied to the body is c energy.	hole of the heat energy 20	. Assertion	: Zeroth law of thermodynamic explain the concept of energy.
	Reason : According to the first lav	w of thermodynamics	Reason	: Energy is dependent on temperature.
	$\Delta Q = \Delta U + p \Delta V .$	[AllMS 1997] 21.	Assertion	: Efficiency of a Carnot engine increase on reducing
8.	without giving (or taking	ne temperature of a body g) heat to (or from) it. of conservation of energy,	Reason	the temperature of sink.The efficiency of a Carnot engine is defined as ratio of net mechanical work done per cycle by the gas to
		should remains conserved.		the amount of heat energy absorbed per cycle from

22.

Assertion

Reason

: The specific heat of a gas is an adiabatic process is

: Specific heat of a gas in directly proportional to

: Work and heat are two equivalent form of energy.

: Work is the transfer of mechanical energy irrespective of temperature difference, whereas heat is the transfer of thermal energy because of

: The heat supplied to a system is always equal to the

change of heat in system and inversely proportional

zero and in an isothermal process is infinite.

to change in temperature.

temperature difference only.

increase in its internal energy.

9.

10.

11.

Assertion

Reason

Assertion

Assertion

Reason



3

the source.

First Law of Thermodynamics ($\triangle Q = \triangle U + \triangle W$)

: The entropy of the solids is the highest

: Atoms of the solids are arranged in orderly manner.

5

6	b	7	b	8	d	9	а	10	d
11	С	12	а	13	d	14	а	15	b
16	b	17	С	18	d	19	d	20	b
21	а	22	d	23	b	24	а	25	d
26	d	27	а	28	b	29	d	30	а
31	b	32	С	33	С	34	а	35	а
36	b	37	С	38	С	39	b	40	а
41	С	42	а	43	а	44	С	45	С
46	а	47	С	48	b				

Isothermal Process

1	С	2	а	3	С	4	d	5	b
6	b	7	С	8	d	9	а	10	С
11	а	12	b	13	а	14	а	15	С
16	С	17	а	18	С	19	а	20	С
21	b	22	b	23	а	24	а	25	а
26	С	27	b	28	b	29	b	30	а
31	d								

Adiabatic Process

1	С	2	С	3	b	4	d	5	С
6	d	7	С	8	b	9	а	10	а
11	С	12	d	13	b	14	d	15	d
16	b	17	а	18	С	19	а	20	С
21	d	22	С	23	b	24	С	25	а
26	b	27	d	28	а	29	d	30	d
31	а	32	С	33	d	34	С	35	а
36	d	37	b	38	d	39	а	40	d
41	С	42	С	43	а	44	а	45	b
46	d	47	а	48	b	49	b	50	а
51	b	52	d	53	b	54	b	55	d
56	b	57	С						

Isobaric and Isochoric Processes

1	а	2	С	3	С	4	а	5	а
6	С	7	С	8	b	9	d	10	С
11	С	12	а	13	b	14	a	15	d
16	b	17	а	18	d	19	С	20	d
21	d	22	d	23	а				

Heat Engine, Refrigerator and Second Law of Thermodynamics

1	I	d	2	С	3	b	4	С	5	b
6	3	d	7	b	8	d	9	b	10	b
1	11	С	12	b	13	С	14	а	15	а
1	16	а	17	b	18	d	19	b	20	а

21	b	22	b	23	С	24	а	25	b
26	а	27	d	28	а	29	b	30	С
31	а	32	а	33	С	34	d		

Critical Thinking Questions

1	d	2	С	3	bc	4	С	5	d
6	d	7	С	8	d	9	С	10	a
11	b	12	С	13	b	14	С	15	a
16	С	17	b	18	а	19	b	20	С
21	b	22	d	23	d	24	b	25	b
26	b	27	d	28	С	29	С	30	d

Graphical Questions

1	С	2	b	3	а	4	С	5	а
6	а	7	а	8	b	9	С	10	d
11	d	12	а	13	b	14	С	15	а
16	С	17	а	18	a	19	С	20	d
21	d	22	d	23	С	24	С	25	d
26	b	27	d	28	С	29	С	30	С
31	d	32	b	33	d	34	а	35	d
36	d	37	а	38	a	39	С	40	b
41	d	42	d	43	a	44	а		

Assertion & Reason

1	а	2	а	3	а	4	а	5	е
6	d	7	е	8	d	9	а	10	а
11	d	12	d	13	а	14	d	15	С
16	С	17	а	18	b	19	С	20	е
21	b	22	а						

Answers and Solutions

First Law of Thermodynamics ($\Delta Q = \Delta U + \Delta W$)

- (a) $\Delta Q = \Delta U + \Delta W$ and $\Delta W = P\Delta V$
- **2.** (c)
- 3. (b) $\Delta Q = \Delta U + \Delta W$ $\Rightarrow \Delta U = \Delta Q - \Delta W = Q - W$ (using proper sign)
- **4.** (b) $\Delta U = \Delta Q W = 35 15 = 20 J$
- $\textbf{5.} \hspace{0.5in} \textbf{(c)} \hspace{0.2in} \textbf{Internal energy depends only on the temperature of the gas.} \\$
- **6.** (b)

7. (b) (i) Case
$$\rightarrow$$
 Volume = constant $\Rightarrow \int PdV = 0$

(ii) Case
$$\rightarrow P = \text{constant} \implies \int_{V_1}^{2V_1} P dV = P \int_{V_1}^{2V_1} dV = P V_1$$

8. (d)
$$\Delta Q = \Delta W + \Delta U \Rightarrow 35 = -15 + \Delta U \Rightarrow \Delta U = 50J$$

9. (a)
$$J\Delta Q = \Delta U + \Delta W$$
, $\Delta U = J\Delta Q - \Delta W$
$$\Delta U = 4.18 \times 300 - 600 = 654 \ Joule$$

10. (d) Work done
$$=\int_1^2 P dV$$
, which is state dependent as well as path dependent.

11. (c)
$$\Delta Q = \Delta U + \Delta W :: \Delta W = 0 \Rightarrow \Delta Q = \Delta U = \frac{f}{2} \mu R \Delta T$$

$$= \frac{3}{2} \times 2R(373 - 273) = 300R.$$

12. (a)
$$\Delta Q = 2k \ cal = 2 \times 10^3 \times 4.2 J = 8400 J \ and \ \Delta W = 500 J.$$
Hence from $\Delta Q = \Delta U + \Delta W$, $\Delta W = \Delta Q - \Delta U = 8400 - 500 = 7900 J$

13. (d) Change in internal energy
$$(\Delta U)$$
 depends upon initial an find state of the function while ΔQ and ΔW are path dependent

14. (a) This is the case of free expansion and in this case
$$\Delta W=0$$
 ,
$$\Delta U=0 \ \ {\rm so\ temperature\ remains\ same}\ \ \emph{i.e.}\ \ 300\ \ \emph{K}.$$

15. (b)
$$\Delta Q = \Delta U + \Delta W \Rightarrow \Delta W = \Delta Q - \Delta U = 100 - 40 = 70J$$

17. (c)
$$\Delta Q = \Delta U + \Delta W = 167 + 333 = 500 \ cal$$

19. (d) Change in internal energy does not depend upon path so
$$\Delta U = \Delta Q - \Delta W \ \ {\rm remain\ constant}.$$

20. (b)
$$\Delta Q = \Delta U + \Delta W$$
; $\Delta Q = 200J$ and $\Delta W = -100J$
$$\Rightarrow \Delta U = \Delta Q - \Delta W = 200 - (-100) = 300J$$

21. (a) During free expansion of a perfect gas no, work is done and also no heat is supplied from outside. Therefore, no change in internal energy. Hence, temperature remain constant.

22. (d)
$$\Delta Q = \Delta U + \Delta W \Rightarrow \Delta U = \Delta Q - \Delta W = 150 - 110 = 40 J$$

23. (b) From FLOT $\Delta O = \Delta U + \Delta W$

: Heat supplied to the system so $\Delta Q \to {\sf Positive}$ and work is done on the system so $\Delta W \to {\sf Negative}$ Hence + $\Delta Q = \Delta U - \Delta W$

24. (a)

25. (d) State of a thermodynamic state cannot determine by a single variable (*P* or *V* or *T*)

26. (d) *R* is the universal gas constant.

27. (a) From FLOT $\Rightarrow dU = dQ - dW \Rightarrow dU = dQ(<0) \quad (\because dW = 0)$ $\Rightarrow dU < 0 \text{ So temperature will decrease.}$

28. (b) From FLOT $\Delta Q = \Delta U + \Delta W$

Work done at constant pressure $(\Delta W)_P = (\Delta Q)_P - \Delta U$

$$(\Delta Q)_P - (\Delta Q)_V$$
 (As we know $(\Delta Q)_V = \Delta U$)

Also
$$(\Delta Q)_P = mc_P \Delta T$$
 and $(\Delta Q)_V = mc_V \Delta T$

$$\Rightarrow (\Delta W)_P = m(c_P - c_V) \Delta T$$

$$\Rightarrow (\Delta W)_P = 1 \times (3.4 \times 10^3 - 2.4 \times 10^3) \times 10 = 10^4 cal$$

29. (d)

30. (a) Ideal gas possess only kinetic energy.

31. (b) The internal energy and entropy depend only on the initial and final states of the system and not on the path followed to attain that state.

32. (c) $\Delta Q = \Delta U + \Delta W$ $\therefore \Delta Q = 200 cal = 200 \times 4.2 = 840 J \text{ and } \Delta W = 40 J$ $\Rightarrow \Delta U = \Delta Q - \Delta W = 840 - 40 = 800 J$

33. (c)
$$\Delta Q = \Delta U + \Delta W = (U_f - U_i) + \Delta W$$

 $\Rightarrow 30 = (U_f - 40) + 10 \Rightarrow U_f = 60J$

34. (a) With rise in temperature, internal energy also increases.

35. (a)

36. (b) Heat supplied to a gas raise its internal energy and does some work against expansion, so it is a special case of law of conservation of energy.

37. (c) Change in internal energy is always equal to the heat supplied

i.e.
$$\Delta U = (\Delta Q)_V = \mu C_V \Delta T$$
.

For monoatomic gas $C_V = \frac{3}{2} R$

$$\Rightarrow \Delta U = \mu \left(\frac{3}{2}R\right) \Delta T = 1 \times \frac{3}{2} \times 8.31 \times (100 - 0)$$

$$=12.48 \times 10^2 J$$

38. (c) $\Delta U = \mu C_V \Delta T = n \left(\frac{R}{\gamma - 1} \right) \Delta T$ $\Rightarrow \Delta U = \frac{P \Delta V}{(\gamma - 1)} = \frac{P(2V - V)}{\gamma - 1} = \frac{PV}{(\gamma - 1)}$

39. (b) $\Delta U = \mu C_V \Delta T = 2 \times 4.96 \times (342 - 340) = 19.84 \ cal$

40. (a)

41. (c) According to FLOT $\Delta Q = \Delta U + P(\Delta V) \Rightarrow \Delta U = \Delta Q - P(\Delta V)$ $= 1500 - (2.1 \times 10^5)(2.5 \times 10^{-3}) = 975 \text{ loule}$

42. (a) $\Delta Q = \Delta U + \Delta W \Rightarrow \Delta U = \Delta Q - \Delta W$ = $6 \times 4.18 - 6 = 19.08kJ \approx 19.1kJ$

43. (a) Given $\Delta Q=-20J$, $\Delta W=-8J$ and $U_i=30J$ $\Delta Q=\Delta U+\Delta W\Rightarrow \Delta U=(\Delta Q-\Delta W)$ $\Rightarrow (U_f-U_i)=(U_f-30)=-20-(-8)\Rightarrow U_f=18J$

44. (c) Change in internal energy $\Delta U = \mu \, C_V \Delta T$ it doesn't depend upon type of process. Actually it is a state function

45. (c)

46. (a) In first process using $\Delta Q = \Delta U + \Delta W$ $\Rightarrow 8 \times 10^5 = \Delta U + 6.5 \times 10^5 \Rightarrow \Delta U = 1.5 \times 10J$ Since final and initial states are same in both process
So ΔU will be same in both process
For second process using $\Delta Q = \Delta U + \Delta W$ $\Rightarrow 10^5 = 1.5 \times 10^5 + \Delta W \Rightarrow \Delta W = -0.5 \times 10^5 J$

47. (c) $\Delta W = P\Delta V$; here ΔV is negative so ΔW will be negative

48. (b) Entropy is related to second law of thermodynamics.

Isothermal Process

1. (c) In isothermal process temperature remains constant.

2. (a) If isothermal curves cut each other then at equilibrium two temperature will be there which is impossible.

3. (c) In isothermal expansion temperature remains constant, hence no change in internal energy.

4. (d) $W = \mu RT \log_e \frac{V_2}{V_1}$ $= \left(\frac{m}{M}\right) RT \log_e \frac{V_2}{V_1} = 2.3 \times \frac{m}{M} RT \log_{10} \frac{V_2}{V_1}$ $= 2.3 \times \frac{96}{32} R (273 + 27) \log_{10} \frac{140}{70} = 2.3 \times 900 R \log_{10} 2$

5. (b) $0.8 \times 5 = P \times (3+5) \Rightarrow P = 0.5 m$

6. (b) Differentiate $PV = \text{constant } w.r.t \ V$ $\Rightarrow P\Delta V + V\Delta P = 0 \Rightarrow \frac{\Delta P}{R} = -\frac{\Delta V}{V}$

7. (c)

8. (d)
$$W = -\mu RT \log_e \frac{V_2}{V_1} = -1 \times 8.31 \times (273 + 0) \log_e \left(\frac{22.4}{11.2}\right)$$

= $-8.31 \times 273 \times \log_e 2 = -1572.5J$ [: $\log_e 2 = 0.693$]

9. (a) $E_{\theta}=P$, if $P={\rm constant},\ E_{\theta}={\rm constant}$

10. (c) For isothermal process
$$PV = RT \Rightarrow P = \frac{RT}{V}$$

$$\therefore W = PdV = \int_{V_1}^{V_2} \frac{RT}{V} dV = RT \log_e \frac{V_2}{V_1}$$

II. (a) $E_{\theta} = P$

12. (b) For such a case, pressure
$$=\frac{1}{\text{Compressibility}}$$

13. (a) $E_{\theta} = P = 1.013 \times 10^5 \, N/m^2$

14. (a) In isothermal process, compressibility $E_{\theta} = \rho$.

15. (c) In isothermal process, exchange of energy takes place between system and surrounding to maintain the system temperature constant.

16. (c) No change in the internal energy of ideal gas but for real gas internal energy increases because work is done against intermolecular forces.

17. (a) In isothermal process temperature remains constant. i.e., $\Delta T=0 \ . \ \ \text{Hence according to} \ \ C=\frac{Q}{m\Delta T} \Rightarrow C_{iso}=\infty$

18. (c) This is the case of free expansion of gas. In free expansion $\Delta U=0 \Longrightarrow$ Temp. remains same.

19. (a) An isothermal process takes place at constant temperature, must be carried out in a vessel with conducting wall so that heat generated should go out at once.

20. (c) For isothermal process

$$dU = 0$$
 and work done $= dW = P(V_2 - V_1)$
 $V = V$

$$\because V_2 = \frac{V_1}{2} = \frac{V}{2} \therefore dW = -\frac{PV}{2}$$

21. (b) In isothermal process, temperature remains constant.

22. (b) In isothermal process, heat is released by the gas to maintain the constant temperature.

23. (a) In isothermal compression, there is always an increase of heat. which must flow out the gas.

$$\Delta Q = \Delta U + \Delta W \Rightarrow \Delta Q = \Delta W \ (\because \Delta U = 0)$$

$$\Rightarrow \Delta Q = -1.5 \times 10^4 J = \frac{1.5 \times 10^4}{4.18} cal = -3.6 \times 10^3 cal$$

24. (a) In isothermal change, temperature remains constant, Hence ΔU = 0.

Also from $\Delta Q = \Delta U + \Delta W \implies \Delta Q = \Delta W$

25. (a) It is an isothermal process. Hence work done = $P(V_2-V_1)$ = $1\times10^5\times(1.091-1)\times10^{-6}=0.0091~J$

26. (c) $\Delta Q = \Delta U + \Delta W \implies \Delta U = \Delta Q - \Delta W = 2240 - 168 = 2072 J$.

27. (b) Amount of heat given = $540 \ calories$ Change in volume $\Delta V = 1670 \ c.c$

Atmospheric pressure $P = 1.01 \times 10^6 \ dyne \ / \ cm^2$

Work done against atmospheric pressure

$$W = P\Delta V = \frac{1.01 \times 10^6 \times 1670}{4.2 \times 10^7} \approx 40 \ cal$$

28. (b) $W_{iso} = \mu RT \log_e \frac{V_2}{V_1} = 1 \times 8.31 \times 300 \log_e \frac{20}{10} = 1728 J$

29. (b)
$$W = \mu RT \log_e \left(\frac{V_2}{V_1}\right) = 0.2 \times 8.3 \times \log_e 2 \times (27 + 273)$$

= $0.2 \times 8.3 \times 300 \times 0.693 = 345 J$

30. (a) For isothermal process $P_1V_1 = P_2V_2$

$$\Rightarrow P_2 = \frac{P_1 V_1}{V_2} = \frac{72 \times 1000}{900} = 80 \text{ cm}$$

Stress $\Delta P = P_2 - P_1 = 80 - 72 = 8cm$

31. (d) During isothermal change $T = \text{constant} \Rightarrow \Delta U = 0$ also from FLOT, $\Delta Q = \Delta W$.

Adiabatic Process

 (c) Gas cylinder suddenly explodes is an irreversible adiabatic change and work done against expansion reduces the temperature.

2. (c) Work done in adiabatic change $=\frac{\mu R(T_1-T_2)}{\gamma-1}$

3. (b) In case of adiabatic expansion ΔW = positive and $\Delta Q=0$ from FLOT $\Delta Q=\Delta U+\Delta W$ \Rightarrow $\Delta U=-\Delta W$ i.e., ΔU will be negative.

4. (d) For adiabatic process $\frac{T^{\gamma}}{P^{\gamma-1}} = \text{constant}$

$$\Rightarrow \frac{T_2}{T_1} = \left(\frac{P_1}{P_2}\right)^{\frac{1-\gamma}{\gamma}} \Rightarrow \frac{T_2}{300} = \left(\frac{4}{1}\right)^{\frac{(1-1.4)}{1.4}} \Rightarrow T_2 = 300(4)^{-\frac{0.4}{1.4}}$$

5. (c)
$$PV^{\gamma} = \text{constant} \Rightarrow \frac{P_2}{P_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} \Rightarrow \frac{P_2}{1} = \left(\frac{V_1}{V_1/4}\right)^{3/2} = 8$$

 $\Rightarrow P_2 = 8 \text{ atm.}$

6. (d)
$$PV^{\gamma} = \text{constant} \Rightarrow \frac{P_2}{P_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} \Rightarrow P_2 = (8)^{5/3} P_1 = 32 P_1$$

7. (c) Volume of the gas $V = \frac{m}{d}$ and using $PV^{\gamma} = \text{constant}$

We get
$$\frac{P'}{P} = \left(\frac{V}{V'}\right)^{\gamma} = \left(\frac{d'}{d}\right)^{\gamma} = (32)^{7/5} = 128$$

8. (b)
$$\frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} \Rightarrow T_2 = 300 \left(\frac{27}{8}\right)^{\frac{3}{3} - 1} = 300 \left(\frac{27}{8}\right)^{\frac{2}{3}}$$

$$= 300 \left\{ \left(\frac{27}{8}\right)^{1/3} \right\}^2 = 800 \left(\frac{3}{2}\right)^2 = 675 K$$

$$\Rightarrow \Delta T = 675 - 300 = 375 K$$

(a) In thermodynamic processes.Work done = Area covered by PV diagram with V-axis

From graph it is clear that (Area)_{iso} > (Area)_{adi}

 $\Rightarrow W_{iso} > W_{adi} \qquad P$ | Isothermal | Adiabatic |

10. (a) Since PV = RT and T = constant; $\therefore PV = \text{constant}$.

11. (c) For Isothermal process PV= constant $\Rightarrow \left(\frac{dP}{dV}\right) = \frac{-P}{V} = \text{Slope of Isothermal curve}$

For adiabatic $PV^{\gamma} = \text{constant}$

$$\Rightarrow \frac{dP}{dV} = \frac{-\gamma P}{V} = \text{Slop of adiabatic curve slope}$$

Clearly, $\left(\frac{dP}{dV}\right)_{\text{adiabatic}} = \gamma \left(\frac{dP}{dV}\right)_{\text{Isothermal}}$

12. (d) $PV^{\gamma} = \text{constant} \Rightarrow P\left(\frac{RT}{P}\right)^{\gamma} = \text{constant}$ $\Rightarrow P^{1-\gamma}T^{\gamma} = \text{constant}$

13. (b) $W_{adi} = \frac{R}{\gamma - 1} (T_i - T_f) = \frac{R}{\gamma - 1} (T - T_1)$

14. (d) $dQ = 0 = -2 + dW \Rightarrow dW = 2 J$ \Rightarrow Work done by the gas = 2 J \Rightarrow Work done on the gas = -2 J

15. (d) $E_{\phi} = \gamma P = 1.4 \times (1 \times 10^5) = 1.4 \times 10^5 N / m^2$

16. (b) Slope of adiabatic curve = $\gamma \times$ (Slope of isothermal curve)

17. (a) Due to compression the temperature of the system increases to a very high value. This causes the flow of heat from system to the surroundings, thus decreasing the temperature. This decrease in temperature results in decrease in pressure.

18. (c) $\Delta Q = \Delta U + \Delta W = 0 \Rightarrow \Delta W = -\Delta U$ if ΔW is positive *i.e.*, gas does work then ΔU should be negative meaning internal energy is used in doing work.

19. (a) $W = \frac{R}{\gamma - 1} (T_1 - T_2)$ $= \frac{8.31 \times \{(273 + 27) - (273 + 127)\}}{1.4 - 1} = -2077.5 \text{ joules}$

20. (c) Pressure is reduced, so the temperature falls.

21. (d) Adiabatic Bulk modulus $E_{\phi} = \gamma P$

22. (c) In adiabatic process, no heat transfers between system and

23. (b) $\frac{P_2}{P_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} \Rightarrow P_2 = P_1 \left(\frac{V_1}{V_2}\right)^{\gamma} = P_0(8)^{4/3} = 16P_0.$

24. (c) In adiabatic process $PV^{\gamma} = \text{constant}$

 $\Rightarrow \left(\frac{RT}{V}\right) V^{\gamma} = \text{constant} \Rightarrow TV^{\gamma-1} = \text{constant}$

25. (a) $TV^{\gamma-1} = \text{constant} \Rightarrow \frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} \Rightarrow T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma}$ $\Rightarrow T_2 = 300 \left(\frac{1}{2}\right)^{0.4} = 227.36 \text{ K}$

26. (b) In adiabatic change $Q = \text{constant} \Rightarrow \Delta Q = 0$ So $\Delta W = -\Delta U$ (: $\Delta Q = \Delta U + \Delta W$)

27. (d) For adiabatic process from FLOT $\Delta W = -\Delta U \qquad (\because \Delta Q = 0)$ $\Rightarrow \Delta W = -(-100) = +100 J$

28. (a) $\Delta U = -\Delta W = -\frac{R(T_1 - T_2)}{(\gamma - 1)} = \frac{R(T_2 - T_1)}{\gamma - 1}$

29. (d) $TV^{\gamma-1} = \text{constant} \Rightarrow T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma-1} = 927^{\circ} C$

30. (d) The process is very fast, so the gas fails to gain or lose heat. Hence this process in adiabatic

31. (a) $\Delta U = \mu C_V \Delta T = 1 \times C_V (T_f - T_i) = -C_V (T_i - T_f)$ $\Rightarrow |\Delta U| = C_V (T_f - T_f)$

32. (c) $T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma - 1} = 273(2)^{0.41} = 273 \times 1.328 = 363 K$ $W = \frac{R(T_1 - T_2)}{\gamma - 1} = \frac{8.31(273 - 363)}{1.41 - 1} = -1824$ $\Rightarrow |W| \approx 1815 J$

33. (d)

34. (c) $TV^{\gamma-1} = \text{constant}$

$$\Rightarrow T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma - 1} = (273 + 18) \left(\frac{V}{V/8}\right)^{0.4} = 668K$$

35. (a) $\Delta Q = mc\Delta\theta$. Here $\Delta Q = 0$, hence c = 0

36. (d) In adiabatic process, no transfer of heat takes place between system and surrounding.

37. (b) $W = \frac{\mu R(T_1 - T_2)}{(\gamma - 1)} = \frac{\mu RT_1}{(\gamma - 1)} \left[1 - \frac{T_2}{T_1} \right]$ $= \frac{\mu RT_1}{(\gamma - 1)} \left[1 - \left(\frac{V_1}{V_2} \right)^{\gamma - 1} \right]$ $= \frac{2 \times 8.31 \times 300}{\left(\frac{5}{2} - 1 \right)} \left[1 - \left(\frac{1}{2} \right)^{\frac{5}{3} - 1} \right] = +2767.23 J$

38. (d) $T^{\gamma}P^{1-\gamma} = \text{constant} \Rightarrow T \propto P^{\frac{\gamma-1}{\gamma}}$ $\Rightarrow \frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} = \left(\frac{1}{8}\right)^{\frac{5/3-1}{5/3}}$

$$T_2 = 300 \times \left(\frac{1}{8}\right)^{0.4} = 131K = -142^{\circ}C$$

39. (a) In adiabatic process $\Delta Q = 0 \Rightarrow \Delta U + \Delta W = 0$ $(\because \Delta Q = \Delta U + \Delta W)$

40. (d) Using relation
$$\frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} = (8)^{\frac{3/2-1}{3/2}} = 2$$
.

$$\Rightarrow T_2 = 2T_1 \Rightarrow T_2 = 2(273 + 27) = 600K = 327^{\circ}C$$

41. (c)
$$\frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} \Rightarrow \frac{T_2}{T_1} = \left(\frac{1}{8}\right)^{\frac{1.5-1}{1.5}} = \left(\frac{1}{8}\right)^{\frac{1}{3}} = \frac{1}{2}$$

$$\Rightarrow T_2 = \frac{T_1}{2} = \frac{300}{2} = 150K.$$

42. (c)
$$\frac{P_2}{P_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} \Rightarrow \frac{P'}{P} = (8)^{5/2} \Rightarrow P' = P \times (2)^{15/2}$$

43. (a)

44. (a) Given $P \propto T^3$, but we know for an adiabatic process, the pressure $P \propto T^{\gamma/\gamma-1}$

So
$$\frac{\gamma}{\gamma - 1} = 3 \Rightarrow \gamma = \frac{3}{2} \Rightarrow \frac{C_P}{C_V} = \frac{3}{2}$$

45. (b)

46. (d)
$$W = \frac{R(T_i - T_f)}{\gamma - 1} \Rightarrow 6R = \frac{R(T - T_f)}{\left(\frac{5}{3} - 1\right)} \Rightarrow T_f = (T - 4)K.$$

47. (a)
$$TV^{\gamma-1} = \text{constant}$$
 $T_1V_1^{\gamma-1} = T_2V_2^{\gamma-1}$
$$\Rightarrow T_2 = T_1 \left(\frac{V_1}{V_2}\right)^{\gamma-1} = T_1(4)^{1.5-1} = 2T_1$$

: change in temperature

$$=T_2-T_1=2T_1-T_1=T_1=273 K$$

48. (b)
$$\therefore PV^{\gamma} = k \text{ (constant)} \Rightarrow P_1V_1^{\gamma} = P_2V_2^{\gamma}$$

$$\Rightarrow P_2 = P_1 \left(\frac{V_1}{V}\right)^{\gamma} = 10^5 \times (2)^{1.3} \quad (\because V_2 = \frac{V_1}{2})$$

49. (b) In adiabatic process $\Delta U = -\Delta W$. In compression ΔW is negative, so ΔU is positive *i.e.* internal energy increases.

50. (a) According to the first law of thermodynamics $\Delta Q = \Delta U + \Delta W$

In adiabatic process $\Delta Q=0$, hence $\Delta U=-\Delta W$

51. (b)
$$PV^{\gamma} = \text{constant} \Rightarrow \frac{P_2}{P_1} = \left(\frac{V_1}{V_2}\right)^{\gamma} = \left(\frac{V_1}{V_1/4}\right)^{\gamma} = 4^{\gamma}$$

$$\Rightarrow P_2 = 4^{\gamma} P$$

As γ is always greater than one so $4^{\gamma} > 4 \implies P_2 > 4P$

52. (d)
$$P_1 V_1^{\gamma} = P_2 V_2^{\gamma} \Rightarrow \frac{P_2}{P_1} = \left[\frac{V_1}{V_2} \right]^{\gamma} = \left[\frac{4}{1} \right]^{3/2} = \frac{8}{1}$$

53. (b) Change in internal energy of the gas $\Delta U = -\Delta W \frac{R}{\gamma - 1} \left[T_2 - T_1 \right] = \frac{8.3}{(1.4 - 1)} [308 - 300] = 166J$

54. (b) For adiabatic change $TV^{\gamma-1}$ = constant

$$\Rightarrow \frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} \Rightarrow T_2 = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} \times T_1$$

$$\Rightarrow T_2 = \left(\frac{V}{V/4}\right)^{1.4 - 1} \times 300 = 300 \times (4)^{0.4} K$$

55. (d) For adiabatic forces $\Delta W = -\Delta U$ (: $\Delta Q = 0$) $\Rightarrow \Delta W = -(-50) = +50J$

56. (b) Adiabaticelasticity
$$(E_{\phi})$$

$$= \gamma \Rightarrow E_{\theta} = \frac{E_{\phi}}{\gamma}$$

$$\Rightarrow E_{\theta} = \frac{2.1 \times 10^{5}}{1.4} = 1.5 \times 10^{5} \, N/m^{2}$$

57. (c) $PV^{\gamma} = \text{constant}: \text{ Differentiating both sides}$ $P\gamma V^{\gamma-1} dV + V^{\gamma} dP = 0 \Rightarrow \frac{dP}{P} = -\gamma \frac{dV}{V}$

Isobaric and Isochoric Processes

1. (a) Work done $= P\Delta V = P(V_2 - V_1)$

2. (c) When heat is supplied at constant pressure, a part of it goes in the expansion of gas and remaining part is used to increase the temperature of the gas which in turn increases the internal energy.

3. (c) For isobaric process
$$\frac{V_2}{V_1}=\frac{T_2}{T_1}\Rightarrow V_2=V\times\frac{274}{273}$$
 Increase $=\frac{274\ V}{273}-V=\frac{V}{273}$

4. (a) From FLOT $\Delta Q = \Delta U + \Delta W = \Delta U + P\Delta V$ $\Rightarrow 100 = \Delta U + 50 \times (4 - 10) \Rightarrow \Delta U = 400 J$

5. (a) $W = P \times \Delta V = 2 \times 10^5 (150 - 50) \times 10^{-3} = 2 \times 10^4 J$

6. (c) $W = P\Delta V = nR\Delta T = 0.1 \times 2 \times 300 = 60 \ cal$

7. (c) $\Delta Q = \Delta V + P\Delta V \Rightarrow mL = \Delta U + P(V - V)$ $\Rightarrow \Delta U = L - P(V - V)$ (: m = 1)

8. (b) $\Delta W = P\Delta V = 10^3 \times 0.25 = 250 J$

9. (d)
$$W = P\Delta V = 1.01 \times 10^5 (3.34 - 2 \times 10^{-3})$$

= $337 \times 10^3 J \approx 340 \text{ KJ}$

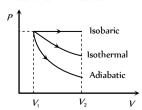
10. (c)
$$\frac{T_2}{T_1} = \frac{V_2}{V_1} = 2 \Rightarrow T_2 = 2 \times T_1 = 2 \times 300 = 600 \ K = 327^{\circ} \ C$$

11. (c) $V \propto T$ at constant pressure $\Rightarrow \frac{V_1}{V_2} = \frac{T_1}{T_2} \Rightarrow V_2 = \frac{V_1 T_2}{T_1} = \frac{300 \times 280}{300} = 280 \, ml.$

12. (a) In thermodynamic process, work done is equal to the area covered by the PV curve with volume axis.

Hence, according to graph shown

$$W_{adiabatic} < W_{isothermal} < W_{isobaric}$$



- 13. (b) (Similar to previous question)
- **14.** (a)
- 15. (d) $W = P\Delta V = 2.4 \times 10^{-4} \times 1 \times 10^{5} = 24/$
- **16.** (b) At constant pressure

$$W = P\Delta V = \mu R\Delta T = 1 \times 8.31 \times 100 = 831 \approx 814J$$

- 17. (a) $\Delta V = 0 \Rightarrow P\Delta V = 0 \Rightarrow \Delta W = 0$
- **18.** (d) Entropy of a reversible process does not change.
- 19. (c) $W = P\Delta V = 0$

$$(As \Delta V = 0)$$

- **20.** (d)
- **21.** (d) At constant volume $P \propto T \Rightarrow \frac{P_1}{P_2} = \frac{T_1}{T_2} \Rightarrow \frac{P_1}{P_2} = \frac{300}{400} = \frac{3}{4}$
- **22.** (d) In isothermal process $\Delta Q \neq 0$.
- **23.** (a) For isochoric process $\Delta V = 0 \Rightarrow \Delta W = 0$

From FLOT $\Delta Q = \Delta U + \Delta W \Rightarrow \Delta Q = \Delta U$

Heat Engine, Refrigerator and Second Law of Thermodynamics

1. (d)
$$\eta = \frac{T_1 - T_2}{T_1} - \frac{W}{Q} \Rightarrow Q = \left(\frac{T_1}{T_1 - T_2}\right) W$$

$$= \frac{600}{(600 - 300)} \times 800 = 1600 J$$

2. (c) Coefficient of performance

$$K = \frac{T_2}{T_1 - T_2} = \frac{273}{303 - 273} = \frac{273}{30} = 9$$

- **3.** (b) In a refrigerator, the heat dissipated in the atmosphere is more than that taken from the cooling chamber, therefore the room is heated if the door of a refrigerator is kept open.
- 4. (c) Internal energy is a state function.
- **5.** (b)
- **6.** (d) For a reversible process $\int \frac{dQ}{T} = 0$
- 7. (b) For cyclic forces $\Delta U = 0$ So, $\Delta Q = \Delta W$

8. (d)
$$\eta = 1 - \frac{T_2}{T_1} = 1 - \frac{400}{500} = \frac{1}{5} : : \eta = \frac{W}{Q} \implies \frac{1}{5} = \frac{W}{Q}$$

$$\Rightarrow W = \frac{Q}{5} = \frac{6}{5} \times 10^4 = 1.2 \times 10^4 J$$

9. (b)
$$\eta = 1 - \frac{T_2}{T_1} \Rightarrow \frac{30}{100} = 1 - \frac{350}{T_1}$$

 $\Rightarrow \frac{350}{T_1} = 1 - \frac{50}{100} = \frac{70}{100} = \frac{7}{10} \Rightarrow T_1 = 500 K = 227^{\circ}C$

10. (b)
$$\eta = 1 - \frac{T_2}{T_1}$$
 for 100% efficiency $\eta = 1$ which gives $T = 0$ K.

11. (c)
$$\eta = 1 - \frac{T_2}{T_1} = 1 - \frac{(273 + 69)}{(273 + 411)} = 0.5$$

 \Rightarrow Work done = $\eta \times Q = 0.5 \times 1000 = 500 J$

12. (b)
$$: \eta = 1 - \frac{T_2}{T_1} = \frac{W}{Q_1} = \frac{Q_1 - Q_2}{Q_1}$$

where Q_1 = heat absorbed, Q_2 = heat rejected

$$\Rightarrow 1 - \frac{T/3}{T} = \frac{W}{Q_1} \Rightarrow \frac{2}{3} = \frac{W}{Q_1} = \frac{Q_1 - Q_2}{Q_1}$$

$$\Rightarrow \frac{2}{3} = 1 - \frac{Q_2}{Q_1} \Rightarrow \frac{Q_2}{Q_1} = \frac{1}{3} \Rightarrow Q_2 = \frac{Q_1}{3} = \frac{Q}{3}$$

13. (c)
$$\eta = 1 - \frac{T_2}{T_1} \Rightarrow \frac{25}{100} = 1 - \frac{300}{T_1} \Rightarrow \frac{1}{4} = 1 - \frac{300}{T_1}$$

$$T_1 = 400 K = 127^{\circ}C$$

14. (a)
$$\eta = 1 - \frac{T_2}{T_1} \Rightarrow \frac{70}{100} = 1 - \frac{T_2}{1000} \Rightarrow T_2 = 300 \text{ K}$$

15. (a)
$$\eta = 1 - \frac{T_2}{T_1} = \frac{T_1 - T_2}{T_1} \implies \eta_1 = \frac{(473 - 273)}{473} = \frac{200}{473}$$

and
$$\eta_2 = \frac{273 - 73}{273} = \frac{200}{273}$$

So required ratio
$$\frac{\eta_1}{\eta_2} = \frac{273}{473} = 0.577$$

16. (a)
$$\eta = 1 - \frac{T_2}{T_1} = 1 - \frac{(273 + 123)}{(273 + 27)} = 1 - \frac{150}{300} = \frac{1}{2} = 50\%$$

17. (b)
$$\eta = 1 - \frac{T_2}{T_1} = 1 - \frac{300}{500} = \frac{2}{5}$$

- **18.** (d
- 19. (b) In first case, $(\eta_1) = 1 \frac{500}{800} = \frac{3}{8}$

and in second case, $(\eta_2) = 1 - \frac{600}{x}$

Since
$$\eta_1 = \eta_2$$
, therefore $\frac{3}{8} = 1 - \frac{600}{r}$

or
$$\frac{600}{x} = 1 - \frac{3}{8} = \frac{5}{8}$$
 or $x = \frac{600 \times 8}{5} = 960K$

20. (a)
$$\eta_{\text{max}} = 1 - \frac{T_2}{T_1} = 1 - \frac{300}{400} = \frac{1}{4} = 25\%$$

So 26% efficiency is impossible

21. (b) In first case
$$\eta_1 = 1 - \frac{T_2}{T_1} = 1 - \frac{(273 + 0)}{(273 + 200)} = \frac{200}{473}$$

In second case
$$\eta_2 = 1 - \frac{(273 - 200)}{(273 + 0)} = \frac{200}{273}$$

$$\Rightarrow \frac{\eta_1}{\eta_2} = \frac{1}{\left(\frac{473}{273}\right)} = 1:1.73$$

22. (b)
$$\eta = 1 - \frac{T_2}{T} \Rightarrow \frac{1}{2} = 1 - \frac{500}{T} \Rightarrow \frac{500}{T} = \frac{1}{2}$$
(i)

$$\frac{60}{100} = 1 - \frac{T_2}{T_1} \Rightarrow \frac{T_2}{T_1} = \frac{2}{5}$$
(ii)

Dividing equation (i) by (ii), $\frac{500}{T_2'} = \frac{5}{4} \Rightarrow T_2' = 400K$

- **23.** (c)
- **24.** (a)

25. (b)
$$\eta = 1 - \frac{T_2}{T_1} = \frac{W}{Q} \Rightarrow W = \left(1 - \frac{T_1}{T_2}\right)Q = \left\{1 - \frac{(273 + 27)}{(273 + 627)}\right\}$$

$$\Rightarrow W = \left(1 - \frac{300}{900}\right) \times 3 \times 10^6 = 2 \times 10^6 \times 4.2 \ J = 8.4 \times 10^6 \ J$$

- **26.** (a
- **27.** (d) $\eta = 1 \frac{T_2}{T_1}$; for η to be max. ratio $\frac{T_2}{T_1}$ should be min.
- **28.** (a)
- **29.** (b) In first case $\eta_1 = \frac{T_1 T_2}{T_1}$ In second case $\eta_2 = \frac{2T_1 2T_2}{2T_2} = \frac{T_1 T_2}{T_2} = \eta$
- 30. (c) Coefficient of performance $K = \frac{T_2}{T_1 T_2} \Rightarrow 5 = \frac{(273 13)}{T_1 (273 13)} = \frac{260}{T_1 260}$ $\Rightarrow 5T_1 1300 = 260 \Rightarrow 5T_1 = 1560$ $\Rightarrow T_1 = 312K \rightarrow 39^{\circ}C$
- 31. (a) Coefficient of performance $K = \frac{T_2}{T_1 T_2}$ $= \frac{(273 23)}{(273 + 27) (273 23)} = \frac{250}{300 250} = \frac{250}{20} = 5$
- **32.** (a) $\eta = \frac{T_1 T_2}{T_1} = \frac{(273 + 727) (273 + 227)}{273 + 727} = \frac{1000 500}{1000} = \frac{1}{2}$
- 33. (c) $\eta = \frac{T_1 T_2}{T_1} = \frac{W}{Q} \Rightarrow W = \frac{Q(T_1 T_2)}{T_1}$ $= \frac{6 \times 10^4 \left[(227 + 273) (273 + 127) \right]}{(227 + 273)}$ $= \frac{6 \times 10^4 \times 100}{500} = 1.2 \times 10^4 cal$
- **34.** (d) Slow isothermal expansion or compression of an ideal gas is reversible process, while the other given process are irreversible in nature.

Critical Thinking Questions

 $\begin{tabular}{ll} \textbf{(d)} & Fraction of supplied energy which in creases the internal energy is given by \\ \end{tabular}$

$$f = \frac{\Delta U}{(\Delta Q)_P} = \frac{(\Delta Q)_V}{(\Delta Q)_P} = \frac{\mu C_V \Delta T}{\mu C_P \Delta T} = \frac{1}{\gamma}$$

For diatomic gas $\gamma = \frac{7}{5} \Rightarrow f = \frac{5}{7}$

2. (c) $\Delta Q = \Delta U + \Delta W$ $\therefore \Delta U = \Delta Q - \Delta W = 540 - \frac{P(V_2 / V_1)}{J}$

$$=540 - \frac{1.013 \times 10^5 \times [(1671 - 1) \times 10^{-6}]}{4.2}$$

$$= 540 - 39.7 = 500$$
 calories

(b,c) There is a decrease in volume during melting on an ice slab at 273K. Therefore, negative work is done by ice-water system on the atmosphere or positive work is done on the ice-water system by the atmosphere. Hence option (b) is correct. Secondly heat is absorbed during melting (i.e. ΔQ is positive) and as we have seen, work done by ice-water system is negative (ΔW is negative). Therefore, from first law of thermodynamics $\Delta U = \Delta Q - \Delta W$.

Change in internal energy of ice-water system, ΔU will be positive or internal energy will increase.

4. (c) Process is isothermal. There fore, T = constant,

 $\left(P \propto \frac{1}{V}\right)$ volume is increasing, therefore pressure will

In chamber A:

$$\Delta P = P_i - P_f = \frac{\mu_A RT}{V} - \frac{\mu_A RT}{2V} = \frac{\mu_A RT}{2V} \qquad(i)$$

In chamber B:

$$1.5\Delta P = P_i - P_f = \frac{\mu_B RT}{V} - \frac{\mu_B RT}{2V} = \frac{\mu_B RT}{2V} \qquad(ii)$$

from equations (i) and (ii) $\frac{\mu_A}{\mu_B} = \frac{1}{1.5} = \frac{2}{3}$

$$\Rightarrow \frac{m_A / M}{m_B / M} = \frac{2}{3} \Rightarrow 3m_A = 2m_B.$$

- $\mathbf{5.} \qquad \text{(d)} \quad T_1 V_1^{\gamma 1} = T_2 V_2^{\gamma 1} \Rightarrow \frac{T_1}{T_2} = \left(\frac{V_2}{V_1}\right)^{\gamma 1} = \left(\frac{L_2 A}{L_1 A}\right)^{\frac{5}{3} 1} = \left(\frac{L_2}{L_1}\right)^{\frac{2}{3}}$
- **5.** (d) Using Boyle's law, we have $\frac{V}{T} = \text{constant}$

$$\Rightarrow \frac{\frac{l}{2} + 5}{373} = \frac{\frac{l}{2} - 5}{273}$$

As the piston moves 5 *cm*, the length of one side will be $\left(\frac{l}{2} + 5\right)$ and other side $\left(\frac{l}{2} - 5\right)$. On solving this equation, we get l = 64.6 *cm*.

(c) $\Delta Q = \Delta U + \Delta W \Rightarrow \Delta W = (\Delta Q)_P - \Delta U = (\Delta Q)_P \left| 1 - \frac{(\Delta Q)_V}{(\Delta Q)_P} \right|$ 7.

$$= (\Delta Q)_P \left[1 - \frac{C_V}{C_P} \right] = Q = \left[1 - \frac{3}{5} \right] = \frac{2}{5} Q$$

 $\therefore (\Delta Q)_P = Q$ and $\gamma = \frac{5}{3}$ for monatomic gas

8. (d) Oxygen is diatomic gas, hence its energy of two moles $=2\times\frac{5}{2}RT=5RT$

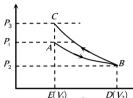
> Argon is a monoatomic gas, hence its internal energy of 4 moles = $4 \times \frac{3}{2}RT = 6RT$

Total Internal energy = (6+5)RT = 11RT

(c) From graph it is clear that $P_3 > P_1$. 9.

> Since area under adiabatic process (BCED) is greater than that of isothermal process (ABDE). Therefore net work done

$$W = W_i + (-W_A) :: W_A > W_i \implies W < 0$$



(a) According to given Vander Waal's equation 10.

$$P = \frac{nRT}{V - n\beta} - \frac{\alpha n^2}{V^2}$$

Work done,
$$W = \int_{V_1}^{V_2} P dV = nRT \int_{V_1}^{V_2} \frac{dV}{V - n\beta} - cn^2 \int_{V_1}^{V_2} \frac{dV}{V^2}$$

$$= nRT \left[\log_e (V - n\beta) \right]_{V_1}^{V_2} + cn^2 \left[\frac{1}{V} \right]_{V_1}^{V_2}$$

$$= nRT \log_e \frac{V_2 - n\beta}{V_1 - n\beta} + cn^2 \left(\frac{V_1 - V_2}{V_1 V_2} \right)$$

(b) Volume of the gas is constant $V = \text{constant} : P \propto T$ 11. i.e., pressure will be doubled if temperature is doubled

$$\therefore P = 2P_0$$

Now let *F* be the tension in the wire. Then equilibrium of any one piston gives $P_{QA} = P_{QA}$



$$F = (P - P_0)A = (2P_0 - P_0)A = P_0A$$

(c) $dU = C_V dT = \left(\frac{5}{2}R\right) dT$ or $dT = \frac{2(dU)}{5R}$ 12.

From first law of thermodynamics

$$dU = dQ - dW = Q - \frac{Q}{4} = \frac{3Q}{4}$$
. Now molar heat capacity
$$C = \frac{dQ}{dT} = \frac{Q}{\frac{2(dU)}{5R}} = \frac{5RQ}{2\left(\frac{3Q}{4}\right)} = \frac{10}{3}R$$
.

 $Q = \Delta U = U_f - U_i$ = [internal energy of 4 moles of a 13. monoatomic gas + internal energy of 2 moles of a diatomic gas [internal energy of 4 moles of a diatomic gas]

$$= \left(4 \times \frac{3}{2}RT + 2 \times \frac{5}{2}RT\right) - \left(4 \times \frac{5}{2}RT\right) = RT$$

Note: (a) 2 moles of diatomic gas becomes 4 moles of a monoatomic gas when gas dissociated into atoms.

(b) Internal energy of μ moles of an ideal gas of degrees of freedom F is given by $U = \frac{f}{2} \mu RT$

F = 3 for a monoatomic gas and 5 for diatomic gas.

- (c) $PV^{\gamma} = K \text{ or } P^{\gamma} V^{\gamma-1} dV + dP$, $V^{\gamma} = 0$ or $\frac{dP}{P} = -\gamma \frac{dV}{V}$ or $\frac{dP}{P} \times 100 = -\gamma \left(\frac{dV}{V} \times 100 \right)$
- (a) $TV^{\gamma-1} = \text{consta}$ 15. $\therefore \frac{T_1}{T_2} = \left(\frac{V_2}{V_1}\right)^{\gamma - 1} \text{ or } \left(\frac{1}{2}\right)^{\gamma - 1} = \sqrt{\frac{1}{2}}$
 - $\therefore \gamma 1 = \frac{1}{2} \text{ or } \gamma = \frac{3}{2}$ $\therefore PV^{3/2} = \text{constant}$
- (c) $\eta_A = \frac{T_1 T_2}{T} = \frac{W_A}{Q} \Rightarrow \eta_B = \frac{T_2 T_3}{T_1} = \frac{W_B}{Q_2}$ $\therefore \ \frac{Q_1}{Q_2} = \frac{T_1}{T_2} \times \frac{T_2 - T_3}{T_1 - T_2} = \frac{T_1}{T_2} \quad \therefore \ W_A = W_B$ $T_2 = \frac{T_1 + T_3}{2} = \frac{800 + 300}{2} = 550K$
- 17. $\gamma = \frac{C_P}{C_V} = \frac{5}{3}$ we know $\Delta Q = nC_P \Delta T$ and $\Delta U = nC_V \Delta T \Rightarrow \frac{\Delta U}{\Delta Q} = \frac{C_V}{C_P} = \frac{3}{5}$

i.e. fraction of heat energy to increase the internal energy be

- (a) $\Delta Q = \Delta U + \Delta W \Rightarrow \frac{\Delta W}{\Delta Q} = 1 \frac{\Delta U}{\Delta Q} = 1 \frac{nC_V dT}{nC_D dT}$ 18. $\Rightarrow \frac{\Delta W}{\Delta Q} = 1 - \frac{C_V}{C_R} = 1 - \frac{3}{5} = \frac{2}{5} = 0.4$
- (b) $\Delta U = \mu C_V \Delta T = \frac{m}{M} C_V \Delta T = \frac{N}{N} C_V \Delta T$ 19. $\Rightarrow (\Delta U)_N = \frac{56 \times 10^3}{14} \times \frac{5}{2} R \times 300$ and $(\Delta U)_A = \frac{6 \times 10^{26}}{6 \times 10^{23}} \times \frac{3}{2} R \times 900 \Rightarrow (\Delta U)_N > (\Delta U)_A$
- 20. $P_1 \cdot V(i) = P_2 \cdot \frac{V}{2} \Rightarrow P_2 = 2P_1$

and B is compressed adiabatically, hence

$$P_1 V^{\gamma} = P_2 \left(\frac{V}{2}\right)^{\gamma} \Rightarrow P_2 = (2)^{\gamma} P_1$$

Since $\gamma > 1$, hence P_2 '> P_2 or $P_2 < P_2'$

(b) In isothermal process $P_1V_1 = P_2V_2$ 21.

or
$$PV = P_2 \times 4V$$
 $\therefore P_2 = \frac{P}{A}$

In adiabatic process

$$P_2 V_2^{\gamma} = P_3 V_3^{\gamma} \Rightarrow \frac{P}{4} \times (4V)^{1.5} = P_2 V^{1.5} \Rightarrow P_3 = 2P_3 + P_3 + P_$$

- **22.** (d) Volume of the ideal gas is constant so $W = P\Delta V = 0$ using FLOT $\Delta Q = \Delta U \Rightarrow \Delta U = i^2 Rt = 1^2 \times 100 \times 5 \times 60$ $= 30 \times 10^3 = 30 KI$
- **23.** (d) Initially $\eta = \left(1 \frac{T_2}{T_1}\right) = \frac{W}{Q} = \frac{1}{6}$...(i) Finally $\eta' = \left(1 \frac{T_2}{T_1}\right) = \left(1 \frac{(T_2 62)}{T_1}\right) = 1 \frac{T_2}{T_1} + \frac{62}{T_1}$ (ii) $= \eta + \frac{62}{T_1}$ (iii)

It is given that $\eta' = 2\eta$. Hence solving equation (i) and (ii) $\Rightarrow T_1 = 372 \ K = 99^{\circ}C$ and $T_2 = 310 \ K = 37^{\circ}C$

24. (b) Input energy = $\frac{1g}{\sec} \times \frac{2kcal}{g} = 2kcal/\sec$.

Output energy = $10 \ KW = 10 \ K \ J/S = \frac{10}{4.2} kcal/sec$.

 $\Rightarrow \eta = \frac{\text{output energy}}{\text{input energy}} = \frac{10}{4.2 \times 2} > 1, \text{ it is impossible.}$

25. (b) Gain of entropy of ice

$$S_1 = \frac{\Delta Q}{T} = \frac{mL}{T} = \frac{80 \times 100}{(0 + 273)} = \frac{8 \times 10^3}{273} cal/K$$

Loss of entropy of water = $S_2 = -\frac{\Delta Q}{T} = -\frac{mL}{T}$

$$= \frac{80 \times 100}{(273 + 50)} = \frac{8 \times 10^3}{323} cal/K$$

Total change of entropy

$$S_1 + S_2 = \frac{8 \times 10^3}{273} - \frac{8 \times 10^3}{323} = +4.5 \ cal/K$$

- 26. (b) $PV^2=$ constant represents adiabatic equation. So during the expansion of ideal gas internal energy of gas decreases and temperature falls.
- 27. (d) Initially $\eta = \frac{T_1 T_2}{T_1} \Rightarrow 0.5 = \frac{T_1 (273 + 7)}{T_1}$ $\Rightarrow \frac{1}{2} = \frac{T_1 280}{T_1} \Rightarrow T_1 = 560K$ Finally $\eta_1' = \frac{T_1' T_2}{T_1'} \Rightarrow 0.7 = \frac{T_1' (273 + 7)}{T_1'} \Rightarrow T_1' = 933K$

 \therefore increase in temperature = $933 - 560 = 373K \approx 380K$

28. (c) *P-V* diagram of the gas is a straight line passing through origin. Hence $P \propto V$ or $PV^{-1} = \text{constant}$

Molar heat capacity in the process $PV^x = \text{constant is}$ R R

$$C = \frac{R}{\gamma - 1} + \frac{R}{1 - x}$$
; Here $\gamma = 1.4$ (For diatomic gas)

$$\Rightarrow C = \frac{R}{1.4 - 1} + \frac{R}{1 + 1} \Rightarrow C = 3R$$

29. (c) As finally the piston is in equilibrium, both the gases must be at same pressure P_f . It is given that displacement of piston be in final state x and if A is the area of cross-section of the piston. Hence the final volumes of the left and right part finally can be given by figure as

$$V_L = \frac{V_0}{2} + Ax \text{ and } V_R = \frac{V_0}{2} - Ax$$

As it is given that the container walls and the piston are adiabatic in left side and the gas undergoes adiabatic expansion and on the right side the gas undergoes adiabatic compressive. Thus we have for initial and final state of gas on left side

$$P_{1}\left(\frac{V_{0}}{2}\right)^{\gamma} = P_{f}\left(\frac{V_{0}}{2} + Ax\right)^{\gamma}$$
(i)

Similarly for gas in right side, we have

$$P_2 \left(\frac{V_0}{2}\right)^{\gamma} = P_f \left(\frac{V_0}{2} - Ax\right)^{\gamma}$$
(ii)

From eq. (i) and (ii)

$$\frac{P_1}{P_2} = \frac{\left(\frac{V_0}{2} + Ax\right)^{\gamma}}{\left(\frac{V_0}{2} - Ax\right)^{\gamma}} \implies Ax = \frac{V_0}{2} \begin{bmatrix} P_1^{1/\gamma} - P_2^{1/\gamma} \\ P_1^{1/\gamma} + P_2^{1/\gamma} \end{bmatrix}$$

Now from equation (i) $P_f = \frac{P_1 \left(\frac{V_0}{2}\right)^{\gamma}}{\left[\frac{V_0}{2} + Ax\right]^{\gamma}}$

30. (d) In both cylinders A and B the gases are diatomic ($\gamma = 1.4$). Piston A is free to move i.e. it is isobaric process. Piston B is fixed i.e. it is isochoric process. If same amount of heat ΔQ is given to both then

 $(\Delta Q)_{\text{isobaric}} = (\Delta Q)_{\text{isochoric}} \Rightarrow \mu C_p(\Delta T)_A = \mu C_v(\Delta T)_B$

$$\Rightarrow (\Delta T)_B = \frac{C_p}{C_p} (\Delta T)_A = \gamma (\Delta T)_A = 1.4 \times 30 = 42 \text{ K}.$$

Graphical Questions

- (c) As internal energy is a point function therefore change in internal energy does not depends upon the path followed i.e. $\Delta U_{\rm I} = \Delta U_{\rm II}$
- **2.** (b) Work done by the system = Area of shaded portion on *P-V* diagram

$$=(300-100)10^{-6}\times(200-10)\times10^{3}=20 J$$

- 3. (a) Work done = Area enclosed by triangle $ABC = \frac{1}{2}AC \times BC = \frac{1}{2} \times (3V V) \times (3P P) = 2PV$
- **4.** (c) Area enclosed between *a* and *f* is maximum. So work done in closed cycles follows *a* and *f* is maximum.
- **5.** (a) Initial and final states are same in all the process.

Hence $\Delta U = 0$; in each case.

By FLOT; $\Delta Q = \Delta W = \text{Area enclosed by curve with volume axis.}$ $\therefore \text{ (Area)} < \text{ (Area)} < \text{ (Area)} \Rightarrow Q < Q < Q.$

6. (a) For an isothermal process PV = constant

$$\Rightarrow PdV + VdP = 0 \Rightarrow -\frac{1}{V} \left(\frac{dV}{dP} \right) = \frac{1}{P}$$

So, $\beta = \frac{1}{D}$: graph will be rectangular hyperbola.

7. (a) By adjoining graph $W_{AB}=0$ and

$$W_{BC} = 8 \times 10^4 [5 - 2] \times 10^{-3} = 240 J$$

$$\therefore W_{AC} = W_{AB} + W_{BC} = 0 + 240 = 240 J$$

Now,
$$\Delta Q_{AC} = \Delta Q_{AB} + \Delta Q_{BC} = 600 + 200 = 800 J$$

From FLOT
$$\Delta Q_{AC} = \Delta U_{AC} + \Delta W_{AC}$$

$$\Rightarrow 800 = \Delta U_{AC} + 240 \Rightarrow \Delta U_{AC} = 560 J.$$

(b) In adiabatic process, slope of PV-graph. 8.

$$\frac{dP}{dV} = -\gamma \frac{P}{V} \Longrightarrow |\text{Slope}| \propto \gamma$$

From the given graph (Slope) > (Slope) $\Rightarrow \gamma_2 > \gamma_1$

therefore 1 should correspond to $O(\gamma = 1.4)$ and 2 should correspond to $He(\gamma = 1.66)$

As we know that slope of isothermal and adiabatic curves are 9. always negative and slope of adiabatic curve is always greater than that of isothermal curve

> Hence in the given graph curve A and B represents adiabatic and isothermal changes respectively.

- Process CD is isochoric as volume is constant, Process DA is 10. isothermal as temperature constant and Process AB is isobaric as pressure is constant.
- 11. (d) Heat given $\Delta Q = 20 \ cal = 20 \times 4.2 = 84 \ J$.

Work done $\Delta W = -50$ / [As process is anticlockwise] By first law of thermodynamics

$$\Rightarrow \Delta U = \Delta Q - \Delta W = 84 - (-50) = 134J$$

(a) For cyclic process. Total work done $= W_{AB} + W_{BC} + W_{CA}$ 12.

$$\Delta W_{M} = P\Delta V = 10(2-1) = 10J$$
 and $\Delta W_{K} = 0$

(as V = constant)

From FLOT, $\Delta Q = \Delta U + \Delta W$

 $\Delta U = 0$ (Process ABCA is cyclic)

$$\Rightarrow \Delta Q = \Delta W_1 + \Delta W_2 + \Delta W_3$$

$$\Rightarrow$$
 5 = 10 + 0 + $\Delta W \Rightarrow \Delta W = -5 J$

- (b) The cyclic process 1 is clockwise where as process 2 is 13. anticlockwise. Clockwise area represents positive work and anticlockwise area represents negative work. Since negative area (2) > positive area (1), hence net work done is negative.
- Process *AB* is isochoric, $W_{AB} = P \Delta V = 0$ 14.

 $\therefore W_{BC} = RT_2 \cdot \ln \left(\frac{V_2}{V_1} \right)$ Process BC is isothermal

Process CA is isobaric

$$W_{CA} = -P\Delta V = -R\Delta T = -R(T_1 - T_2) = R(T_2 - T_1)$$

(Negative sign is taken because of compression)

- AB is isobaric process, BC is isothermal process, CD is 15. isometric process and DA is isothermal process These process are correctly represented by graph (a).
- Work done by the gas (as cyclic process is clockwise) $\therefore \Delta W =$ 16. Area ABCD

So from the first law of thermodynamics ΔQ (net heat absorbed) = ΔW = Area ABCD

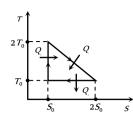
As change in internal energy in cycle $\Delta U = 0$.

(a) $Q_1 = T_0 S_0 + \frac{1}{2} T_0 S_0 = \frac{3}{2} T_0 S_0$ 17.

$$Q_2 = T_0 S_0 \text{ and } Q_3 = 0$$

$$\eta = \frac{W}{Q_1} = \frac{Q_1 - Q_2}{Q_1}$$

$$=1-\frac{Q_2}{Q_1}=1-\frac{2}{3}=\frac{1}{3}$$



(a) Work done = Area of closed PV diagram 18.

$$= (2V - V) \times (2P - P) = PV$$

From the given VT diagram,

In process AB, $V \propto T \Rightarrow$ Pressure is constant (As quantity of the gas remains same)

In process BC, V = Constant and in process CA,

T = constant

... These processes are correctly represented on PV diagram by graph (c).

(d) $\Delta Q = \Delta U + \Delta W$; ΔU does not depend upon path. 20.

$$\because \ \Delta W_A > \Delta W_B \ \Rightarrow \ \Delta Q_A > \Delta Q_B$$

- (d) Work done = Area under curve = $\frac{6P_1 \times 3V_1}{2}$ = 9 PV21.
- (d) Work done = $\frac{1}{2} \times 2P_1 \times 2V_1 = 2P_1V_1$ 22.
- (c) In a cyclic, $\Delta U = 0$ 23.

From FLOT, $\Delta Q = \Delta U + \Delta W = 0 + \Delta W = \text{Area of closed curve}$

$$\Rightarrow \Delta Q = \pi r \ \pi \left(\frac{20}{2}\right)^2 k P_a \times litre$$

$$=100 \pi \times 10^3 \times 10^{-3} J = 100 \pi J$$

- The work done in cyclic process is equal to the area enclosed 24. by the PV diagram
- In all given cases, process is cyclic and in cyclic process $\Delta U = 0$. 25.
- In cyclic process ΔQ = Work done = Area inside the closed 26.

Treat the circle as an ellipse of area $=\frac{\pi}{4}(P_2-P_1)(V_2-V_1)$

$$\Rightarrow \Delta Q = \frac{\pi}{4} \{ (150 - 50) \times 10^3 \} = \frac{\pi}{2} J$$

(d) $W_{\text{ex}} = -$ Area of triangle $BCO = -\frac{P_0 V_0}{2}$ 27.

$$W_{\text{\tiny asso}}$$
 + Area of triangle $AOD = + \frac{P_0 V_0}{2}$

- (c) AD and BC represent adiabatic process (more slope) 28. AB and DC represent isothermal process (less slope)
- Work done = Area of curve enclosed 29.

(c) Work done = Area of PV graph (here trapezium) 30.

$$= \frac{1}{2} (1 \times 10^5 + 5 \times 10^5) \times (5 - 1) = 12 \times 10^5 J$$

(d) For path $ab: (\Delta U)_{ab} = 7000 J$ 31.

By using
$$\Delta U = \mu C_V \Delta T$$

$$7000 = \mu \times \frac{5}{2} R \times 700 \Rightarrow \mu = 0.48$$

For path ca:

$$(\Delta Q)_{ca} = (\Delta U)_{ca} + (\Delta W)_{ca} \qquad(i)$$

$$\therefore (\Delta U)_{ab} + (\Delta U)_{bc} + (\Delta U)_{ca} = 0$$

:.
$$7000 + 0 + (\Delta U)_{ca} = 0 \Rightarrow (\Delta U)_{ca} = -7000 J$$
(ii)

Also
$$(\Delta W)_{cq} = P_1(V_1 - V_2) = \mu R(T_1 - T_2)$$

$$=0.48 \times 8.31 \times (300-1000) = -2792.16 J$$
(iii)

on solving equations (i), (ii) and (iii)

$$(\Delta Q)_{ca} = -7000 - 2792.16 = -9792.16 J = -9800 J$$

32. (b) Work done = Area enclosed by indicator diagram

$$= \frac{1}{2} \times (3V - V)(4P - P) = 3PV$$

33. (d) ΔU , remains same for both path

For path *iaf*: $\Delta U = \Delta Q - \Delta W = 50 - 20 = 30J$.

For path $fi: \Delta U = -30 J$ and $\Delta W = -13 J$

$$\Rightarrow \Delta Q = -30 - 13 = -43 J$$
.

- **34.** (a) $\Delta E_{\rm int}=0$, for a complete cycle and for given cycle work done is negative, so from first law of thermodynamics Q will be negative *i.e.* Q<0.
- **35.** (d) Work done = Area enclosed by the curve

$$=\frac{1}{2}(3V-V)(2P-P)=PV$$

36. (d) W = Area bonded by the indicator diagram with V-axis)

$$= \frac{1}{2} (P_A + P_B)(V_B - V_A)$$

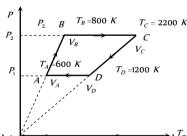
37. (a) Heat given $\Delta Q = 40 J$ and Work done $\Delta W = 30 J$

 $\Rightarrow \Delta U = \Delta Q - \Delta W = 40 - 30 = 10 J.$

- **38.** (a) As the volume is continuously increasing and the work of expansion is always positive, so the work done by the system continuously increases.
- **39.** (c) Processes *A* to *B* and *C* to *D* are parts of straight line graphs of the form v = mx

Also
$$P = \frac{\mu R}{V} T$$
 $(\mu = 6)$

 $\Rightarrow P \propto T$. So volume remains constant for the graphs AB and CD



So no work is done during processes for A to B and $\stackrel{T}{C}$ to D i.e., $\stackrel{W}{W} = 0$ and $\stackrel{W}{W} = P(V - V) = \mu R (T - T)$

$$=6R(2200-800)=6R\times1400J$$

Also
$$W_{a} = P(V - V) = \mu R(T - T)$$

$$=6R(600-1200)=-6R\times600J$$

Hence work done in complete cycle

$$W = W + W + W + W$$

$$= 0 + 6R \times 1400 + 0 - 6R \times 600$$

$$= 6R \times 900 = 6 \times 8.3 \times 800 \approx 40 \text{ k/}$$

40. (b) In isothermal process $P \propto \frac{1}{V}$.

Hence graph between P and V is a hyperbola.

- **41.** (d) Adiabatic curves are more stepper than isothermal curves.
- **42.** (d) During process A to B, pressure and volume both are decreasing. Therefore, temperature and hence, internal energy of the gas will decrease $(T \propto PV)$ or $\Delta U_{A \to B} =$ negative. Further $\Delta W_{A \to B}$ is also negative as the volume of the gas is decreasing. Thus $\Delta Q_{A \to B}$ is negative.

In process B to C pressure of the gas is constant while volume is increasing. Hence temperature should increase or $\Delta U_{B \to C}$ = positive. During C to A volume is constant while pressure is increasing. Therefore, temperature and hence, internal energy of the gas should increase or $\Delta U_{C \to A}$ = positive. During process CAB volume of the gas is decreasing. Hence, work done by the gas is negative.

43. (a) $\Delta W_{AB} = 0$ as V = constant

$$\therefore \Delta Q_{AB} = \Delta U_{AB} = 50J$$
 (Given)

$$U_A = 1500J$$
 : $U_B = (1500 + 50)J = 1550J$

$$\Delta W_{BC} = -\Delta U_{BC} = -40J \tag{Given}$$

$$\Delta U_{BC} = 40J$$
 $D_{C} = (1550 + 40)J = 1590J$

44. (a) For adiabatic process $T_1 V_b^{\gamma - 1} = \text{Constant}$

For *bc* curve
$$T_1 V_b^{\gamma - 1} = T_2 V_c^{\gamma - 1}$$
 or $\frac{T_2}{T_1} = \left(\frac{V_b}{V_c}\right)^{\gamma - 1}$ (i)

For ad curve
$$T_1 V_a^{\gamma - 1} = T_2 V_d^{\gamma - 1}$$
 or $\frac{T_2}{T_1} = \left(\frac{V_a}{V_d}\right)^{\gamma - 1}$ (ii)

From equation (i) and (ii)
$$\frac{V_b}{V_c} = \frac{V_a}{V_d}$$

Assertion and Reason

- 1. (a) In a perfectly reversible system, there is no loss of energy. Losses can be minimised, friction can be reduced, the resistance in L-C oscillating system can also be negligible. But one cannot completely eliminate energy losses. This makes a perfectly reversible system, an ideal.
- **2.** (a) Adiabatic expansion produces cooling.
- 3. (a) In reversible process, there always occurs some loss of energy. This is because energy spent in working against the dissipative force is not recovered back. Some irreversible process occur in nature such as friction where extra work to cancel the effect of friction. Salt dissolves in water but a salt does not separate by itself into pure salt and pure water.
- 4. (a) When a bottle of cold carbonated drink is opened. A slight fog forms around the opening. This is because of adiabatic expansion of gas causes lowering of temperature and condensation of water vapours.
- **5.** (e) As isothermal processes are very slow and so the different isothermal curves have different slopes so they cannot intersect each other.
- **6.** (d) Adiabatic compression is a rapid action and both the internal energy and the temperature increases.
- 7. (e) As there is no change in internal energy of the system during an isothermal change. Hence, the energy taken by the gas is

utilised by doing work against external pressure. According to FLOT $\Delta O = \Delta U + P\Delta V$

Hence $\Delta Q = \Delta U = P\Delta V$

can have values for 0 to ∞ .

Therefore, reason is true and assertion is false.

- **8.** (d) We can change the temperature of a body without giving (or taking) heat to (or from) it. For example in an adiabatic compression temperature rises and in an adiabatic expansion temperature false, although no heat is given or taken from the system in the respective changes.
- 9. (a) $c = \frac{Q}{m.\Delta\theta}$; a gas may be heated by putting pressure, so it

 C_P and C_V are it's two principle specific heats, out of infinite possible values.

In adiabatic process C = 0 and in isothermal process $C = \infty$.

- 10. (a) Heat is similar to work in that both represent ways of transferring energy. Neither heat nor work is an intrinsic property of a system, that is, we cannot say that a system contains a certain amount of heat or work.
- 11. (d) According to first law of thermodynamics, $\Delta Q = \Delta U + \Delta W = \Delta U + P\Delta V. \text{ If heat is supplies in such a}$ manner that volume does not change $\Delta V = 0$ *i.e.,* isochoric process, then whole of the heat energy supplied to the system will increase internal energy only. But, in any other process it is not possible.

Also heat may absorbed or evolved when state of thermal equilibrium changes.

- 12. (d) When the door of refrigerator is kept open, heat rejected by the refrigerator to the room will be more than the heat taken by the refrigerator from the room (by an amount equal to work done by the compressor). Therefore, temperature of room will increase and so it will be warmed gradually. As according to 2- law of thermodynamics, heat cannot be transferred on its own, from a body at lower temperature to another at higher temperature.
- (a) Second law of thermodynamics can be explained with the help of example of refrigerator, as we know that refrigerator, the working substance extracts heat from colder body and rejects a large amount of heat to a hotter body with the help of an external agency i.e., the electric supply of the refrigerator. No refrigerator can ever work without external supply of electric energy to it.
- 14. (d) If an electric fan is switched on in a closed room, the air will be heated because due to motion of the fan, the speed of air molecules will increase. In fact, we feel cold due to evaporation of our sweat.
- 15. (c) The internal energy of system depends only on its temperature. In isothermal process temperature does not change, therefore, internal energy of the system remains the same.
- 16. (c) In an adiabatic process, no exchange of heat is permissible i.e., $\Delta Q = 0 \; . \label{eq:deltaQ}$

As, $\Delta Q = \Delta U + \Delta W = 0 \implies \Delta U = -\Delta W$.

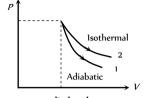
Also in adiabatic process, temperature of gas changes.

- 17. (a) Change in entropy, $\Delta S = \frac{\Delta Q}{T}$. In an adiabatic change, heat transfer $\Delta Q = 0$. \therefore $\Delta S = 0$, or S = constant i.e., entropy remains constant in an adiabatic process, or an adiabatic process is an isoentropic process.
- **18.** (b) As we know, in thermodynamic processes work done = Area covered by *P-V* diagram with volume axis.

Hence, according to following graph.

 $(Area) < (Area) \Rightarrow W < W$

Also in isothermal changes temperature remains same but in adiabatic changes temperature also changes.



- 19. (c) First law of thermodynamics is restatement of the
- principal of conservation of energy as applied to heat energy.

 20. (e) Zeroth law of thermodynamics explain the concept of temperature. According to which there exist a scalar quantity called temperature which is property of all thermodynamic system.
- **21.** (b) Efficiency of cannot cycle $\eta=\frac{W}{Q_1}=1-\frac{T_2}{T_1}$, for Carnot engine when T_2 decrease η increases.
- **22.** (a) Entropy is a measure of the disorder or randomness of the system. Greater the randomness, greater the entropy.

Self Evaluation Test -14

- The P-V diagram of 2 gm of helium gas for a certain process $A \rightarrow B$ is shown in the figure. What is the heat given to the gas during the process $A \rightarrow B$
 - (a) $4P_{o}V_{o}$
 - $6P_{o}V_{o}$
 - $4.5P_{o}V_{c}$
 - (d) $2P_{o}V_{o}$
- $$V_{\rm o}$$ $2\,V_{\rm o}$ V A certain mass of gas at 273 $\it K$ is expanded to 81 times its volume 2. under adiabatic condition. If $\gamma = 1.25$ for the gas, then its final temperature is [Pb. PET 1997]
 - (a) $-235^{\circ}C$
- (b) $-182^{\circ}C$
- (c) 91°C
- (d) 0°C
- In an adiabatic process 90/ of work is done on the gas. The change in internal energy of the gas is [CPMT 1996]
 - (a) -90 J
 - (b) +90 *]*

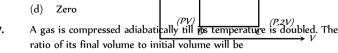
 - Depends on initial temperature
- If a Carnot's engine functions at source temperature 127°C and at sink temperature 87°C, what is its efficiency

[DCE 1997]

- (a) 10%
- (b) 25%
- (c) 40%
- (d) 50%
- In the case of diatomic gas, the heat given at constant pressure is that part of energy which is used for the expansion of gas, is

- An ideal monoatomic gas is taken round the cycle ABCDA shown in 6. the PV diagram in the given fig. The work done during the cycle is

 - (c)

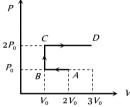


- 7.
 - (a) 1/2
- (b) More than 1 / 2
- (c) Less than 1 / 2
- (d) Between 1 and 2

- A tyre filled with air (27° C, and 2 atm) bursts, then what is temperature of air ($\gamma = 1.5$) [RPMT 2002]
 - (a) $-33^{\circ} C$
- (b) 0° C
- (c) 27° C
- (d) $240^{\circ} C$
- A gas expands adiabatically at constant pressure such that its 9. temperature $T \propto \frac{1}{\sqrt{V}}$, the value of C_P / C_V of gas is

[RPMT 2002; MHCET 2004]

- (a) 1.30
- (b) 1.50
- (c) 1.67
- (d) 2.00
- P-V diagram of an ideal gas is as shown in figure. Work done by the 10. gas in process ABCD is
 - (a) $4 P_0 V_0$
 - $2 P_0 V_0$
 - $3 P_0 V_0$
 - (d) P_0V_0



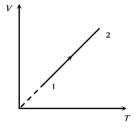
- 11. An engineer claims to have made an engine delivering 10 kW power with fuel consumption of $1 g s^{-1}$. The calorific value of fuel is 2kcallg. His claim [] & K CET 2000]
 - (a) Is non-valid
- (b) Is valid
- (c) Depends on engine
- (d) Depends on load
- An ideal gas heat engine operates in a Carnot cycle between 27°C 12. and 127°C. It absorbs 6 kcal at the higher temperature. The amount of heat (in kcal) converted into work is equal to
 - (a) 3.5

(b) 1.6

(c) 1.2

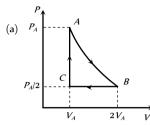
- (d) 4.8
- A gas expands with temperature according to the relation 13. $V = kT^{2/3}$. What is the work done when the temperature changes by 30° C
 - [**UPSEAT 1998**] (a)
- (b) 20 R
- (c) 30 R
- (d) 40 R
- An ideal gas ($\gamma = 1.5$) is expanded adiabatically. How many times 14. has the gas to be expanded to reduce the root mean square velocity of molecules 2.0 times
 - (a) 4 times
- (b) 16 times
- (c) 8 times
- (d) 2 times
- Three samples of the same gas A, B and $C(\gamma=3/2)$ have initially equal volume. Now the volume of each sample is doubled. The 15. process is adiabatic for A isobaric for B and isothermal for C. If the final pressures are equal for all three samples, the ratio of their initial pressures are

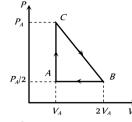
- (a) $2\sqrt{2}:2:1$
- (b) $2\sqrt{2}:1:2$
- (c) $\sqrt{2}:1:2$
- (d) $2:1:\sqrt{2}$
- **16.** Volume versus temperature graph of two moles of helium gas is as shown in figure. The ratio of heat absorbed and the work done by the gas in process 1-2 is
 - (a) 3
 - (b) $\frac{5}{2}$
 - (c) $\frac{5}{3}$
 - (d) $\frac{7}{2}$

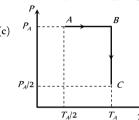


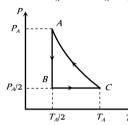
- 17. In the P-V diagram shown in figure ABC is a semicircle. The work done in the process ABC is
 - (a) Zero
- (b) $\frac{\pi}{2}atm-lt$
- (c) $-\frac{\pi}{2}atm-lt$
- (d) 4 *atm-lt*
- 1 2 V(litre)
- **18.** Heat is supplied to a diatomic gas at constant pressure. The ratio of $\Delta Q: \Delta U: \Delta W$ is
 - (a) 5:3:2
- (b) 5:2:3
- (c) 7:5:2
- (d) 7:2:5
- **19.** A gas undergoes a change of state during which 100 *J* of heat is supplied to it and it does 20 *J* of work. The system is brought back to its original state through a process during which 20 *J* of heat is released by the gas. The work done by the gas in the second process is
 - (a) 60 J
- (b) 40 *J*
- (c) 80 /
- (d) 20 *I*
- **20.** *N* moles of an ideal diatomic gas are in a cylinder at temperature *T*. suppose on supplying heat to the gas, its temperature remain constant but *n* moles get dissociated into atoms. Heat supplied to the gas is
 - (a) Zero
- (b) $\frac{1}{2}nRT$
- (c) $\frac{3}{2}nRT$
- (d) $\frac{3}{2}(N-n)RT$
- **21.** Three moles of an ideal gas $\left(C_P = \frac{7}{2}R\right)$ at pressure P_A and

temperature T_A is isothermally expanded to twice its initial volume. It is then compressed at constant pressure to its original volume. Finally the gas is compressed at constant volume to its original pressure P_A . The correct P-V and P-T diagrams indicating the process are









- **22.** A cylinder of mass 1kg is given heat of 20000 J at atmospheric pressure. If initially temperature of cylinder is $20^{\circ}C$, then work done by the cylinder will be (Given that Specific heat of cylinder = 400 J kg, Coefficient of volume expansion = $9 \times 10^{\circ}$ °C, Atmospheric pressure = $10 \ N/m$ and density of cylinder $9000 \ kg/m$)
 - (a) 0.02 /
- (b) 0.05 J
- (c) 0.08 J
- (d) 0.1 /
- 23. In a thermodynamic process pressure of a fixed mass of a gas is changed in such a manner that the gas releases 30 *joules* of heat and 10 joules of work was done on the gas. If the initial internal energy of the gas was 30 *joules*, then the final internal energy will be [CPMT 1986]
 - (a) 2 *J*

- (b) 18 *J*
- (c) 10 J
- (d) 58 J
- 24. In an adiabatic change, the pressure P and temperature T of a monoatomic gas are related by the relation $P \propto T^C$, where C equals [CBSE PMT 1994;

BHU 1997; AllMS 2001; MH CET 2000]

- (a) 5/3
- (b) 2/5
- (c) 3/5
- (d) 5/2
- 25. The internal energy of an ideal gas increases during an isothermal process when the gas is [SCRA 1998]
 - (a) Expanded by adding more molecules to it
 - (b) Expanded by adding more heat to it
 - (c) Expanded against zero pressure
 - (d) Compressed by doing work on it

Answers and Solutions

(SET -14)

I. (b) Change in internal energy from $A \rightarrow B$ is

 $\Delta U = \frac{f}{2} \mu R \Delta T = \frac{f}{2} (P_f V_f - P_i V_i)$

$$= \frac{3}{2}(2P_0 \times 2V_0 - P_0 \times V_0) = \frac{9}{2}P_0V_0$$

Work done in process $A \to B$ is equal to the Area covered by the graph with volume axis i.e.,

$$\begin{split} W_{A\to B} &= \frac{1}{2}(P_0 + 2P_0) \times (2V_0 - V_0) = \frac{3}{2}\,P_0\,V_0 \\ \text{Hence, } \Delta Q &= \Delta U + \Delta W = \frac{9}{2}\,P_0\,V_0 + \frac{3}{2}\,P_0\,V_0 = 6\,P_0\,V_0 \end{split}$$

2. (b) For adiabatic process $TV^{\gamma-1} = \text{constant}$

$$\Rightarrow \frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} \Rightarrow T_2 = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} \times T_1$$

$$\Rightarrow T_2 = \left(\frac{1}{81}\right)^{1.25 - 1} \times 273 = \left(\frac{1}{81}\right)^{0.25} \times 273$$

$$= \frac{273}{3} = 91K \rightarrow -182^{\circ}C$$

3. (b) For adiabatic process $\Delta Q=0$ From $\Delta Q=\Delta U+\Delta W\Rightarrow 0=\Delta U-90 \Rightarrow \Delta U=+90J$

4. (d)
$$\eta = \frac{T_1 - T_2}{T_1} = \frac{(127 + 273) - (87 + 273)}{(127 + 273)}$$

$$= \frac{400 - 360}{400} = 0.1 \rightarrow 10\%$$

5. (c) $\Delta W = \text{energy used for expansion} = PdV = RdT$ $\Delta Q = \text{heat supplied to diatomic gas at constant } P$ $= C_p dT = \frac{7}{2} RdT \quad (\because C_p = \frac{7}{2} R) \quad \because \frac{\Delta W}{\Delta Q} = \frac{RdT}{\frac{7}{2} RdT} = \frac{2}{7} RdT$

7. (c)
$$\frac{T_2}{T_1} = \left(\frac{V_1}{V_2}\right)^{\gamma - 1} = 2 \Rightarrow \left(\frac{V_2}{V_1}\right)^{\gamma - 1} = \frac{1}{2} \Rightarrow \frac{V_2}{V_1} = \left(\frac{1}{2}\right)^{\frac{1}{\gamma - 1}} < \frac{1}{2}$$

$$\Rightarrow V_2 < \frac{V_1}{2}$$

8. (a)
$$\frac{T_2}{T_1} = \left(\frac{P_2}{P_1}\right)^{\frac{\gamma-1}{\gamma}} \Rightarrow \frac{T_2}{(273+27)} = \left(\frac{1}{2}\right)^{\frac{1.5-1}{1.5}} = \left(\frac{1}{2}\right)^{\frac{1}{3}} = \frac{1}{2.5}$$

$$\Rightarrow T_2 = \frac{T_1}{1.25} = \frac{(273+27)}{1.25} = 238 K = -34.8^{\circ}C$$

9. (b) $TV^{\gamma-1}={\rm constant}\Rightarrow T\propto V^{1-\gamma}$ According to question $T\propto V^{-\frac{1}{2}}$ Hence $1-\gamma=-\frac{1}{2}\Rightarrow \gamma\,\frac{3}{2}=1.5$

10. (c) $W_{AB}=-P_0V_0$, $W_{BC}=0$ and $W_{CD}=4P_0V_0$ $\Rightarrow W_{ABCD}=-P_0V_0+0+4P_0V_0=3P_0V_0$

11. (a) Power = $10 \ KW = 10000 \ J/s = \frac{10000}{4.2} = 2.38 \ k \ cal/gm$

But the calorific value of fuel is only 2 $\it k$ $\it call gm$. Hence claim is invalid.

12. (c) Efficiency of a carnot engine is given by $\eta = 1 - \frac{T_2}{T_1}$ or $\frac{W}{Q} = 1 - \frac{T_2}{T_1} \Rightarrow \frac{W}{6} = 1 - \frac{(273 + 127)}{(273 + 227)} \Rightarrow W = 1.2 k cal$

13. (b) $W = \int P dV = \int \frac{RT}{V} dV$ Since $V = kT^{2/3} \implies dV = \frac{2}{3} KT^{-1/3} dT$ Eliminating K, we find $\frac{dV}{V} = \frac{2}{3} \frac{dT}{T}$ Hence $W = \int_{T_0}^{T_2} \frac{2}{3} \frac{RT}{T} dT = \frac{2}{3} R(T_2 - T_1) = \frac{2}{3} R(30) = 20 R$

14. (b)
$$v_{ms} = \sqrt{\frac{3RT}{M}} \implies v_{ms} \propto \sqrt{T}$$

 v_{mns} is to reduce two times *i.e.* temperature of the gas will have to reduce four times or $\frac{T'}{T} = \frac{1}{4}$

During adiabatic process $TV^{\gamma-1} = T'V'^{\gamma-1}$

$$\Rightarrow \frac{V'}{V} = \left(\frac{T}{T'}\right)^{\frac{1}{\gamma - 1}} = (4)^{\frac{1}{1.5 - 1}} = (4)^2 = 16 \Rightarrow V' = 16 V$$

15. (b) Let the initial pressure of the three samples be P_A , P_B and P_C , then $P_A(V)^{3/2}=(2V)^{3/2}P$, $P_B=P$ and $P_C(V)=P(2V)$ $\Rightarrow P_A:P_B:P_C=(2)^{3/2}:1:2=2\sqrt{2}:1:2$

16. (b) $V ext{-}T$ graph is a straight line passing through origin. Hence, $V \propto T$ or P = constant $\therefore \quad \Delta Q = nC_P \Delta T \quad \text{and} \quad \Delta U = nC_V \Delta T$ Also $\Delta W = \Delta Q - \Delta U = \mu (C_P - C_V) \Delta T$ $\therefore \quad \frac{\Delta Q}{\Delta W} = \frac{nC_P \Delta T}{n(C_P - C_V) \Delta T} = \frac{C_P}{C_P - C_V} = \frac{1}{1 - \frac{C_V}{C_P}}$ $\frac{C_V}{C_P} = \frac{3}{5} \quad \text{for helium gas. Hence} \quad \frac{\Delta Q}{\Delta W} = \frac{1}{1 - \frac{3}{5}} = \frac{5}{2}$

17. (b) W_{AB} is negative (volume is decreasing) and W_{BC} is positive (volume is increasing) and since, $\left|W_{BC}\right| > \left|W_{AB}\right|$

 \therefore net work done is positive and area between semicircle which is equal to $\frac{\pi}{2}atm-lt$.

3. (c)
$$\Delta Q = \mu C_P \Delta T = \frac{7}{2} \mu R \Delta T$$
 $\left(C_P = \frac{7}{2} R\right)$

$$\Delta U = \mu C_V \Delta T = \frac{5}{2} \mu R \Delta T \qquad \left(C_V = \frac{5}{2} R\right)$$
and $\Delta W = \Delta Q - \Delta U = \mu R \Delta T$

$$\Rightarrow \Delta Q : \Delta U : \Delta W = 7 : 5 : 2$$

19. (a) In a cyclic process
$$\Delta U = 0 \Rightarrow \Delta Q = \Delta W$$

$$\Rightarrow (100 - 20) = 20 + W_2 \Rightarrow W_2 = 60 J$$

20. (b) Since the gas is enclosed in a vessel, therefore, during heating process, volume of the gas remains constant. Hence, no work is done by the gas. It means heat supplied to the gas is used to increase its internal energy only.

Initial internal energy of the gas is
$$\ U_1 = N \left(\frac{5}{2} \, R \right) T$$

Since n moles get dissociated into atoms, therefore, after heating, vessel contains (N-n) moles of diatomic gas and 2n moles of a mono-atomic gas. Hence the internal energy for the gas, after heating, will be equal to

$$U_2 = (N - n) \left(\frac{5}{2}R\right) T + 2n \left(\frac{3}{2}R\right) T = \frac{5}{2}NRT + \frac{1}{2}nRT$$

Hence, the heat supplied = increase in internal energy

$$=(U_2-U_1)=\frac{1}{2}nRT$$

21. (a) Let the process start from initial pressure P_A , volume V_A and temperature T_A .

$$A(P_A, V_A, T_A) \longrightarrow B\left(\frac{P_A}{2}, 2V_A, T_A\right)$$

$$C\left(\frac{P_A}{2}, V_A, \frac{T_A}{2}\right)$$

- (i) Isothermal expansion ($PV = {\rm constant}$) at temperature T_A to twice the initial volume V_A
- (ii) Compression at constant pressure $\frac{P_A}{2}$ to original volume $V_A~(i.e.~V \propto T)$
- (iii) Isochoric process (at volume V_A) to initial condition (i.e. $P \propto T$)

22. (b)
$$\Delta Q = mc\Delta T \Rightarrow \Delta T = \frac{20000J}{1kg \times (400J/kg^{\circ}C)} = 50^{\circ}C$$

$$\Rightarrow T = 70^{\circ}C$$
Hence $W = P_{atm}\Delta V = P_{atm}V_{0}\gamma \Delta T$

=
$$(10^5 N/m^2) \left(\frac{1}{9 \times 10^3} m^3\right) (9 \times 10^{-5} / {}^{\circ}C) (50^{\circ}C) = 0.05 J$$

23. (c)
$$\Delta Q = \Delta U + \Delta W = (U_f - U_i) + \Delta W$$

 $\Rightarrow -30 = (U_f - 30) - 10 \Rightarrow U_f = 10 J$

24. (d)
$$T^{\gamma}P^{1-\gamma} = \text{constant} \Rightarrow P \propto T^{\frac{\gamma}{\gamma-1}}$$
Comparing above equation with given equation
$$P \propto T^{C} \Rightarrow C = \frac{\gamma}{\gamma-1} = \frac{5/3}{5/3-1} = \frac{5}{2}$$

25. (a) Internal energy of an ideal gas is given by

$$U = \frac{f}{2} \mu RT = \frac{f}{2} \left(\frac{N}{N_A} \right) RT \implies U \propto NT.$$

In isothermal process T = constant $\Rightarrow U \propto N$. *i.e.* internal energy increases by increasing number of molecules (N).