THERMODYNAMICS

CALORIMETRY

Calorimetry is the study of the measurement of quantities of heat

- HEAT:
 - It is the energy exhange between two systems due to the temperature difference.
 - S.I unit is joule
 - The C.G.S unit is calorie
 - The dimensional formula is [ML² T⁻²]
 - The amount of heat gained or lost depends upon the nature of material and its state.
 - The heat flows from higher temperature region to lower temperature region.
 - Calorie: It is the quantity of heat required to raise the temperature of 1g. of water from 14.5°C to 15.5°C.
 - 1calorie = 4.186J.
 - Heat is measured by using calorimeter

• SPECIFIC HEAT:

 It is the amount of heat required to raise the temperature of 1g of a substance

through 1°C.
$$s = \frac{1}{m} \frac{\Delta Q}{\Delta T}$$

- The S.I unit is Jkg⁻¹ K⁻¹
- The C.G.S unit is $Cal.g^{-1} \circ C^{-1}$
- Dimensional formula is $U^2T^{-2}V^{-1}$

S.No.	Substance	CGS Value Specific heat	S.I Value
1.	Water	1 cal g ^{-1 0} C ⁻¹	4186 JKg ⁻¹ K ⁻¹
2.	ice	0.5 cal g ^{-1 o} c ⁻¹	2100 Jkg ⁻¹ k ⁻¹

• HEATCAPACITY(OR) THERMAL CAPACITY:

• It is the amount of heat required to raise the temperature of the body by 1°C

$$c = \frac{\Delta Q}{\Delta T}$$

- The S.I.unit is JK⁻¹
- The C.G.S unit is Cal °C⁻¹
- Dimensional formula : ML² T⁻² K⁻¹
- Thermal capacity depends on mass and nature of substance.

• WATER EQUIVALENT:

- Mass of water which has same thermal capacity as that of the substance is called water equivalent
- It is numerically equal to heat capacity (ms)
- The S.I unit is kg
- The C.G.S unit is g

DULONG AND PETIT'S LAW :

The product of specific heat of an element in the solid state and its atomic weight is a constant and is approximately equal to 6.4 cal/gram atom / $^{\circ}C$

CHANGE OF STATE:

- Matter exists in three states or phases such as solid, liquid and gas.
- Matter can exist in all these three phases simultaneously at a particular pressure and temperature.
- Melting of ice or vapourisation of water is an example for change of state. During change of state, temperature remains constant.

MELTING:

- Every solid melts at a definite temperature called melting point.
- In case of substances which expand on freezing, the melting point is lowered due to increase of pressure.

Ex: Ice, cast Iron, type metal etc.,

- In case of substances which contract on freezing, the melting point rises due to increase of pressure.
 Ex: Brass, Cu.
- Addition of impurities (or) presence of impurities generally reduces the melting point of a substance.
- The freezing point, melting point and the boiling point of a substance depends onthe external pressure and on the purity of the substance.
- The melting point of the substance which contracts on liquification decreases with increase of pressure. Ex: cast iron, ice and type metal. the lowering of melting point of ice is 0.0072°C /atmosphere.
- The melting point of those substances which expand on liquification increases with the increase of pressure. Ex: wax, gold, silver, copper etc
- The melting of ice when pressure is appiled and resolidification on removal of pressure is called regelation.
- Snow-ball preparation is due to regelation.

BOILING AND EVAPORATION:

- It is the change of state from liquid to vapour at a particular temperature.
- Increase of pressure increases the boiling point of a liquid.
- On mountains, it is difficult to cook food because with increase in height, pressure decreases and hence the boiling point of water decreases.
- Boiling point of a liquid depends on nature of the liquid, applied pressure and presence of impurities.
 - the escape of molecules from the free

surface of a liquid is called evaporation

- Refrigerators, air coolers etc., work on the principle of cooling produced by evaporation.
- Body temperature is also controlled by evaporation of sweat
- Evaporation is a slow process where as boiling is a quick process
- Evaporation takes place at all temperatures where as boiling takes place at a particular temperature
- Evaporation takes place only at the surface of a liquid where as boiling occurs through out the liquid.
- The rate of evaporation depends on the nature of the liquid
- The rate of evaporation is more when the difference in temperatures of the liquid and the surrounding air is high

• SUPER HEATING OF A LIQUID :

A Liquid can be slowly heated beyond its boilingpoint.

In side a pressure cooker the boiling point of water is greater than 100°C

water free from dust and air can be heated up to 137°C with out changing to the vapour state. Super-heated water is unstable. stirring or even a slight mechanical disturbance causes the water to boil immediately.

HOAR-FROST AND SUBLIMATION :-

It is a process of conversion from gaseous state to solid state directly on cooling a material. The reverse process of the hoar -frost is sublimation.

• LATENT HEAT :

It is the amount of heat required to change unit mass of a substance from one state to another state without any change of temperature.

LATENT HEAT OF FUSION OF ICE:

- It is the amount of heat required to convert 1gram of ice at 0°c to 1 gram of water at 0°c.
- The S.I value is 0.335X10⁶ J/kg
- The C.G.S value is 80 cal/g

13. LATENT HEAT OF VAPOURISATION:

- It is the amount of heat required to convert 1gram of water at 100°c to 1gram of steam at 100°C.
- The S.I value is 2.26x10⁶ J/kg.
- The C.G.S value is 540 Cal/g
- Latent heat of vapourisation decreases with increase in temperature.
- Latent heat of a substance becomes zero at critical temperature..
- Latent heat depends on the nature of a substance and pressure.
- The product of the latent heat of fusion and the atomic weight of a substance is known as atomic latent heat.

LAW OF MIXTURES (OR) CALORIMETRY PRINICPLE:

When three substances of different masses m₁, m₂ and m₃ specific heats s₁, s₂, s₃ and at different temperatures t₁,t₂, and t₃ respectively are mixed, then the resultant temperature is

$$t = \frac{m_1 s_1 t_1 + m_2 s_2 t_2 + m_3 s_3 t_3}{m_1 s_1 + m_2 s_2 + m_3 s_3}$$

When ice at 0°C and steam at 100°C are mixed, mass of ice that melts is m_{ice} = 8 m_{steam}

When "x" gram of steam is mixed with "y" gram of ice, the resultant temperature is

$$t = \frac{80(8x - y)}{(x + y)}$$

 When ice and water are mixed the mass of ice that just melts is

$$m_{ice} = \frac{ms\theta}{80}$$

Where m= mass of water.

 When ice and steam are mixed, the amount of steam that just condenses into water at 100°C.

$$m_{steam} = \frac{m_{ice}}{3}$$

- When ice at 0°C and steam at 100°C are mixed then resultant temperature must be between 0°C and 100°C. The resultant temperature cannot be less than 0°C or more than 100°C.
- Steam is used in heat engines as working substance because of its high latent heat.
- Heavy water is used as coolant in nuclear reactors because of its high specific heat.
- In extinguishing fire hot water is preferred than cold water since hot water becomes vapour quickly and vapours do not allow fire.
- Inside a deep mine boiling point of water is greater than 100°C.
- Steam at 100°C produces more severe burns than water at 100°C as steam gives out more heat due to its latent heat.

CALORIFIC VAULE :

Calorific value of a fuel is the quantity of heat liberated when one gram of the fuel is burnt completely.

unit: cal/g or J/kg

It can be determined by using Bomb Calorimeter or Bell Calorimeter

Note : Calorific value of a food stuff is the quantity of heat liberated when a unit mass

of the food stuff is completely utilised by the body.

- THERMODYNAMICS: It is the branch of physics that deals with the conversion of heat into other forms of energy.
- SYSTEM:
 - A particular portion of matter or a restricted region of space under investigation is called system.
 - If the state of a system is represented by pressure (P), volume (V), temperature (T) and Entropy (S) then it is called a thermodynamic system.
- TYPES OF SYSTEMS:
 - Open system: It is the system in which both heat and matter can be exchanged with surroundings.
 - Ex: Plants, Animals etc.,
 - Closed system: It is the system in which only heat is exchanged with the surroundings.
 - Ex: An electric vapour lamp.
 - Isolated system: It is the system in which neither heat nor matter is exchanged with surroundings.
 Ex: Thermos Flask.
- **THERMAL EQUILIBRIUM:** If two bodies are at the same temperature then they are said to be in thermal equilibrium. When bodies are in equilibrium, no exchange of heat takes place
- **ZEROTH LAW OF THERMODYNAMICS:** If two bodies A and B are in thermal equilibrium with a third body C then the bodies A and B will also be in thermal equilibrium with each other. Zeroth law gives the concept of temperature.
- **INTERNAL WORK:** It is the work done by one part of a system on its another parts. Example: work done by a gaseous system against intermolecular forces.
- **EXTERNAL WORK:** When work is done by the system or on the system by the surroundings then it is called external work.
- WORKDONE BY A GAS DURING EXPANSION:
 - When a gas expands at constant pressure then for a small change in volume dv workdone is dw=pdv
 - If the volume changes from v₁ to v₂ at constant pressure p, the work done is w= p(v₂-v₁)
- SIGN CONVENTION:
 - If the work is done by the system , then work done is (+) ve.

 $(\therefore dv > 0, w > o)$

- If work is done by the system, heat is released and the temperature of the system decreases.
- If the work is done on the system, then it is (-) ve.

(: dv < o, w < o)

- If work is done on the system then heat is absorbed and consequently the temperature of the system increases.
- When work is done by the system against external pressure then dw=pdv

$$w = \int_{v_1}^{v_2} p_{\text{dv}}$$

- Area under p-v graph is equal to the work done
 INTERNAL ENERGY: It is the energy possessed by the system due to molecular motion and molecular configuration. Internal energy of a system is a thermodynamic function. It is of two types.
 - Internal Potential Energy: It is the energy present in a system on account of its molecular configuration is called internal potential energy.
 - Internal Kinetic Energy: It is the energy present in a system on account of motion of its molecules is defined as internal kinetic energy.
 - Change in internal engery

$$\Delta U = \Delta U_{K} + \Delta U_{P}$$

 The change in internal energy depends upon the initial and final states only.

 $\Delta U = U_{final} - U_{initial}$

- For an ideal gas since there are no intermolecular forces of attraction there is no potential energy. For such a gas the internal kinetic energy depends only on temperature.
- Internal energy of real gases depends upon temperature and volume.
- Change in internal energy is zero in a cyclic process.
- Real gases consist of both kinetic energy and potential energy due to intermoleculer forces.
- Ex: If a fan is turned on in a closed room then the temperature of the room increases because the molecular speeds increases.

FIRST LAW OF THERMODYNAMICS:

- All the heat added to a system is partly utilised to do the mechanical work and remaining to increase its internal energy.
- The differential form of first law of thermodynamics is dQ=dU+dW, where dQ = heat added, dU = Increase in internal energy.
 - dW= work done

- It defines the property of system called internal energy.
- It is a consequence of law of conservation of energy.
- SIGN CONVENTIONS:
 - When heat is added to the system dQ is + ve(+dQ)
 - When heat is taken from the system dQ is -ve (-dQ)
 - When gas expands work is done by the gas dw is positive (+ dW)
 - When gas contaracts work is done on the gas dw is negative.(-dW)
 - When internal energy of system increases and dU is +ve (+dU)
 - When internal energy of system decreases and dU is -ve(-dU)

 RELATION BETWEEN WORK AND HEAT (JOULE'SLAW): The amount of heat poduced is directly proportional to the amount of work done.

 $H\alpha W$ or W=JH.

 $J = \frac{W}{H}$ where J= Mechanical equivalent of heat.

• MECHANICAL EQUIVALENT OF HEAT (J):

- It is the amount of work necessary to produce unit amount of heat engegy.
- J is not a constant or physical quantity. It is simply a conversion factor which is used to convert joule or erg into calorie or kilocalorie.

• VALUES OF 'J':

- J = 4.186 J/cal =4.186X10 ⁷ erg/cal =4186 J/K.cal
- When heat and work are in Joule then J
 = 1
- The height from which ice is to be dropped to melt it completely is

h=
$$\frac{JL}{g}$$

where L= Latent heat of ice.

 The rise in temperature of water when it falls from a height h to the ground is,

$$\Delta \theta = \frac{gh}{Js}$$

where 's' is specific heat of water

 When a body of mass m moving with a velocity v is stopped and all of its energy is retained by it, then the increase in temperature is.

$$\Delta\theta = \frac{v^2}{2Js}$$

• When a block of ice of mass M is dragged with constant velocity on a rough horizontal surface of coefficient of friction

 μ , through a distance d, then the mass of ice melted is,

$$m = \frac{\mu Mgd}{JL}$$

Where m=mass of ice melted.

When a block of mass m is dragged on a rough horizontal surface of coefficient of friction µ, then the rise in temparature of block is,

$$\Delta \theta = \frac{\mu g d}{Js}$$

 If a bullet just melts when stopped by an obstacle and if all the heat produced is absorbed by the bullet then

$$ms\Delta\theta + mL = \frac{1}{2} \frac{mv^2}{J}$$

Where L= Latent heat of the material of the bullet

s= Specific heat

 A metal ball falls from a height 'h₁' and bounces to height 'h₂'. The rise in temperature of the ball is

$$\Delta \theta = \frac{g(h_1 - h_2)}{Jc}$$

JOULE'S LAW OR MAYER'S HYPOTHESIS: It states that there is no change in internal energy during the free expansion of gas.

SPECIFIC HEAT OF GASES:

- A gas will have two specific heats.
 a) Specific heat at constant volume (c_v)
 b) Specific heat at constant pressure(c_v)
- Specific heat depends only on the nature of material and temperature
- Water has largest specific heat among solids and liquids.
- Among solids, liquids, and gases specific heat is maximum for Hydrogen.
- Specific heat slightly increases with increase of temprature.
- In liquids specific heat is minimum for Mercury.
- The value of specific heat may lie between 0 and ∞.
- In isothermal process, the value of specific heat is ∞ but in adiabatic process its value is zero.
- Specific heat of water is maxium at 15°c and minimum at 37°c.
- Specific heat of all substances is zero at 0K.
- Substances with highest specific heat are bad conductors of heat and with low specific heat are good thermal and electrical conductors.
- The substance with large specific heat

warms up slowly and cools down slowly. C is greater than C., and.

$$\frac{C_p}{C_v} = \gamma (C_p, C_v \text{ are molar specific heats })$$

- C_p C_v=R (for 1 mole of gas) where R is universal gas constant R= 8.3 J/ mol-K
- C_p-C_v = r (for 1 gm of gas)
 Where r is specific gas constant

$$C_{p}-C_{v}=\frac{R}{J}$$
 (In heat units).

• Cv, Cp AND VALUES OF DIFFERENT GASES:

S.No.	Nature of gas	C_p	C,	$\gamma = \frac{c_p}{c_v}$
1.	Monoatomic	$\frac{5}{2}R$	$\frac{3}{2}R$	$\frac{5}{3} = 1.67$
2.	Diatomic	$\frac{7}{2}R$	$\frac{5}{2}R$	$\frac{7}{5} = 1.4$
3.	Tri (or)			
	Polyatomic	4R	3R	$\frac{4}{3} = 1.33$
	(or than and l

 γ value is always greater than one.lt depends upon the atomicity of a gas. It decreases with increase in atomicity.

$$C_p = \gamma \frac{R}{\gamma - 1}$$
 and $C_v = \frac{R}{\gamma - 1}$

• γ OF MIXTURE OF GASES: When n₁moles of a gas with specific heat at constant volume C_{v1} is mixed with n₂ moles of another gas of specific heat at constant volume C_{v2} then

$$(C_v)_{\text{mixture}} = \frac{n_1 c_{v1} + n_2 c_{v2}}{n_1 + n_2}$$
$$(C_p)_{\text{mixture}} = (C_v)_{\text{mixture}} + R$$

$$\gamma_{mixture} = \frac{C_{p_{(mixture)}}}{C_{v_{(mixture)}}}$$

Fraction of heat absorbed that is converted into

internal energy is
$$\frac{dU}{dQ} = \frac{C_v}{C_z} = \frac{1}{2}$$

Fraction of heat absorbed that is converted into

workdone=
$$\frac{dW}{dQ} = \frac{R}{C_p} = 1 - \frac{1}{\gamma}$$

ISOTHERMAL PROCESS:

 In this process, the pressure and volume of gas changes but temperature remains constant.

- The system is in thermal equilibrium with the surroundings.
- It is a slow process.
- The internal energy of the system remains constant i.e., du=0
- It obeys the Boyle's law i.e. PV=k
- The workdone during the isothermal expansion at constant temperature is

$$W = 2.303 RT \log_{10} \left(\frac{V_2}{V_1}\right)$$

=2.303 RTlog₁₀
$$\left(\frac{p_1}{p_2}\right)$$

• The isothermal elasticity

$$=-\frac{dp}{dv/v}=p$$

The -ve sign represents, as pressure increases volume decreases.

It takes place in a conducting vessel.

ADIABATIC PROCESS:

- The pressure, volume and temperature of a gas change but total heat remains constant i.e., dQ=0 (Q=constant).
- It is a quick process.
- The internal energy changes as temperature changes.
- The adiabatic process is represented by the equations.

 PV^{γ} = constant

 $TV^{\gamma-1}$ constant

 $P^{1-\gamma}T^{\gamma}$ =constant

The workdone by the system during the adiabatic expansion is

$$w = \frac{R}{\gamma - 1} (T_1 - T_2) = nC_v(T_1 - T_2)$$
$$= n\frac{c_p}{\gamma} (T_1 - T_2) = \frac{p_2 v_2 - p_1 v_1}{\gamma - 1}$$

- The adiabatic elasticity of gas is γ p.
- The slope of adiabatic curve is γ times greater than the isothermal curve.
- It takes place in a non conducting vessel.
- Adiabatic expansion causes cooling and contraction causes heating.
- If two samples of gases are compressed so that their pressures have the same increase, one sample isothermally and the other adia-batically, final volume is more in adiabatic change. If their pressures decrease by the same factor the final volume is more in Isothermal change.

ISOCHORIC PROCESS:

- It is a process in which the volume of the system remains constant.
 - i.e., ΔV =0 for such process ΔW =0
- In this process, the increase in internal energy is maximum where as the work done is zero.
- ISOBARIC PROCESS: It is a process in which the pressure of the system remains constant. i.e., <u>A</u> p=o

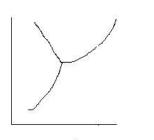
• THE EQUATION OF STATE FOR DIFFERENT TYPES OF PROCESSES:

SI. No.	Name of the process	Quantity remains constant	Quantity which becomes Zero	Result in I law
1. 2. 3. 4.	Isother- mal Isobaric Isochoric Adiabatic		dU None dW dQ	dQ=dW dQ=du+dW dQ=dU dU=-dW

Work done is maximum during isobaric process and minimum in adiabatic process.

• TRIPLE POINT:

- The tempertature and pressure where solid, liquid and vapour states are co-exist is called triple point.
- The triple point of water is 273.16K (0.00750°c) and pressure 613.10 Pa. (0.0459 cm of Hg)



- A graph drawn between the pressure and temperature representing the different states of matter is called the phase diagram.
- OA is the steam line and along this line water and steam are in equilibrium state.
- Above the line water exists and below steam exists.
- The curve has positive slope showing the boiling point increases with pressure.
- OB is called Hoar-frost line. Along this line ice and vapour coexist.
- OB has positive slope.
- OC is called ice line, along this line water

and ice are in equilibrium.

Above the ice line water exists. The curve has negative slope showing the melting point decreases with increase of the pressure.

SECOND LAW OF THERMODYNAMICS:

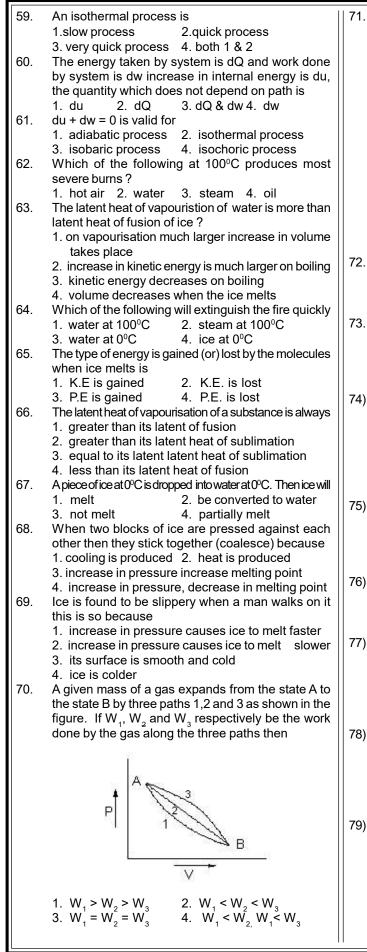
- Claussius statement: It is impossible for a self acting machine unaided by any external agency to transfer heat from a cold reservior to a hot reservior. In other words heat can't by itself flow from a colder to a hotter body.
- Kelvin-Planck Statement: It is impossible for an engine working during a cyclic process to extract heat from a reservoir and convert completely into work. In other words 100% conversion of heat into work is impossible.

CONCEPTUAL QUESTIONS

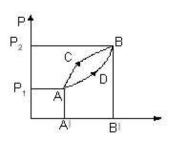
1. System A is in thermal equilibrium with B and B is separately in thermal equilibrium with C. Then A and C are in thermal equilibrium, from which law of thermo dynamics it follows? 1. Zeroth 2. First 3. Second 4. Third 2. Zeroth law of thermodynamics gives the concept of 2. Volume 1. Pressure 3. Temperature 4. Heat 3. Thermodynamic system returns to its original state, which of the following is NOT possible? 1. The work done is Zero 2. The work done is positive 3. The work done is negative 4. The work done is independent of the path followed 4. When we switch on the fan in a closed room. The temperature of the air molecules 1. Increase 2. Decrease 3. Remains Unchanged 4. May increase or decrease depending on the speed of rotation of the fan. 5. The work done by a thermodynamic system depends upon 1. The initial and final states of the system 2. Path along which work is done 3. Only on the initial state of the system 4. Only on the final state 6. Which type of molecular motion does not contribute towards internal energy? 1. Translational 2. Rotational 3. Vibrational 4. All the above 7. In which of the process the internal energy of the system remains constant? 1. Adiabatic 2. Isochoric 3. Isobaric 4. Isothermal 8. The internal energy of a perfect gas is independent of 1. Pressure 2. Temperature 3. Volume 4. Specific heat

9.	The internal energy of a perfect gas is	22.	Heat capacity of a substance is infinite. It means
	1. Complete kinetic		1. heat is given out
	2. Complete potential		2. heat is taken in
	3. Sum of potential and kinetic energy of the		3. no change in temperature whether heat is taken
	molecules 4. Difference of kinetic and potential energies of the		in (or) given out
	molecules		4. all of the above
10.	The internal energy of a real gas is independent of	23.	The temperature range in the definition of calorie is
	1. pressure 2. temperature		1. 14.5° c to 15.5° c 2. 15.5° c to 16.5° c
	3. volume 4. latent heat		3. 1° c to 2° c 4. 13.5° c to 14.5° c
11.	How does the internal energy change when the ice	24.	The heat capacity of material depends upon 1. the structure of a matter
	and wax melt at their normal melting points? 1. Increases for ice, decreases for wax		2. temperature of matter
	2. Decreases for ice and increases for wax		3. density of matter
	3. Decreases both for ice and wax		4. specific heat of matter
	4. Increases both for ice and wax	25.	We need mechanical equivalent of heat because
12.	Internal energy per mole of gas depends on		1. it converts work into heat
	1. viscosity 2. density		2. in C.G.S system, heat is not measured in the
13.	3. temperature 4. thermal conductivity In the free expansion of a gas, its internal energy		units of work
	1. remains constant 2. increases		3. in S.I., heat is measure in the units of work
Í	3. decreases		4. of some reason other than those mentioned above
1	4. sometimes increases, sometimes decreases	26.	The mechanical equivalent of heat J is
14.	The internal energy of an ideal gas depends upon		1. A constant 2. A physical quantity
1	 only its pressure only its volume only its temperature 		3. A conversion factor
1	4. its pressure and volume	27.	4. A dimensional quantity The bullet fired from a gun gets heated on striking a
15.	Heat is	21.	target because
	 kinetic energy of molecules 		1. it loses energy
	2. potential and kinetic energy of molecules		2. mechanical work is converted in to heat
	3. Energy in transit		3. of latent heat
16.	work done on the systemThe temperature determines the direction of net		4. specific heat
	change of	28.	The first law of thermodynamics is based on the law
1	1. gross kinetic energy		of conservation of
	2. intermolecular kinetic energy		1. energy 2. mass
1	3. gross potential energy		3. momentum 4. pressure
17.	 intermolecular potential energy The thermal motion means 	29.	First law of thermodynamics is a special case of
	1. motion due to heat engine		 law of conservation of energy Charle's law
1	2. disorderly motion of the body as a whole		3. law of conservation of mass
1	3. motion of the body that generates heat		4. Boyle's law
18.	4. random motion of molecules	30.	Which of the following states of matter have two
10.	Heat required to raise the temperature of one gram of water through 1°c is		specific heats ?
	1. 0.001 Kcal 2. 0.01 Kcal		1. solid 2. gas 3. liquid 4. vapour
1	3. 0.1 Kcal 4. 1.0 Kcal	31.	The specific heat of a gas in an isothermal process
19.	The direction of flow of heat between two bodies is		is
1	determined by		1. infinity 2. zero 3.negative 4.remains constant
1	1. Average kinetic energy2. total energy3. internal energy4. potential energy	32.	In defining the specific heat, temperature is
20.	Heat is absorbed by a body . But its temperature		represented in °F instead of °C. Then the value of
 	does not raised which of the following statement		specific heat will 1. decreases 2. increases
	explains the phenomena?		3. remain constant
1	1. only K.E. of vibration increases		4. be converted to heat capacity
Í	2. only P.E. of inter molecular force changes	33.	The ratio of the relative rise in pressure for adiabatic
	 no increase in internal energy takes place increase in K.E. is balanced by decrease in P.E. 		compression to that for isothermal compression is
21.	When heat is added to a system at constant		
	temperature, which of the following is possible.		1. γ 2. $\frac{1}{\gamma}$ 3. $1-\gamma$ 4. $\frac{1}{1-\gamma}$
	1. internal energy of system increases		$ \begin{array}{c} \ddots & \gamma \\ \gamma \\ \end{array} \begin{array}{c} 0 \\ 1 \\ \gamma \\ \end{array} \begin{array}{c} 0 \\ 1 \\ \gamma \\ \end{array} \begin{array}{c} 1 \\ -\gamma \\ \end{array} \right $
	2. work is done by system	34.	Ratio of isothermal elasticity of gas to the adiabatic
	neither internal energy increases nor work done by system		elasticity is
	4. internal energy increases and also work done by		1 1
Í	system		1. γ 2. $\frac{1}{\gamma}$ 3. $1-\gamma$ 4. $\frac{1}{1-\gamma}$
			$\gamma \gamma \gamma - \gamma \gamma$
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35.	Why the specific heat at a constant pressure is more	47.	For the Boyle's law to hold good, the necessary
	than that at constant volume 1. there is greater inter molecular attraction at		condition is 1. Isobaric 2. Isothermal
	constant pressure		3. Isochoric 4. Adiabatic
	2. at constant pressure molecular oscillation are	48.	Two samples of gas A and B, initially at same
	more violent		temperature and pressure, are compressed to half
	3. rational work need to be done for allowing		their initial volume, A isothermally and B adiabatically.
	expansion of gas at constant pressure		The final pressure in
	4. due to more reasons other than those mentioned		1. A and B will be same 2. A will be more than in B
	in the above		3. A will be less than in B
36.	The ratio $[C_p / C_v]$ of the specific heats at a constant		4.A will be double that in B
	pressure and at a constant volume of any perfect gas	49.	In which of the following processes all three
	1. can't be greater than 5/4		thermodynamic variables, that is pressure volume
	 can't be greater than 3/2 can't be greater than 5/3 		and temperature can change
	4. can have any value		1. Isobaric 2. Isothermal
37.	Which of the following formulae is wrong ?	50	3. Isochoric 4. Adiabatic
07.		50.	During adiabatic expansion the increase in volume associated with
1	1. $C_v = \frac{R}{\gamma - 1}$ 2. $\frac{C_p}{C_v} = \gamma$		1. increase in pressure and temperature
1	1. $C_v = \gamma - 1$ 2. $\overline{C_v} = \gamma$		2. decrease in pressure and temperature
1	· - v		3. increase in pressure and decrease in temperature
1	$\gamma.R$		4.Decrese in pressure and increase in
1	3. $C_p = \frac{\gamma \cdot R}{\gamma - 1}$ 4. $C_p - C_v = 2R$		temperature
20	/	51.	A gas is being compressed adiabatically. The specific
38.	Which of the following laws of thermodynamics leads to the interference that it is difficult to convert		heat of the gas during compression is 1. zero 2. infinite
	whole of heat into work		3. finite but non zero 4. undefined
	1. zeroth 2. second 3. first 4. third	52.	The process in which no heat enters or leaves the
39.	The process, in which the internal energy of the		system is termed as
	system remains constant ?		1. Isochoric 2. Isobaric
	1. adiabatic 2. isobaric		3. Isothermal 4. adiabatic
	3. isochoric 4. isothermal		$\begin{bmatrix} pv \end{bmatrix}$
40.	The temperature at which adiabatic change is	53.	The gas law $\left \frac{pv}{T} \right $ = constant is true for
	equivalent to the isothermal change?		1. isothermal change only
	 zero degree celsius 2. zero kelvin critical temperature 		2. adiabatic change only
	4. above critical temperature		3. Both isothermal & adiabatic
41.	Two identical sample of gases are allowed to expand		4. neither isothermal nor adiabatic
	(i) isothermally (ii) adiabatically work done is	54.	During adiabatic compression of a gas, its
1	 more in the isothermal process 		temperature
1	2. more in the adiabatic process		1. falls2. rises3. remains constant4. becomes zero
1	3. equivalent in both process	55.	On compressing a gas suddenly, its temperature
	4. equal in all process		1. increases 2. decreases
42.	In the isothermal expansion of a gas		3. remains constant 4. all the above
	 the work done by gas is zero heat is taken from the gas 	56.	The work done on the system in an adiabatic
1	3. heat is neither given nor thaken from gas		compression depends on
1	4. internal energy stored in the gas is constant		1. the increase in internal energy of the system
43.	The conversion of water into ice is an		 the decrease in internal energy the change in volume of the system
1	1. isothermal process 2. isochoric process		4. all the above
	3. isobaric process 4. entropy process	57.	The internal energy of compressed real gas, as
44.	The internal energy of an isolated system		compared to that of the normal gas at the same
1	1. remains constant 2. keeps on changing		temperature is
15	3. zero 4. depending on gas		1. less 2. more
45.	When an ideal gas is expands isothermally, its internal energy will		 sometimes less, sometimes more maximum
1	1. increases 2 decreases	58.	The ratio of slopes of adiabatic and isothermal curves is
1	3. remains constant 4. becomes zero		1
46.	For an isothermal process		1. γ 2. $\frac{1}{\gamma}$ 3. γ^2 4. γ^3
1	1. $dQ = dw$ 2. $dQ = du$, - γ · γ · γ
1	3. $dw = du$ 4. $dQ = du + dw$		



71. 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



- 1. P₁ACBP₂P₁ 2. ACBB'A'A 3. ACBDA 4. ADBB'A'A
- In isothermal expansion the pressure is determined by
- 1. temperature only 2. compressibility only
- 3. both temperature and compressibility
- 4. latent heat
- 73. A given system undergoes a change in which the work done by the system equals the decrease in its internal energy. The system must have undergone an
 - 1. Isothermal change 2. Adiabatic change
 - 3. Isobaric change4. Isochoric change
- 74) A closed vessel contains some gas at a given temperature and pressure. If the vessel is given a very high velocity, the temperature of the gas is 1. increases 2. decreases
 - 3. may increase or decrease depending upon the nature of the gas
 - 4. does not change
- 75) Cooking is difficult on mountains because1) water boils at low temperature
 - 2) water boils at high temperature
 - 2) water boils at high temperatul3) water does not boil
 - 4) it is cool there
- 76) Paraffin wax contracts on solidification. So its melting point with pressure
 - 1) increases 2) decreases
 - 3) remains same 4) we can't say
- 77) In a pressure cooker cooking is done quickly because1) the cooker doesnot absorb any heat
 - 2) it has a safety valve
 - 3) boiling point of water rises due to increased pressure
 - 4) it is a prestige to cook in a cooker
- 78) A large block of ice is placed on a table when the surroundings are at 0°C
 - 1) ice melts at the sides
 - 2) ice melts at the top
 - 3) ice melts at the bottom
 - 4) ice does not melt at all
- 79) What energy transformation takes place when ice is converted into water
 - 1) heat energy to kinetic energy
 - 2) kinetic energy to heat energy
 - 3) heat energy to latent heat
 - 4) heat energy to potential energy

			1
80)	Heat is supplied to a diatomic gas at constant	6.	The amount of heat required to convert 5g of ice at 0% into water at 4% is
	pressure. The ratio of $\Delta Q : \Delta U : \Delta W$ is		0°C into water at 40°C is
	1) 5 : 3 : 2 2) 5 : 2 : 3 3) 7 : 5 : 2 4) 7 : 2 : 5	_	1. 400 cal 2. 200 cal 3. 600 cal 4.1000 cal
81)	Unit mass of liquid of volumeV ₁ completely turns into a	7.	The thermal capacity of 100g lead shot is (specific
	gas of volume V_2 at constant atmospheric pressure P		heat of lead is 0.03 cal/g/°C)
	and temperature T. The latent heat of vaporization is		1. 3 cal/°C 2. 30 cal/°C 3. 300 cal/°C 4. 0.3 cal/°C
	"L". Then the change in internal energy of the gas is $(1) = 0$ (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	8.	3. 300 cal/ ⁰ C 4. 0.3 cal/ ⁰ C Specific heat of aluminium is 0.25 cal/g/ ⁰ c. The wa-
	1) L 2) L+P($V_2 - V_1$) 3) L - P($V_2 - V_1$) 4) Zero	0.	ter equivalent of an aluminium vessel of mass one
82)	In a isobaric (constant pressure) process. the correct ratio is		kilogram is
	1) $\Delta Q : \Delta U = 1 : 1$ 2) $\Delta Q : \Delta U = 1 : \gamma - 1$		1. 40 cal/°C 2. 250 cal/°C
	3) $\Delta Q : \Delta U = \gamma - 1: 1$ 4) $\Delta Q : \Delta U = \gamma : 1$		3. $25 \text{ cal/}^{\circ}\text{C}$ 4. $2.5 \text{ cal/}^{\circ}\text{C}$
83)	In an isobaric process, the correct ratio is	9.	Two liquids A and B of equal volumes have their spe-
	1) $\Delta Q : \Delta W = 1 : 1$ 2) $\Delta Q : \Delta W = \gamma : \gamma - 1$		cific heats in the ratio 2:3. If they have same ther-
	3) $\Delta Q : \Delta W = \gamma - 1; \gamma$ 4) $\Delta Q : \Delta W = \gamma : 1$		mal capacity, then the ratio of their densities is
	KEY		1. 1 : 1 2. 2 : 3 3. 3 : 2 4. 5 : 6
	01) 1 02) 3 03) 4 04) 1 05) 2	10.	Two spheres have their radii in the ratio 1 : 2, the
	06) 2 07) 4 08) 3 09) 1 10) 2,3		densities of their materials are in the ratio 2 : 3 and
	11) 1 12) 3 13) 1 14) 3 15) 3		their specific heats are in the ratio 3:4, then the ra-
	16) 2 17) 4 18) 1 19) 1 20) 2		tio of their thermal capacities is
	21) 2 22) 3 23) 1 24) 4 25) 2		1. 3:4 2. 1:16 3. 2:3 4. 1:1
	26) 3 27) 2 28) 1 29) 1 30) 2	11.	The densities of two substances are in the ratio 3 : 4
	31) 1 32) 1 33) 1 34) 2 35) 3		and their specific heats are in the ratio 2 : 3. The
	36) 3 37) 4 38) 3 39) 4 40) 2		ratio of thermal capacities per unit volume of these substances is
	41) 1 42) 4 43) 3 44) 1 45) 3		1. 2:3 2. 3:4 3.1:1 4. 1:2
	46) 1 47) 2 48) 3 49) 4 50) 2	12.	1kg of water and 1kg of steel are heated through 1K.
	51) 1 52) 4 53) 3 54) 2 55) 1	'2.	The change in their internal energies are (Specific
	56) 1 57) 1 58) 1 59) 1 60) 1 60) 4 60) 4 60) 4 60) 1		heat of steel = 460 J/kg/K; Specific heat of water =
	61) 1 62) 3 63) 1 64) 1 65) 3 60) 4 67) 9 60) 4 70) 9		4200J/kg./K)
	66) 1 67) 3 68) 4 69) 1 70) 3 71) 2 72) 2 73) 2 74) 4 75) 1		1. 460 J, 4200 J 2. 4200 J, 460 J
	71) 3 72) 2 73) 2 74) 4 75) 1 76) 1 77) 3 79) 3 70) 4 90) 3		3. 1000 J, 4200 J 4. 460 J, 1000 J
	76) 1 77) 3 78) 3 79) 4 80) 3 81) 3 82) 4 83) 2	13.	Two liquids A and B are at 30°C and 20°C respec-
	01/0 02/4 00/2		tively. When they are mixed in equal masses the
	LEVEL-1		temperture of the mixture is found to be 26°C. The
			ratio of specific heat is
1.	The ratio of densities of two substances is 2:3 and		1. 4:3 2. 3:4 3. 2:3 4. 3:2
	that of specific heats is 1 : 2. The ratio of thermal	14.	M kg of ice at 0°C is mixed with M kg of water at
	capacities per unit volume is		10ºc. The final temperature is 1. 8ºC 2. 6ºC 3. 4ºC 4. 0ºC
	1.1:2 2.2:1 3.1:3 4.3:1	15.	1. 8°C 2. 6°C 3. 4°C 4. 0°C 5 g of steam at 100°C is mixed with 5g of the ice at
2.	The density of a substance is 400 kgm ⁻³ and that of	13.	0°C. The final temperature of the mixture is
	another substance is 600 kgm ⁻³ . The heat capacity		1. 100°C 2. 95°C 3. 90°C 4. 80°C
	of 40 c.c of first substance is equal to that of 30 c.c	16.	A beaker contains 200g of water. The heat capacity
	of second substance. The ratio of their specific		of the beaker is equal to that of 20g water. The
	heats is		initial temperature of water in the beaker is 20°C. If
	1.1:6 2.6:1 3.9:8 4.8:9		440g of hot water at 92°C is poured in, the final tem-
3.	A body absorbs 1000 calories of heat when it is		perature (neglecting radiation loss) will be nearly
	heated from 20°C to 70°C. The water equivalent of the body is		1. 58°C 2. 68°C 3. 73°C 4. 78°C
	1. 10 g 2. 15 g 3. 20 g 4. 25 g	17.	If 10g of the ice at 0°C is mixed with 10g of water at
4.	Two spheres of copper of diameters 10cm and 20		10°C, then the final temperature of the mixture will
	cm will have thermal capacities in the ratio		
		10	1. 5°C 2. 0°C 3. 100 K 4. 40°C
	1. $\frac{1}{8}$ 2. $\frac{1}{2}$ 3. $\frac{1}{4}$ 4. $\frac{1}{6}$	18.	One gram of water at 80° C is mixed with one gram of ice at 0° C. The resultant temperature is
1	8 2 4 6		1. 100°C 2. 0°C 3. 80°C 4. 40°C
5.	When 5 moles of gas is heated from 100°C to 120°C	19.	If 20g of water at 60° C is mixed with 60g of water at
	at constant volume, the change in internal energy is		20° C. The resultant temperature is
	200 J. The specific heat capacity of the gas is		1. 60°C 2. 20°C 3. 30°C 4. 0°C
	1. $5 \text{ J mol}^{-1}\text{K}^{-1}$ 2. $4 \text{ J mol}^{-1}\text{K}^{-1}$		
	3. $2 \text{ J mol}^{-1}\text{K}^{-1}$ 4. $1 \text{ J mol}^{-1}\text{K}^{-1}$		

20. 21.	The heat energy required to vapourise 5kg of water at 373 K is 1. 2700 K.cal 2. 1000 K.cal 3. 27 K.cal 4. 270 K.cal The amount of heat required to convert 1g of ice at - 10°C into steam at 100°C is (specific heat of ice = 0.5 col/a (specific heat of i	32.	In a thermodynamic process with 2 moles of gas 30 J of heat is released and 22 J of work is done on the gas. Given that initial internal energy of the sample was 20 J. The final internal energy is 1.72 J 2.32 J 3.28 J 4.12 J
	$0.5 \text{ cal/g}/^{\circ}\text{C}$, latent heat of ice = 80 cal/g , latent heat of steam = 540 cal/g)	33.	A gas for which γ is $\frac{4}{3}$ is heated at constant
22.	1. 3045 J 2. 735 J 3. 1000 J 4. 4200 J A liquid of specific heat 0.3 cal/g/°C at 90°C is mixed with another liquid of specific heat 0.5 cal/g°C at 15°C. If the resultant temperature of mix- ture is 60°C then the ratio of their mases is		pressure. The percentage of heat supplied used for external work is 1. 25% 2. 75% 3. 60% 4. 40%
23.	1. $5:2$ 2. $2:5$ 3. $1:1$ 4. $3:4$ Two liquids A and B are at temperatures of 75° C and 150° C respectively. Their masses are in the ratio of 2 : 3 and specific heats are in the ratio	34.	The molar specific heat of hydrogen at constant volume is 5 cal. Heat required to raise the temp. of 1 gm H_2 gas by 1°C at constant volume is 1. 2 cal 2. 2.2 cal 3. 2.5 cal 4. 4 cal
	3:4. The resultant temperature of the mixture, when the above liquids, are mixed (Neglect the water equivalent of container) is $1.125^{\circ}C$ 2. $100^{\circ}C$ 3. $50^{\circ}C$ 4. $150^{\circ}C$	35.	The molar specific heat of helium at constant volume is 3 cal. Heat energy required to raise the temperature of 1gm helium gas by 1°C at constant pressure is
24.	It takes 15 minutes for an electrical kettle to heat a certain quantity of water in grams from 0°c to boiling point 100°C. If takes 81 minutes to boil water into steam, the latent heat of steam is 1. 80 cal/g 2.540 cal/g 3. 100 cal/g 4.336 cal/g	36.	1. 1.2 Cal 2. 1.25 Cal 3. 3 Cal 4. 4 Cal A quantity of heat 'Q' is supplied to a monoa tanic ideal gas which expands at constant pressure. The fraction of heat that goes into work done by the gas is
25.	A copper weight and iron weight of the same mass were dropped from the same height on the ground. Which of the weights had the higher temperature after the impact? (specific heat of Cu = 0.1 Kcal/kg/		1. $\frac{2}{5}$ 2. $\frac{5}{2}$ 3. $\frac{4}{5}$ 4. 1
	 ^oC ;specific heat of iron = 0.11 Kcal/kg/^oC) 1. Iron weight 2. Copper weight 3. Both iron and copper 4. Cannot be answered FIRST LAW OF THERMODYNAMICS 	37.	3 moles of gas requires 60 cal of heat for 5° C rise of temperature at constant volume. Then heat required for 5 moles of same gas under constant pressure for 10° C raise of temperature is
	(dQ = dU + dW)		1. 200 Cal 2. 400 Cal 3. 100 Cal 4. 300 Cal
26.	A gas absorbs 80 cal of heat. If the work done by the gas is 150J the change in internal engery is 1. 150 J 2. 80 J 3. 186 J 4. 156 J	38.	For a gas the ratio of the two specific heats is $\frac{5}{3}$. If
27.	Heat energy of 2100 J is given to a gas at a con- stant pressure of 10^5 Pa. The increase in internal energy if the change in volume is $5 \times 10^{-3} \text{m}^3$		$\label{eq:R} \begin{array}{l} R = 2 \mbox{ cal } / \mbox{ mol/K} \mbox{ then the values of } C_{p} \mbox{ and } \mathbb{C} v \mbox{ are } \\ 1. 5 \ ,3 \mbox{ cal } / \mbox{ mol } / \ ^0 \mbox{K} \mbox{ 2. 3 } \ ,4 \mbox{ cal } / \mbox{ mol } / \ ^0 \mbox{K} \\ 3. 4 \ ,3 \mbox{ cal } / \mbox{ mol } / \ ^0 \mbox{K} \mbox{ 4. 3.5 } \ , \ 7 \ \mbox{ cal } / \mbox{ mol } / \ ^0 \mbox{K} \end{array}$
28.	1. 1500 J 2. 1400 J 3. 1600 J 4. 800 J When 20 cal of heat is supplied to a system, the in-	39.	If for a gas $\frac{R}{C}$ = 0.67, this gas is made up of mol-
	crease in internal energy is 50 J . If the external work done is 35 J, the mechanical equivalent of heat is 1. 4.25 J/cal 2. 1.26 J/cal		ecules which are 1. Diatomic 2. Monoatomic 3. Poly atomic
29.	 3. 4.92 J/cal 4. 2.1 J/cal 100 J work is done on a gas to reduce its volume under adiabatic conditions. The change in internal energy of the gas is 	40.	4. Mixture of diatoamic & Polyatomic One mole of an ideal gas requires 207 J heat to raise the temperature by 10K when heated at con-
30.	1. +100 J 2100 J 3200 J 4. +200 J In a thermodynamic process the pressure of fixed mass of gas is changed. In this process gas re- leases 20 J heat and 8 J work is done on the gas. If initial internal energy of the gas is 30 J, then final internal energy is	41.	stant pressure. If the same gas is heated at con- stant volume to raise the temperature by the same 10K, the heat required is (R=8.3 J/mol/K) 1. 187J 2. 29J 3. 215.3J 4. 124 J If the ratio of sp.heat of a gas at constant pressure to that at constant volume is γ , the change in in-
31.	1. 2 J 2. 42 J 3. 18 J 4. 58 J 70 cal heat is required to raise the temperature of 2 moles of an ideal gas at constant pressure from 25°C to 30°C, then the amount of heat required to raise the temperature of same gas through same raise of temp. at constant volume is 1. 50 cal 2. 70 cal 3. 90 cal 4. 60 cal		ternal energy of gas, when the volume changes from V to 2V at constant pressure P is 1. $\frac{R}{\gamma - 1}$ 2. PV 3. PV / ($\gamma - 1$) 4. $\gamma PV / (\gamma - 1)$

52. In an adiabatic expansion, the temperature of 5 For a certain gas, the ratio of specific heats is $\frac{3}{2}$ moles of gas $\gamma = 1.5$ falls from 87°C to 27°C, then 42. the work done is The value of C_{D} for the gas is 1. 2400 Cal 2. 4980 Cal 1. R 2. 2R 3. 3R 4. 5R 3. 1200 Cal 4. 3000 Cal 43. For hydrogen gas C_p-C_y = a and for Oxygen gas C_p-C = b, Then the relation between 'a' and 'b'. is If a monoatomic gas is suddenly compressed to 53. 1. a = 16b 2. b = 16a 3. a = 4b 4. a = b 44. One gram of water on evaporation at atmospheric of its volume adiabatically, then the pressure of pressure forms 1671 cm³ of steam. Heat of the gas is vaporisation at this pressure is 540 cal gm⁻¹. The 1. 8 times the initial 2. 16 times the initial increase in internal energy is 3. 32 times the initial 4. 128 times the initial 1.250cal 2.500cal 3.1000cal 4.1500cal 54. An ideal monoatomic gas is taken round the cycle 45. A cylinder of fixed capacity 67.2 litres contains ABCDA as shown in the diagram. The work done helium gas at S.T.P. The amount of heat required during the cycle is to raise the temperature of the gas by 15°C is (R =8.31 J/mol/K) 1. 520 J 2. 560.9 J 3. 620 J 4. 621.2 J B(2P V) C(2P,2V) ADIABATIC AND ISOTHERMAL P For a gas $\gamma = \frac{5}{3}$. 800c.c. of this gas is suddenly 46. D(P,2V) A(P,V) compressed to 100c.c. If the initial pressure is P, then the final pressure will be 1. $\frac{P}{32}$ 2. $\frac{24}{5}$ 3. 8P 4. 32P 1. PV 2. 2PV 3. 3PV 4. 4PV 55. Agas is compressed isothermally and adiabatically. The corresponding change in volume are found to be 47. In an adiabatic change, the pressure P and tem-51 c.c and 34 c.c. The value of γ for the gas is perature T of a mnoatomic gas are related as P α T^c 1. 1.67 2. 1.4 3. 1.33 4.1.5 where C equals. 56. Waterfall is falling from 160m height. Assuming 1. $\frac{5}{3}$ 2. $\frac{2}{3}$ 3. $\frac{3}{5}$ 4. $\frac{5}{2}$ that half the K.E. of falling water gets converted into heat, the rise in temperature of water is appoximately. A polyatomic gas ($\gamma = \frac{4}{3}$) is compressed to $\frac{1}{8}$ of 1. 0.1°C 2. 0.2°C 3. 0.3°C 4. 0.4°C 48. 57. How much will the temperature of 100g of water be raised by doing 4200 J of work is stirring the water? its volume adiabatically. If it's initial pressure is P, 1. 0.01°C 2. 0.1°C 3. 1°C 4. 10°C the new pressure will be From what minimum height a block of ice has to be 58. 4. 2P 1.8P 2. 16P 3. 6P dropped in order that it may melt completely on hit-49. During an adiabatic change the density becomes ting the ground $\frac{1}{16}$ th of initial value, then $\frac{P_1}{P_2} = (\gamma = 1.5)$ 1. mgh 2. $\frac{mgh}{J}$ 3. $\frac{JL}{g}$ 4. $\frac{J}{L\sigma}$ 1.16 2.4 3. 32 4.64 59. A bullet travelling at 100 ms⁻¹ suddenly hits a con-50. A fixed amount of dry air at temperature of 27°C is crete wall. If its K.E. is converted completely into heat , the raise in temperature is compressed to $\frac{1}{\Omega}$ of original volume. Its final tem- $(s = 100 J kg^{-1} K^{-1})$ perature is ($\gamma_1 = 1.5$) 1. 20K 2. 40K 3. 50K 4. 60K 1. 627°C 2. 600°C 3. 158°C 4. 527°C 60. To metal balls of same material having masses 50gm and 100gm collides with a target with same velocity. The pressure and density of a diatomic gas $\left(\gamma = \frac{\gamma}{5} \right)$ Then the ratio of their rise in temperature is 51. 1. 1:2 2. 4:1 3. 2:1 4. 1:1 61. The difference between the two specific heat of a gas C₂- C₂ =2 cal. If R = 8.31 \times 10⁷ erg/k/gm,the where $\frac{\rho_1}{\rho_2} = 32$ then $\frac{p_2}{p_1}$ should be value of mechanical equivalent of heat is 1. 4.15J/cal 2. 42J/cal 2. 128 3. 146 4. 124 1. 126 3. 420J/cal 4. 4200J/cal

62.					ning that the
					perfectly in- ater is dissi-
	pated a	s heat. Fin	d the rise	in tempara	ature of the
	water. (g = 10 m/s C ⁻¹ , J = 42	² , Specific	c heat of w	ater = 1000
	1. 0.1°	C , 3 – 42 C 2. 0.5 ⁰	00 J / R . Ca	ⁿ) °C 4.0.	25⁰C
63.	From wi	hich height	a block of	ice must b	e dropped in
					that all the = 10 ms^{-2} ,
		al gm ⁻¹ an			
		Km 2.100			
64.					essure. The ncrease the
		energy is	at energy is	5 0300 10 1	
05	1. 3/5	2. 3/7			
65.		•			to increase
		nal energy		99 10 4004	
66	1.0%	2. 14			100 %
66.		ertain gas			fic heats is
	1. R	2. 2R	3. ŠI	R 4.5	
67.	Four stu	udents fou	nd set of	C _p and C	[in cal/deg
	correct	is given be			owing set is
	1. C _v =	4, $C_p = 2$ 3, $C_p = 4$	2. C	= 4 , C _p = p = 5 , C _v =	= 3
	3. C _v =	3 , C _p = 4	4. C KEY	_p = 5 , C _v =	= 3
	01)3	02)3	03)3	04)1	05)3
	06)3	07)1	08)2	09)3	10)2
	11)4 16)2	12)2 17)2	13)4 18)2	14)4 19)3	15)1 20)1
	21)1	22)1	23)1	24)2	25)2
	26)3	27)3	28)1	29)1	30)3
	31)1 36)1	32)4 37)4	33)1 38)1	34)3 39)2	35)4 40)4
	41)3	42)3	43)4	44)2	45)2
	46)4	47)4	48)2	49)4	50)1
	51)2 56)2	52)3 57)4	53)3 58)3	54)1 59)3	55)4 60)4
	61)1	62)2	63)3	64)3	65)1
	66)3	67)4			
			HINTS		
1.	$\frac{H_1}{H} = \frac{a}{h}$				
	H_2 a	$l_2 c_2$			
	$\frac{c_1}{v_2} = \frac{v_2}{v_2}$	d.			
2.	$\frac{c_1}{c_2} = \frac{v_2}{v_1}$				
3.		-			
3.		mc (70-20)			
	$w = \frac{100}{100}$	$\frac{00}{0} = 20 \text{ gr}$	m		
	5	0	-		
	m.	r_{1}^{3} 1			
4.	<u> </u>	$\frac{1}{3} = \frac{1}{2}$			
	m_{2}	r. 8			
	$\frac{m_1}{m_2} = \frac{1}{2}$	r_2^{5} 8			

$$\frac{H_1}{H_2} = \frac{m_1}{m_2} = \frac{1}{8}$$
5. $dQ = du = mc_{\Delta}t \text{ since } dQ = 200J$
 $\Delta t = 20K \text{ and } M = 5 \text{ mole}, \text{ hence}$
 $c = 2J \text{ mol}^{+K^{-1}}$
6. $Q = ms \Delta \theta$
7. $Q = ms$
8. water equivalent = ms
9. $m_1 s_1 = m_2 s_2$, $v_1 p_1 s_1 = v_2 p_2 s_2$
 $\frac{\rho_1}{\rho_2} = \frac{v_2 s_2}{v_1 s_1} = \frac{s_2}{s_1}$
10. $\frac{C_1}{C_2} = \frac{r_1^{-3}}{r_2^{-3}} \times \frac{d_1}{d_2} \times \frac{s_1}{s_2}$
11. $\frac{T_1}{T_2} = \frac{\rho_1 s_1}{\rho_2 s_2}$
12. $Q = ms \Delta t$
16. Applying principle of calorimitry
(200 + 20) × 1 × $\theta = 68^{\circ}c$
19. $20(60 \cdot t) = 60 (t \cdot 20)$
22. $ms\Delta t_1 = ms\Delta t_2$
26. $du = dQ - dw$
27. $du = dQ - pdv$
28. $dQ = du + dw$
29. $du = -dw$
30. $U_2 - U_1 = dQ - dW$
32. $dQ = du + dw = u_2 - u_1 + dw$
33. $\frac{dw}{dQ} \times 100 = \left(1 - \frac{1}{\gamma}\right) \times 100$
34. $dQ = nc_y dt$
35. $c_p = c_v + R$
 $dQ = nc_p dt$
36. $\frac{dw}{dQ} = 1 - \frac{1}{\gamma}$
37. $du = nc_v dt$
38. $c_v = \frac{R}{\gamma - 1}$
 $c_p - c_v = R$

39.
$$\frac{R}{w} = \frac{c_p - c_v}{c_v} = \gamma - 1 = 0.67$$

 $\gamma = 01.67$
40.
$$c_p = \frac{5}{2}R$$

$$c_v = \frac{3}{2}R$$

$$\frac{Q_v}{Q_p} = \frac{3}{5} \text{ here } Q_p = 270 \text{ J} , Q_v = 124 \text{ J}$$

41.
$$dv = nc_v dt \text{ also } \frac{c_p}{c_v} = \gamma$$

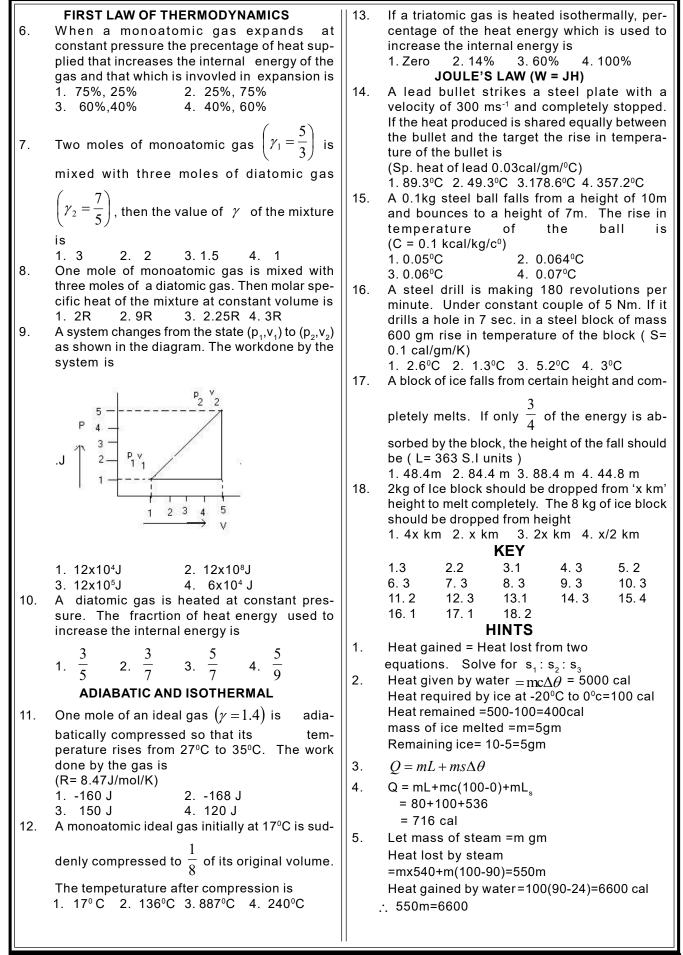
$$\frac{c_p - c_v}{c_v} = \gamma - 1 \text{ i.e. } c_v = \frac{R}{\gamma - 1}$$
42.
$$\frac{c_p}{c_v} = \frac{3}{2} = \frac{c_v + R}{c_v} = \frac{3}{2}$$
hence $c_v = 2R$ and $c_p = 3R$
43. Both are diatomic gases also
 $c_p - c_v = R$ for all gases
44.
$$dQ = du + dw$$
mL = $du + pdv$
du = mL - pdv
45.
$$Q = nc_v dT$$

$$= 3 \times \frac{3}{2} \times 8 .31 \times 15 = 560.9 \text{ J}$$

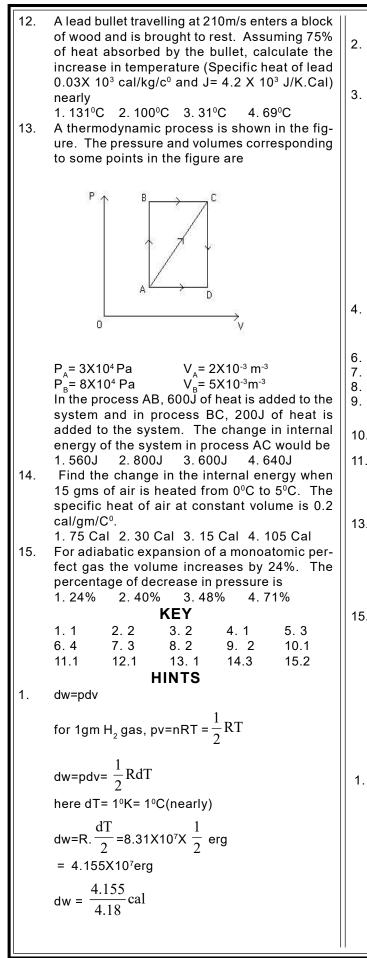
46. $p_1 v_1^{\gamma} = p_2 v_2^{\gamma}$
47. $T^{\gamma} p^{1-\gamma} = const$
48. $p_1 v_1^{\gamma} = p_2 v_2^{\gamma}$
49.
$$\frac{p_1}{p_2} = \left(\frac{d_1}{d_2}\right)^{\gamma}$$

50. $TV^{\gamma - 1} = constant$
51. $pv^{\gamma} = k \quad \frac{p}{\rho^{\gamma}} = k$
 $\frac{P_2}{p_1} = \frac{p_2^{\gamma}}{p_1}$
52.
$$w = \frac{nR}{\gamma - 1} (T_1 - T_2)$$
where R = 2 cal mol^{-1} k^{-1}

53.
$$pv^{\gamma} = k$$
 where $\gamma = \frac{5}{3}$
54. work done = Area under rect angle
55. $\left(\frac{dv}{v}\right)$ Isothermal = $\gamma\left(\frac{dv}{v}\right)$ Adiabatic
56. $\frac{1}{2}$ mgh = msdt
57. w = JH w = Jmc $\Delta \theta$
4200 = 4.2 x 100 x 1 x $\Delta \theta$
 $\Delta \theta = 10^{\circ}$ C
58. w = JH mgh = Jml
59. $\frac{1}{2}$ mv² = ms $\Delta \theta$
60. $\frac{1}{2}$ mv² = ms $\Delta \theta$
61. $c_{p} - c_{v} = \frac{R}{J} = \frac{R}{c_{p} - c_{v}}$
62. mgh = Jms θ
63. mgh = JmL
LEVEL - II
CALORIMETRY
1. Three liquids A,B and C of masses 400gm, 600
gm and 800 gm are at 30°c, 40°c and 50°c re-
spectively. When A and B are mixed resultant
temperature is 36°C when B and C are mixed
resultant temperature is 44°C Then ratio of their
specific heats are
1. 2:1:1 2. 3:2:1 3. 2:2:1 4. 1:4:9
2. 10gm of Ice at -20°C is added to 10g of water
at 50°C. The amount of ice and water that are
present at equilibrium respectively
1. 0,20g 2. 5g, 15g
3. 5g, 10g 4. 10g, 10g
3. A boy eats an ice cube of mass 100gm at 0°C. If the
body temperature of the boy is 37°C the number of
joules of energy lostby the boy is (L_{ice} = 33000J/kg
for water Sp. heat =4200 J-kg^{+(x)})
1. 49140 J 2. 39140 J
3. 49000 J 4. 29140 J
3. 716 cal 4. 450 cal
5. The temperature of 100 gm of water is to be raised
from 24°C to 90°C by adding steam to it. The
mass of the steam required for this purpose is
1. 10gm 2. 12 gm 3. 14gm 4. 16gm



$$\begin{split} \mathbf{m} &= \frac{6600}{550} = 12g \\ \mathbf{m} &= \frac{6}{550} = 12g \\ \mathbf{m} &= \frac{1}{550} = \frac{1}{550} = \frac{1}{550} = \frac{1}{550} = \frac{1}{550} \\ \mathbf{m} &= \frac{1}{550} = \frac{1}{550} = \frac{1}{550} = \frac{1}{550} = \frac{1}{550} = \frac{1}{550} \\ \mathbf{m} &= \frac{1}{550} = \frac{1}{50} = \frac{1}{550} = \frac{1}{50} = \frac{1}{550} =$$



$$\gamma = \frac{n_1 \gamma_1 + n_2 \gamma_2}{n_1 + n_2}$$

$$C_A(16-12) = C_B(19-16) \Rightarrow \frac{C_A}{C_B} = \frac{3}{4}$$

$$C_B(23-19) = C_C(28-23) \Rightarrow \frac{C_B}{C_C} = \frac{5}{4}$$

$$\Rightarrow \frac{C_A}{C_C} = \frac{15}{16}$$
When A andC are mixed
$$C_A(\theta - 12) = C_C(28 - \theta)$$
Where θ is common temperature
 $\theta = 20.2^{\circ}C$

$$m_s L_s = (m_{icc} L_{icc} + m_{icc}s\Delta t)$$
Where m_s mass of steam condensed to rise temperature of ice to 100°c water.
Heat lost = Heat gained
$$Heat lost = Heat gained
Heat lost = Heat gained
Implement = $\frac{gh}{100} = \frac{gh}{12} = \frac{980 \times 10^5}{133} = \frac{1}{33}$
State AB is a isochoric process there work done=P_B
(V_p, V_n)=240J
dQ = 600+200=800J
Now dQ = dU+dW
dU = 800-240=560J
Implement P_0 = $\frac{\Delta p}{p} \times 100 = -\gamma \frac{\Delta v}{v} \times 100$
for monoatomic gas $\gamma = \frac{5}{3}$

$$\frac{\Delta p}{p} \times 100 = \frac{5}{4} \times 24 = 40$$

$$\frac{MEW MODEL QUESTION & ANSWERS}{Match the following:
List - 1
List - 11
a) Isothermal process
f) no heat exchange
c) Isobaric process
f) no heat exchange
c) Isobaric process
f) no heat excha$$$$

2.	Match the following :
<u> </u> ∠.	List - I List - II
	a) Latent heat capacity e) $ML^2T^{-2}K^{-1}$
	b) Specific heat capacity f) $M^{0}L^{2}T^{-2}K^{-1}$
	c) Stefans constant g) L^2T^{-2}
	d) Boltzmaum constant h) $MT^{-3}K^{-4}$
	1) a - g, b - f, c - h, d - e 2) a - f, b - g, c - e, d - h
	3) a - g, b - h, c - f, d - e 4) a - h, b - e, c - f, d - g
3.	Match the following :
	Process Workdone a) Adiabatic process e) zero
	b) Isothermal process f) $P(V_1 - V_2)$
	c) Isochoric process g) $RT \log \left(\frac{P_1}{P_2} \right)$
	d) Isobaric process h) $\frac{R}{\nu - 1}(T_1 - T_2)$
	1) a - h, b - g, c - e, d - f
	2) a - h, b - g, c - f, d - e 3) a - g, b - e, c - f, d - h
	4) a - h, b - f, c - e, d - g
4.	Match the following : List - I List - II
	a) Isothermal process e) $dQ = dU+dW$ b) Adjabatic process f) $dQ = dW$
	b) Adiabatic process f) dQ = dW c) Isobaric process g) dQ = dU
	d) Isochoric process h) dU = -dW 1) a - h, b - e,c - g, d - f
	2) a - f, b - e, c - h, d - g 3) a - f, b - h, c - e, d - g
	4) a - h, b - f, c - g, d - e
5.	Match the following : List - I List - II
	a) SI units of thermal capacity e) Jkg^{-1}
	b) SI units of specific heat f) Jk^{-1}
	c) SI units of latent heat g) $Jkg^{-1}k^{-1}$
	d) Shupit of Internal anargy (b) T
	d) SI unit of Internal energy h) <i>J</i> 1) a - f, b - g, c - e, d - h
	2) a - g, b - e, c - f, d - h 3) a - f, b - e, c - g, d - h
	4) a - e, b - f, c - h, d - g
6.	Match the following : List - I List - II
	a) workdone on the system e) Positive
	b) heat energy given to the f) negative system
	c) change in internal energy g) may be +ve or -ve
	d) during cyclic process h) zero
	change in internal energy 1) a - g, b - e, c - h, d - f
	2) a - f, b - g, c - e, d - h 3) a - f, b - e, c - g, d - h
	4) a - g, b - e, c - h, d - g

7.	Match the following : List - I	List - II
	a) Among solids and liquids	e) Mercury specific heat is maximum for f) Water minimum for
	c) With increaseof pressured) With increase ofpressure	melting point of a solid substance
8.	1) a - f, b - e, c - h 2) a - e, b - f, c - g 3) a - e, b - g, c - f 4) a - f, b - e, c - g Match the following :	, d - h
	Nature of Gas	Specific Heat at Constant volume(C_v)
	a) Monoatomic	e) $\frac{6R}{2}$
	b) Diatomic	f) $\frac{4R}{2}$
	c) Triatomic	g) $\frac{3}{2}R$
	d) Mixture of equal	h) $\frac{5}{2}R$
	moles of di and mono atomic gases	2
9.	1) a - e, b - f, c - g 2) a - g, b - h, c - f 3) a - f, b - g, c - h 4) a - g, b - h, c - e Match the following : Nature of the Gas	, d - e n, d - e e, d - f Fraction of interanl
	a) Triatomic gas	d, ,
10.	4) a - d, b - f, c - e Match the following :	Э,
a) Iso	List - I	List - II one in the process
,	W = nRT	
b) Iso	choric f) PV grap	oh is a straight line par
c) Adi	iabtic g) first law	volume axis of thermo dynamics
d) Iso		<i>du</i> udden and quick
	process 1) a - f, b - g, c - h, c 2) a - g, b - f, c - h, c 3) a - h, b - e, c - f, c 4) a - e, b - g, c - e,c	l - e l - g

11 Match the following : 15. A: Zeroth law of thermodynamics gives us the Process Ex: for the process concept of energy a) Isothermal e) Any process in an open place **B:** Internal energy is dependent on temperature b) Adiabatic f) Melting of gold 1) Both Assertion and Reason are true and c) Isobaric g) Process inside a pressure cooker reason is correct explanation of Assertion. d) Isochoric h) Explosion of a bomb 2) Both Assertion and Reason are true but 1) a - g, b - f,c - e, d - h reason is not correct explanation of 2) a - f, b - h, c - g, d - e Assertion. 3) a - h, b - g, c - f, d - e 3) Assertion is true but reason is false 4) a - f, b - h, c - e, d - g 4) Assertion is false but reason is true 12. Match the following : 16. Assertion (A) : Two systems which are in ther-List-I List - II mal equilibrium with a third system are in thera) Heat cannot flow d) Zeroth law of mal equilibrium with each other. from a body at low thermodynamics Reason (R): The heat flows always from a systemperature to a body tem at higher temperature to a system at a at a higher temperature lower temperature 1) Both Assertion and Reason are true and by itself b) Introduces concept e) Clausius statement reason is correct explanation of Assertion. of temperature 2) Both Assertion and Reason are true bur c) Based on the law of f) First law of reason is not correct explanation of Assertion. conservation of energy thermo dynamics 3) Assertion is true but reason is false 4) Both assertion and reason are false List II 17. 1) a - e, b - d, c - f, Assertion (A) : Specific heat at constant vol-2) a - e, b - f, c - d, ume is greater than the specific heat at con-3) a - f, b - e, c - d, stant pressure. Reason (R) : At constant volume, work will be 4) a - e, b - d, c - d, 13. Match the following : done by the gas. So specific heat at constant List - I List - II volume is more a) food is cooked faster e) melting point is low-1) Both Assertion and Reason are true and in a pressure cooker. ered with rise in reason is correct explanation of Assertion. 2) Both Assertion and Reason are true bur pressure. b) two blocks of ice when f) melting of ice into reason is not correct explanation of Assertion. pressed together forms a water 3) Assertion is true but reason is false single block 4) Both assertion and reason are false c) specific heat infinity g) boiling point is el 18. Assertion (A): In adiabatic compression, the evated with rise in internal energy and temperature of the system get decreased. pressure Reason (R) : The adiabatic compression is a d)specific heat zero h) dQ = 0 (adiabatic slow process process) 1) Both Assertion and Reason are true and reason is correct explanation of Assertion. 1) a - e, b - f, c - h, d - g 2) Both Assertion and Reason are true bur 2) a - g, b - e, c - f, d - h reason is not correct explanation of Assertion. 3) a - g, b - e, c - h, d - f 3) Assertion is true but reason is false 4) a - f, b - e, c - h, d - g 4) Both assertion and reason are false 14. Match the following : 19. Assertion (A): The isothermal curves intersect List - I List - II each other at a certain point. a) Thermos flask containing e) temp increases Reason (R) : The isothermal changes take coffee is vigorously shaken place slowly, so the isothermal curves have very b) door of a refrigerator is f) temp decreases little slope opened while it is working 1) Both Assertion and Reason are true and c) cycle tube gets punctured g) dQ = du, dw = 0reason is correct explanation of Assertion. 2) Both Assertion and Reason are true bur d)gas heated in a vessel h) dQ = 0reason is not correct explanation of Assertion. fitted with a tight piston 3) Assertion is true but reason is false 1) a - h, b - f, c - e, d - g 4) Both assertion and reason are false 2) a - h, b - e, c - f, d - g 3) a - h, b - g, c - e, d - f 4) a - g, b - h, c - e, d - f

20.	Assertion (A) : The ratio of $\frac{C_P}{C_v}$ for a diatomic	LEVEL IV
	gas is more than that for a monoatomic gas	COMPREHENSIVE
	Reason (R) : The molecules of monoatomic gas	1. Read the following passage to answer
	have more degrees of freedom than those of a	questions.
	diatomic gas	The internal energy of gases at room tempera- ture is due to the interaction of molecules and
	1) Both Assertion and Reason are true and	atoms of the gas, the translational, vibrational
	reason is correct explanation of Assertion.	and rotational kinetic enegies of molecules. The
	2) Both Assertion and Reason are true bur	temperature of a substance is a measure of
	reason is not correct explanation of Assertion.	average translational kinetic energy only. At
	3) Assertion is true but reason is false	absolute zero the translational kinetic energy
	4) Both assertion and reason are false	of molecules become zero and it is known as
21.	Assertion (A) : According to the principle of	zero point energy. At a given temperature gas melecules have random velocity and their av-
	conservation of energy all heat can be con-	erage velocity is zero. The rms value of gas
	verted into mechanical work	
	Reason (R) : Due to various losses, it is im-	3 <i>RT</i>
	possible to convert all heat into mechanical work	molecules is given by $\sqrt{\frac{3RT}{M}}$. You are given a sample
	1) Both Assertion and Reason are true and reason is correct explanation of Assertion.	of 2 moles of O_2 gas at 27°C, where the rotational
	2) Both Assertion and Reason are true bur	kinetic energy of each molecule is $18{ imes}10^{-23}J$.
	reason is not correct explanation of Assertion.	1. At zero point energy, a gas may posses
	3) Assertion is true but reason is false	a) vibrational kinetic energy b) elastic potential energy
0.0	4) Both assertion and reason are false	c) rotational kinetic energy
22.	Assertion (A): Cooking in a pressure cooker is faster Reason (R): Boiling point is elevated with rise	d) translational kinetic energy
	in pressure	1) a,c,d are correct 2) a,b,c are correct
	1) Both Assertion and Reason are true and	3) b,c,d are correct 4) a,b,d are correct
	reason is correct explanation of Assertion.	2. The total internal energy of the gas is
	2) Both Assertion and Reason are true bur	1) 7469 J 2) 216.8 J 3) 7685.8 J 4) 6432.2 J
	reason is not correct explanation of Assertion.	3. The average internal energy of each molecule
	3) Assertion is true but reason is false	is
	Both assertion and reason are false	1) $638 \times 10^{-23} J$ 2) $576 \times 10^{-23} J$
23.	Assertion (A) : According to Joule, heat and	
	work are related	3) $576 \times 10^{-23} J$ 4) $216 \times 10^{-23} J$
	Reason (R) : For every 1 cal. of heat we can get 4.186 J of mechanical work.	4. The total energy of gas at absolute zero is
	1) Both Assertion and Reason are true and	1) 0 2) 218.2 J
	reason is correct explanation of Assertion.	3) 816 J 4) 216.8 J
	2) Both Assertion and Reason are true bur	KEY
	reason is not correct explanation of Assertion.	1) 2 2) 3 3) 2 4) 4
	3) Assertion is true but reason is false	II. For 20 g of H ₂ O the phase diagram is given.
	4) Both assertion and reason are false	Using the diagram answer the following
24.	Assertion (A) : During an adiabatic process,	questions.
	an ideal gas expands by decrease in its inter-	Ϋ́
	nal energy only	steamline)
	Reason (R) : During an adibatic process, heat	$10 atm$ $\sqrt{ice line}$
	cannot be exchanged between a system and	
	its surroundings	
	 A is true and R is false Both A and R are true and R is the correct 	150
	explanation of A	4.58 atm
	3) Both A and R are true and R is not correct	3 - hoar/frost line
	explanation of A	2 7
	4) A is false and R is true	
	KEY	$-40^{\circ}C$ $0.01^{\circ}C$
	01.1 02.1 03.1 04.3 05.1 06.3	MAR II 1923.08 76
	07.4 08.4 09.1 10.1 11.4 12.1	1. The maximum temperature at which ice is pos-
	13.2 14.2 15.4 16.1 17.4 18.4	
	19.4 20.4 21.2 22.1 23.1 24.2	1) 0°C 2) 0.01°C 3) 100°C 4) 58.6°C
		·

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2. When the system is moved from -40°C to 40°C
at 5 atm pressure, its phase changes from
1) ice to vapour 2) ice to water to vapour
3) water to vapour 4) vapour to ice
3. When the system is moved from -10°C to 10°C
at 2.4 atm pressure, its phae changes from
1) ice to vapour through water
2) water to vapour gradually decreased from
8 atm to 3 atm. Then the phase changes from
1) water to ice to vapour
2) ice to wapour without entering into water
4) water to ice to vapour
2) ice to water to vapour
3) water to ice to vapour
2) ice to water to vapour
3) water to apour without entering into ice
5. For water at 6 atm pressure and at a tempera-
ture 18°C, the pressure is reduced to 4.58 atm.
The matter that exist at new
coordinates is in the form of
1) loe 2) water 3) vapour
4) co-existance of all three
KEY
1) 2 2) 2 3) 1 4) 3
5) 2
PREVIOUS EAMCET QUESTIONS
1. The tyre of a motor car contains air at 15°C if
the temperature increases to 35°C, the approxi-
mate percentage increase in pressure is (ig-
nore the expansion of tyre) (2005E)
1) 7 2) 9 3) 11 4)13
2. The ratio of specific heats of a gas is
$$\gamma$$
. The
change in internal energy of one mole of the
gas when the volume changes from V to 2V at
constant pressure "P" is (2005E)
1) $\frac{\gamma - 1}{PV}$ 2) PV 3) $\frac{PV}{\gamma - 1}$ 4) V
3. A 42 kg block of ice moving on rough horizon-
tal surface stops due to friction, after some
time. If the initial velocity of the decelerating
block is 4 ms⁻¹, the mass of ice (in kg) that
has melted due to the heat generated by the
friction is (latent heat of ice is $3.36 \times 10^5 Jkg^{-1}$
(2005E)
1) 10^{-3} 2) 1.5×10^{-3}
4. A) First law of thermodynamics specifies the
conditions under which a body can use its heat
energy to produce the work.
B) Second law of thermodynamics states that heat
always flows from hot body to cold body by itself

1) Both A and B are true

2) Both A and B are false

3) A is truee but B is false

4) A is false B is true

If 4 moles of an ideal monoatomic gas at temperature 400K is mixed with 2 moles of another ideal monoatomic gas at temperature 700K, the temperature of the mixture is (Volume constant (2004E)

1) 550°C 2) 500°C 3) 550K 4) 536 K The pressure and density of a given mass of a diatomic gas ($\gamma = 7/5$) change adiabatically

from (P, d) to (P^I, d^I). If
$$\frac{d^{I}}{d} = 32$$
 then $\frac{P^{I}}{P}$ is (γ

=ratio of specific heats)(2004E)

1) $\frac{1}{128}$ 2) $\frac{1}{64}$ 3) 64 4) 128

When a heat of Q is suppiled to one mole of a monoatomic gas ($\gamma = 5/3$), Then the molar heat capacity of the gas at constant volume is (2004M)

1)
$$\frac{3R}{4}$$
 2) $\frac{5R}{4}$ 3) $\frac{7R}{4}$ 4) $\frac{3R}{2}$

The sample of the same gas, x, y and z, for which the ratio of specific heats is $\gamma = 3/2$ have initially the same volume. The volume of the each sample is doubled by adiabatic process in the case of x, by isobaric process in the case of y and by isothermal process in the case of z. If the initial pressure of the sample

of x, y and z are in the ratio $2\sqrt{2}$:1:2 then the

ratio of the their final pressures is

(2004M)

1)2:1:12) 1 : 1 : 1 3) 1 : 2 : 1 4) 1 : 1 : 2

The temperature of 5 moles of a gas at constant volume is changed from 100°C to 120°C. The change in internal energy is 80J. The heat capacity of the gas at constant volume will be in Joule/Kelvin is (2003E)

4) 0.4 1)8 2)4 3) 0.8 During an adiabatic process, the pressure of a gas is proportional to the cube of its adiabatic

temperature. The value of $\frac{C_P}{C_v}$ for that gas (2003M) is

1)
$$\frac{3}{5}$$
 2) $\frac{4}{3}$ 3) $\frac{5}{3}$

4) $\frac{3}{2}$

5 moles of hydreogen $\left(\gamma_1 = \frac{7}{5}\right)$ initially at

S.T.P is compressed adiabatically so that its temperature becomes 400° c. The increase in internal energy in kilojoules is (R = 8.30)J/mole/k) (2002E) 2.41.5 3.65.55 4.80.55 1.21.55

12. A metal sphere of radius 'r ' and specific heat 20. A flask is filled with 13 g of an ideal gas at 27° 's' is rotating about an axis passing through C and its temperature is raised to 52°C. The its centre at a speed of n rotations per sec is mass of the gas that has to be released to suddenly stopped and 50% of its energy is used maintain the temperature of the gas in the flask in increasing its temperature. The rise in temat 52°C and the pressure remaining the same perature of the sphere is (2002 E) is (2000E) 1.2.5g 2.2.0 g 3.1.5 g 4.1.0 g 1. $\frac{2}{5} \frac{\pi^2 r^2 n^2}{s}$ 2. $\frac{1}{10} \frac{\pi^2 n^2}{r^2 s}$ 21. An ideal gas at a pressure of 1 atmosphere and temperature of 27°C is compressed adiabatically until its pressure becomes 8 times the $3.\frac{7}{8}\frac{\pi r^2 n^2 s}{1} \qquad 4.\frac{5}{14}\frac{\pi^2 r^2 n^2}{s}$ initial pressure, then the final temperature is--- $-----\left(\gamma = \frac{3}{2}\right)$ 13. A gas under constant pressure of 4.5×10^5 pa (2000 E) when subjected to 800KJ of heat, changes the volume from 0.5 m³ to 2.0m³. The change in 1.627°C 2.527°C 3.427°C 5. 327°C the internal energy of the gas. 22. Two liquids at temperatures 60°C and 20°C re-(2002 M) spectively have masses in the ratio 3:4 and their 2.5.25 × 10⁵J 1. $6.75 \times 10^5 \text{ J}$ specific heats in the ratio 4:5. If the two liquids 3. 3.25 × 10⁵J 4.1.25 × 10⁵J are mixed, the resultant temperature is (2000E) If for hydrogen $c_p - c_v = m$ and for nitrogen $c_p - c_v = n$, where c_p and c_v refer to specific heats per unit mass respectively at constant pres-14. 1.70°C 2.40°C 3.50°C 4.35°C 23. A steel ball of mass 0.1 kg falls freely from a height of 10m and bounces to a height of 5.4 m sure and constant volume. The relation between from the ground. If the dissipated energy in m and n is (2002 M) this process is absorbed by the ball, the rise 2.n = 7m 1.n = 14m in its temperature is (Specific heat of steel = 4.m= 14n 3.m = 7n $460J/kg/^{\circ}C, g = 10ms^{-2}$ (2000 M) A lead bullet of 10 g travelling with 300 m/s 15. 1. 0.01°C 2. 0.1°C 3. 1°C 4.1.1°C strikes against a block of wood and comes to 50g of copper is heated to increase its tem-24. rest. Assuming 60% of heat is absorbed by the bullet, the increase in its temperature is (Sp. perature by 10°C. If the same quantity jof heat is given to 10 g of water, the rise in its temheat of lead is 150 Jkg⁻¹ k⁻¹) (2001E) 1. 10°C 2. 12.5°C 3. 18°C 4. 20°C (2000M) perature is (Specific heat of copper = 420J/kg/°C 16. The pressure and density of a diatomic Specific heat of water = 4200J/kg/°C $gas\left(\gamma = \frac{7}{5};\right)$ changes adiabatically from (p,d) 2. 6°C 3. 7°C 1. 5⁰C 4.8°C 25. A liquid of mass 'm' and specific heat 'S' is at a temperature '2t'. If another liquid of thermal to (p^1,d^1) . If $\frac{d'}{d}$ = 32 then $\frac{p'}{p}$ is cpacity 1.5 times, at a temperature of t/3 is (2001E) added to it, the resultant temperature will be (1999E) 1. $\frac{1}{128}$ 2. 32 3. 128 4. 256 1. $\frac{4}{3}t$ 2. t 3. $\frac{t}{2}$ 4. $\frac{2}{3}t$ 17. A gas for which $\gamma = 1.5$ is suddenly compressed 26. When heat energy of 1500 J is supplied to a gas at constant pressure is 2.1X10⁵ N/m² there to $\frac{1}{4}$ th of its initial value then the ratio of the was a increase in its volume equal to 2.5 X 10⁻ ³ m³. The increase in interanal energy of the final initial to pressure is gas, in joules is (1999E) (2001 M) 1.450 2.525 3.975 4.2025 1.1:16 2.1:8 3.1:4 4.8:1 27. The relation between melting point of ice and 18. I mole of an ideal gas with $\gamma = 1.4$ is adiabatically pressure is shown by ice line, which will be compressed so that its temperature rises from 27°C (1999E) to 35°C. The change in internal energy of the gas 1. with a positive slope is (R= 8.3 Joule mole-1 k-1) (2001M) 2. with a negative slope 1. -166J 2. + 166J 3. -168 J 4. + 168 3. parallel to pressure axis 19. During an adiabatic process, if the pressure of 4. parallel to temperature axis the ideal gas is proportional to the cube of its 28. For an adiabatic change in a gas, if P, V,T detemperature, the ratio $\gamma = \frac{c_p}{c_y}$ is (C_p-Specific notes pressure, volume and absolute temperature of gas at any time and γ is the ratio of specific heats of gas, which jof the following heat at constant pressure C_v-Specific heat at equation is ture? (1999 M) constant volume) (2000E) 1. $\frac{7}{5}$ 2. $\frac{4}{3}$ 3. $\frac{5}{3}$ 4. $\frac{3}{2}$ ¹· $T^{\gamma}P^{1-\gamma} = \text{const.}$ ²· $T^{\nu\gamma}P^{\gamma} = \text{const.}$ 4. $T^{\gamma}V^{\gamma} = const.$ 3. $T^{\gamma-1}V^{\gamma} = \text{const.}$

JR. PHYSICS

THERMODYNAMICS

29.	Boiling water at 100ºC and cold water at tºC are mixed in the ratio 1:3 and the resultant	41.
	maximum temperature was 37ºC. Assuming no heat losses, the value of 't' is (1999M)	42.
30.	1. 4° C 2. 9° C 3. 12° C 4. 16° C 20g of ice and 20 g of hot water are mixed,	
30.	when the ice is melted the temperature of the	
	mixture was found to be 0° C. The temperature	43.
	of hot water taken should be (L _{ice} =80 cal/g) (1999 M)	
31.	1. 40°C 2. 72°C 3. 80°C 4. 96°C A lead bullet of mass 21 g travelling at a speed	
01.	of 100 ms ⁻¹ comes to rest in a wooden block.	44.
	If no heat is taken away by the wood, the rise in temperature of the bullet in the wood nearly	
	is (Sp. heat of lead 80 cal/kg/ºc)	
	(1998E) 1. 25ºc 2. 28ºC 3. 33ºC 4. 15ºC	
32.	The material that has largest specific heat is (1998 E)	
	1. mercury 2. water	
33.	3. iron 4. diamond First law of thermodynamics states that	
0.05	(1998M)	
	1. system can do work 2. system has temperature	
	3. system has pressure	
34.	4. heat is form of energy An amount of water of mass 20g, at 0°C is	
	mixed with 40 g of water at 100°C, Final tem-	
	perature of mixture is (1996E) 120ºC 2. 6.66ºC 3. 5ºC 4. 0ºC	
35.	A piece of lead falls from a height of 100m on a fixed non-conducting slab which brings it to	
	rest. The temperature of the lead piece imme-	1.
	diately after collision increases by (Sp.heat of lead is 30.6cal/kg/°C and g= 9.8 m/sec ² .)	
	(1996E)	2.
36.	1. 0K 2. 27°C 3. 7.62K 4. 4.2K A diatomic gas molecule has translational, ro-	2.
	tational and vibrational degrees of freedom.	
	Then $\frac{c_p}{c}$ is (1995 E)	3.
	v	
37.	1. 1.67 2. 1.4 3. 1.29 4. 1.33 An iron ball of mass 0.2 kg is heated to 100°C when	
	it is put in ice block at 0 ^o C, 25 g of ice is melted.	4.
	The specific heat of iron is C.G.S Units (1995 E)	.
20	1.1 2.0.1 3.0.8 4.0.08	
38.	The amount of heat required to convert 10g of ice at -10°C into steam at 100°C is (in cal.)	
	(1995 M) 1. 5400 2. 6400 3. 7200 4. 7250	
39.	1g of steam is sent into 1g of ice. The result-	
	ant temperature of the mixture is (1995 M) 1. 270°C 2. 230°C 3. 100°C 4. 50°C	
40.	0.1 moles of a diatomic gas at 27º c is heated	
	at constant pressure, so that the volume is doubled. If R=2 cal/mol, the work done is (1994	
	E) 1. 150cal 2. 60 cal 3. 40 cal 4. 30 cal	
	1. 1900al 2. 00 0al 3. 40 0al 4. 30 0al	
		1

	1. Gay-Lussac's law 2. Charles law 3. Boyle's law 4. Dalton's law In P-V graphs, the ratio of slope of adiabatic to isothermal is (1994M)					
	1. γ-1	2. $\frac{1}{\gamma}$	3.1	4.γ		
	Latent heat of vaporization of water is 538 cal/ g. During the conversion of 1g of water to steam, if the external work done is 168J, the increase in internal energy is (1994 M) 1) 638cal 2) 538cal 3) 498cal 4) 80cal A quantity of heat 'Q' is supplied to a monoatomic ideal gas which expands at con- stant pressure. The fraction of heat that goes into workdone by the gas is (1993 E)					
	1. $\frac{2}{5}$	2. $\frac{3}{5}$	3. $\frac{2}{3}$	4.1		
KEY						
	01)1	02)3	03)1	04)4	05)4	
	06)4	07)4	08)2	09)3	10)4	
	11)2	12)1	13)4	14)3	15)3	
	16)3	17)2	18)2	19)4	20)4	
	21)4	22)4	23)2	24)1	25)2	
	26)3	27)2	28)1	29)4	30)3	
	31)4	32)2	33)1	34)2	35)3	
	36)2	37)2	38)4	39)3	40)1	
	41)3	42)4	43)3	44)1	-	

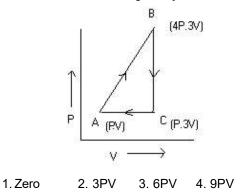
The law obeyed by isothermal process (1994E)

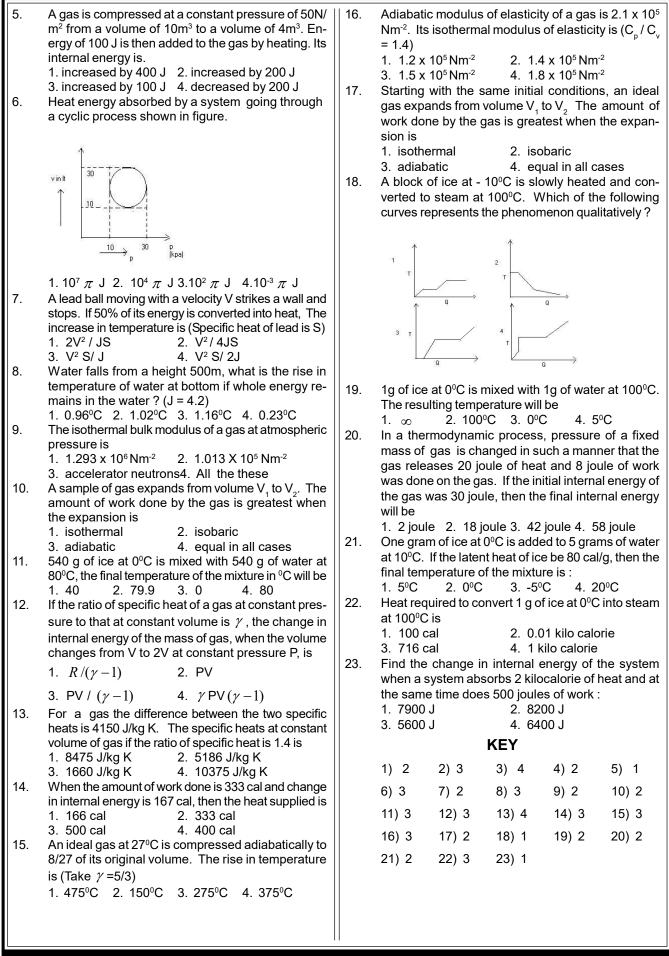
NUMERICALS FROM OTHER COMPETETIVE EXAMINATIONS

. 110 Joules of heat are added to a gaseous system, whose internal energy is 40 J; then the amount of external work done is

1. 150 J 2. 70 J 3. 110 J 4. 40 J At 27°C a gas compressed suddenly such that its pressure becomes 1/32 of original pressure. Final temperature will be ($\gamma = 5/3$)

- temperature will be $(\gamma = 5/3)$ 1. 420°K 2. 300°K 3. 927°C 4. 327°C 3. One mole of an ideal gas requires 207 J heat to raise the temperature by 10k when heated at constant pressure. If the same gas is heated at constant volume to raise the temperature by the same 10k, the heat required is
- 1. 198.7 J 2. 29 J 3. 215.3 J 4. 124J A Sample of ideal monoatomic gas is taken round the cycle ABCA as shown in the figure. The work done during the cycle is





HINTS10.work done is max for isobaric process1.dg=du+dw2.
$$T_r^T P_r^{T_r} P_r^{T_r} P_r^{T_r} P_r^{T_r}$$
3. $C_{p-5}2R_{r}C_{v-3}2R$ $\frac{Q}{QP} = 3$ free $Q_P = 700.Qv=124J(nearly)$ 4.W=area enclosed by the indicatordiagram $= \frac{1}{2}(3V-V)(4P-P)=3PV$ 5.Acc to 1* low of thermodynamics $dQ=4u+Pdv$ since dv is negative, we get du=400.J6.W=area of loop = $r^{r}e=r(10)2$ J7. $\frac{1}{2}mv^2 \frac{1}{2} = JH = Jmc \partial$ $\theta = \frac{y^2}{4Jc}$ 8.mgh=mcd β 9.we know Pv=kPdw+Vdp=0 $P = \frac{dP}{dy/v} = Bi$ $x = \frac{dP}{dy/v} = Bi$ $x = \frac{dP}{dy/v} = Bi$