

SUMMARY

- ⇒ Thermal equilibrium, zeroth law of thermodynamics, Concept of temperature.
- ⇒ Heat, work and internal energy. First law of thermodynamics.
- ⇒ Second law of thermodynamics, reversible and irreversible processes.
- ⇒ Isothermal and adiabatic process.
- ⇒ Carnot engine and its efficiency Refrigerators.

$$\Rightarrow \frac{C - 0}{100^{\circ}} = \frac{F - 32}{180^{\circ}} \Leftrightarrow \frac{\Delta C}{100^{\circ}} = \frac{\Delta F}{180^{\circ}}$$

$$\Rightarrow w = P \Delta V \text{ (Isothermal process)}$$

$$\mu RT \ln \frac{V_2}{V_1} \text{ (Isothermal process)}$$

$$PV = \text{Const (Isothermal process)}$$

$$P V^{\gamma} = \text{Const (adiabatic process)}$$

$$P^{1-\gamma} T^{\gamma} = \text{Const}$$

$$TV^{\gamma-1} = \text{Constant}$$

$$r = \frac{c_p}{c_v}$$

$$\text{Monoatomic gases } C_v = \frac{3}{2} R, C_p = \frac{5}{2} R, \gamma = 1.67$$

$$\text{Diatomic gases } C_v = \frac{5}{2} R, C_p = \frac{7}{2} R, \gamma = 1.4$$

$$\text{Polyatomic gases } C_v = \frac{7R}{2}, C_p = \frac{9R}{2}, \gamma = 1.4$$

$$W = \frac{1}{\gamma-1} (P_1 V_1 - P_2 V_2) \text{ (adiabatic Process)}$$

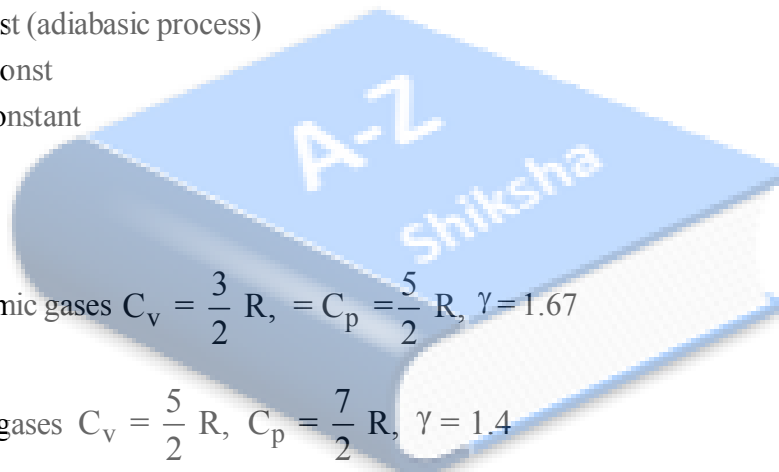
$$= \frac{mR (T_1 - T_2)}{\gamma-1}$$

$$\Rightarrow \Delta Q = \Delta u + \Delta W \text{ (First law of thermodynamics)}$$

$$\Delta Q = \Delta U = \mu C_v \Delta t \text{ (V = Const)}$$

$$\Delta Q = \mu C_p \Delta t \text{ (P = Const)}$$

$$\Delta u = 0 \text{ (isothermal process cyclic process)}$$



⇒ Coefficient of Performance of refrigerators

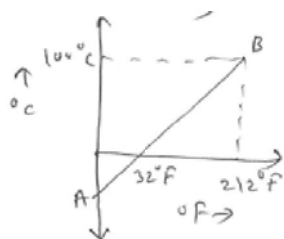
$$n = \frac{W}{Q_1}$$
$$= \frac{Q_1 - Q_2}{Q_1}$$
$$= \frac{T_1 - T_2}{T_1}$$

⇒ Coefficient of performance of refrigerators.

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

⇒ Ideal gas $C_p - C_v = R$

1. A difference of temperature of 25°C is equivalent to a difference of
(A) 72°F (B) 45°F (C) 32°F (D) 25°F
2. What is the value of absolute temperature on the Celsius Scale?
(A) -273.15°C (B) 100°C (C) -32°C (D) 0°C
3. The temperature of a substance increases by 27°C What is the value of this increase of Kelvin scale?
(A) 300K (B) $2-46\text{K}$ (C) 7K (D) 27K
4. At Which temperature the density of water is maximum?
(A) 4°F (B) 42°F (C) 32°F (D) 39.2°F
5. The graph AB Shown in figure is a plot of a temperature of a body in degree Fahrenheit than slope of line AB is



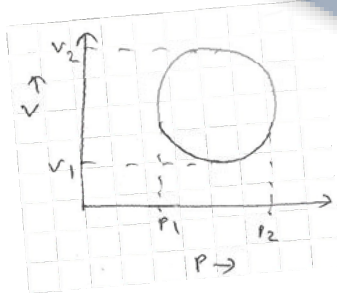
- (A) $\frac{5}{9}$ (B) $\frac{9}{5}$ (C) $\frac{1}{9}$ (D) $\frac{3}{9}$

6. The temperature on celsius scale is 25°C What is the corresponding temperature on the Fahrenheit Scale?
 (A) 40°F (B) 45°F (C) 50°F (D) 77°F
7. The temperature of a body on Kelvin Scale is found to be $x\text{K}$. when it is measured by Fahrenheit thesmometes. it is found to be $x^{\circ}\text{F}$, then the value of x is .
 (A) 313 (B) 301.24 (C) 574-25 (D) 40
8. A Centigrade and a Fahrenheit thesmometes are dipped in boiling wates-The wates temperature is lowered until the Farenhit thesmometes registered 140° what is the fall in thrmometers
 (A) 80° (B) 60° (C) 40° (D) 30°
9. A uniform metal rod is used as a bas pendulum. If the room temperature rises by 10°C and the efficient of line as expansion of the metal of the rod is, $2 \times 10^{-6} 0_c^{-1}$ what will have percentage increase in the period of the pendulum ?
 (A) -2×10^{-3} (B) 1×10^{-3} (C) -1×10^{-3} (D) 2×10^{-3}
10. A gas expands from 1 litre to 3 litre at atmospheric pressure. The work done by the gas is about
 (A) 200 J (B) 2 J (C) 300 J (D) $2 \times 10^5 \text{ J}$

11. Each molecule of a gas has f degrees of freedom. The ratio $\frac{C_P}{C_V} = \gamma$ for the gas is

(A) $1 + \frac{f}{2}$ (B) $1 + \frac{1}{f}$ (C) $1 + \frac{2}{f}$ (D) $1 + \frac{(f-1)}{3}$

12. Is the cyclic Process Shown on the $V \rightarrow P$ diagram, the magnitude of the work done is



(A) $\pi \left(\frac{P_2 - P_1}{2} \right)^2$ (B) $\pi \left(\frac{V_2 - V_1}{2} \right)^2$

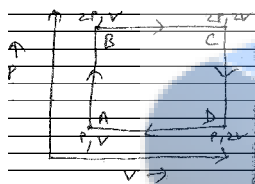
(C) $\pi (P_2 V_2 - P_1 V_1)$ (D) $\frac{\pi}{4} (P_2 - P_1)(V_2 - V_1)$

13. If the ratio of specific heat of a gas at Consgant pressure to that at constant volume is γ , the Change in internal energy of the mass of gas, when the volume changes from V to $2V$ at Constant Pressure p , is

(A) $\frac{PV}{\gamma-1}$ (B) $\frac{R}{\gamma-1}$ (C) PV (D) $\frac{\gamma PV}{\gamma-1}$

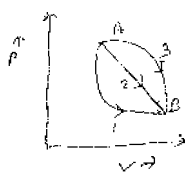
14. The change in internal energy, when a gas is cooled from 927°C to 27°C
 (A) 200% (B) 100% (C) 300% (D) 400%
15. For hydrogen gas $C_p - C_v = a$ and for oxygen gas $C_p - C_v = b$, The relation between a and b is given by
 (A) $a = 4b$ (B) $a = b$ (C) $a = 16b$ (D) $a = 8b$
16. In a thermodynamic process, pressure of a fixed mass of a gas is changed in such a manner that the gas release 20J of heat and 8J of work has done on the gas- If the initial internal energy of the gas was 30J, then the final internal energy will be
 (A) 58 J (B) 2 J (C) 42 J (D) 18 J
17. If for a gas $\frac{C_p}{C_v} = 1.67$, this gas is made up to molecules which are
 (A) diatomic (B) Polytomic
 (C) monoatomic (D) mixnese of diatomic and polytomic molecules

18. An ideal monoatomic gas is taken around the cycle ABCDA as Shown in the $P \rightarrow V$ diagram. The work done during the cycle is given by



- (A) PV (B) $\frac{1}{2} PV$ (C) $2 PV$ (D) $4 PV$

19. A given mass of a gas expands from state A to B by three different paths 1, 2 and 3 as shown in the figure. If W_1, W_2 and W_3 respectively be the work done by the gas along the three paths, then



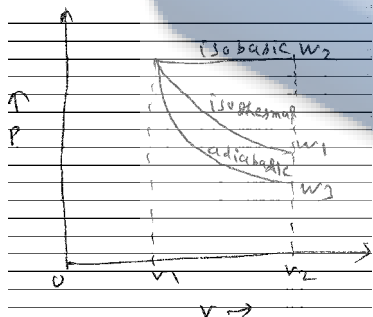
- (A) $W_1 > W_2 > W_3$ (B) $W_1 < W_2 < W_3$ (C) $W_1 = W_2 = W_3$ (D) $W_1 < W_2, W_1 < W_3$
20. One mole of a monoatomic gas $\left(\gamma = \frac{5}{3}\right)$ is mixed with one mole of A diatomic gas $\left(\gamma = \frac{7}{5}\right)$ what will be value of γ for mixture?
 (A) 1-454 (B) 1-4 (C) 1-54 (D) 1-5
21. If du represents the increase in internal energy of a thesmodynamic system and dw the work done by the system, which of the following statement is true?
 (A) $du = dw$ in isothermal process (C) $du = - dw$ in an aidabadic process
 (B) $du = dw$ in aidabadic process (D) $da = - dw$ in an isothermal process

22. One mole of a monoatomic ideal gas is mixed with one mole of a diatomic ideal gas. The molar specific heat of the mixture at constant volume is
- (A) $4R$ (B) $3R$ (C) R (D) $2R$
23. One mole of a monoatomic gas is heated at a constant pressure of 1 atmosphere from 0K to 100K . If the gas constant $R = 8.32\text{ J/mol K}$, the change in internal energy of the gas is approximately?
- (A) 23 J (B) $1.25 \times 10^3\text{ J}$ (C) $8.67 \times 10^3\text{ J}$ (D) 46 J
24. A gas mixture consists of 2 moles of oxygen and 4 moles of argon at temperature T . Neglecting all vibrational modes, the total internal energy of the system is
- (A) $11RT$ (B) $9RT$ (C) $15RT$ (D) $4RT$
25. 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 $\frac{T_1}{T_2}$ is given by

- (A) $\left(\frac{L_1}{L_2}\right)^{2/3}$ (B) $\left(\frac{L_2}{L_1}\right)^{2/3}$ (C) $\frac{L_1}{L_2}$ (D) $\frac{L_2}{L_1}$

26. Starting with the same initial conditions, an ideal gas expands from volume V_1 to V_2 in three different ways. The work done by the gas is W_1 if the process is purely isothermal, W_2 if purely isobaric and W_3 if purely adiabatic. Then



- (A) $W_2 > W_1 > W_3$ (B) $W_2 > W_3 > W_1$
 (C) $W_1 > W_2 > W_3$ (D) $W_1 > W_3 > W_2$
27. In a given process on an ideal gas $dw = 0$ and $dQ < 0$. Then for the gas
- (A) the volume will increase (B) the pressure will remain constant
 (C) the temperature will decrease (D) the temperature will increase
28. A wafer of volume 2 liter in a container is heated with a coil of 1 kW at 27°C . The lid of the container is open and energy dissipates at the rate of $160\frac{\text{J}}{\text{s}}$. In how much time will the temperature rise from 27°C to 77°C . Specific heat of wafers is $4.2\frac{\text{KJ}}{\text{Kg}}$
- (A) 7 min (B) $6\text{ min } 2\text{ s}$ (C) 14 min (D) $8\text{ min } 20\text{ s}$

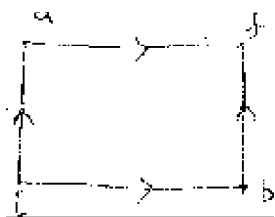
29. 70 calorie of heat are required to raise the temperature of 2 mole of an ideal gas at constant pressure from 30°C to 35°C
The amount of heat required to raise the temperature of the same gas through the same range at constant volume is calorie.
(A) 50 (B) 30 (C) 70 (D) 90
30. When an ideal diatomic gas is heated at constant pressure, the fraction of the heat energy supplied which increases the internal energy of the gas is..
(A) $\frac{3}{7}$ (B) $\frac{3}{5}$ (C) $\frac{2}{5}$ (D) $\frac{5}{7}$
31. Two cylinders A and B fitted with piston 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 30K, then the rise in temperature of the gas in B is.
(A) 30 K (B) 42 K (C) 18 K (D) 50 K
32. An insulated container containing monoatomic gas of molar mass M_0 is moving with a velocity, V . If the container is suddenly stopped, find the change in temperature.
(A) $\frac{M_0 v^2}{5R}$ (B) $\frac{M_0 v^2}{4R}$ (C) $\frac{M_0 v^2}{3R}$ (D) $\frac{M_0 v^2}{2R}$
33. A small spherical body of radius r is falling under gravity in a viscous medium. Due to friction the medium gets heated. How does the rate of heating depend on radius of body when it attains terminal velocity!
(A) r^2 (B) r^3 (C) r^4 (D) r^5
34. The first law of thermodynamics is concerned with the conservation of
(A) momentum (B) energy (C) mass (D) temperature
35. If heat given to a system is 6 kcal and work done is 6 kJ. The change in internal energy is KJ.
(A) 12.4 (B) 25 (C) 19.1 (D) 0
36. The internal energy change in a system that has absorbed 2 Kcal of heat and done 500J of work is
(A) 7900 J (B) 4400 J (C) 6400 J (D) 8900 J
37. Which of the following is not a thermodynamical function.
(A) Enthalpy (B) Work done
(C) Gibbs' energy (D) Internal energy
38. Which of the following is not a thermodynamic co-ordinate.
(A) R (B) P (C) T (D) V
39. The work of 62-25 KJ is performed in order to compress one kilo mole of gas adiabatically and in this process the temperature of the gas increases by 5°C The gas is _____ $R = 8.3 \frac{\text{J}}{\text{mole K}}$
(A) triatomic (C) monoatomic
(B) diatomic (D) a mixture of monoatomic and diatomic

40. C_p and C_v denote the specific heat of oxygen per unit mass at constant Pressure and volume respectively, then

(A) $c_p - c_v = \frac{R}{16}$ (B) $C_p - C_v = R$

(C) $C_p - C_v = 32 R$ (D) $C_p - C_v = \frac{R}{32}$

41. When a System is taken from State i to State f along the path iaf, it is found that $Q = 70$ cal and $w = 30$ cal, along the path ibf. $Q = 52$ cal. W atoug the path ibf is



- (A) 6 cal (B) 12 cal (C) 24 cal (D) 8 cal

42. One kg of adiatomic gas is at a pressure of $5 \times 10^5 \frac{N}{m^2}$ The density of the gas is $\frac{5kg}{m^3}$ what is the energy of the gas due to its thermal motion ?

- (A) $2.5 \times 10^5 J$ (B) $3.5 \times 10^5 J$ (C) $4.5 \times 10^5 J$ (D) $1.5 \times 10^5 J$

43. 200g of water is heated from $25^\circ C$ to $45^\circ C$ Ignoring the slight expansion of the water the change in its internal energy is (Specific heat of wafer $1 \frac{cal}{9^\circ C}$)

- (A) 33.4 KJ (B) 11.33 KJ (C) 5.57 KJ (D) 16.7 KJ

44. During an adiabatic process, the pressure of a gas ifound to be propostional to the fifth power of its absolute temperature. The radio $\frac{c_p}{c_v}$ for the gas is

- (A) $\frac{4}{5}$ (B) $\frac{3}{4}$ (C) $\frac{5}{4}$ (D) 4

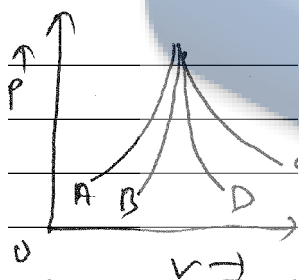
45. One mole of oxygen is heated at constant pressure stasting at $0^\circ C$. How much heat energy in cal must be added to the gas to double its volume ? Take $R = 2 \frac{cal}{molk}$

- (A) 1938 (B) 1920 (C) 1911 (D) 1957

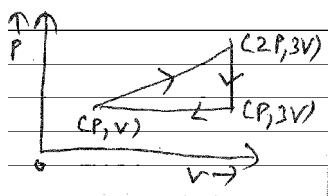
46. μ moles of a gas filled in a containes at temperature T is in equilibrium initially - If the gas is compressed slowly and is thesmally to half its initial volume the work done by the atmosphere on the piston is

- (A) $-\frac{RT}{2}$ (B) $\frac{\mu RT}{2}$ (C) $\mu RT \ln(2 - \frac{1}{2})$ (D) $-\mu RT / n_2$

47. Heat capacity of a body depends on the as well as on
- (A) material of the body, its mass (C) mass of the body, its temperature
 (B) material of the body, its temperature (D) Volume of the body, its mass
48. In thermodynamics, the work done by the system is considered and the work done on the system is Considered
- (A) Positive, zero (B) negative, Positive
 (C) zero, negative (D) Positive, negative
49. A thermodynamic system goes from States
 (i) P, V to $2P, V$ (ii) P, V to $P, 2V$. Then what is work done in the two Cases.
 (A) Zero, PV (B) Zero, Zero (C) PV , Zero (D) PV, PV
50. For free expansion of the gas which of the following is true ?
 (A) $Q = 0, W > 0$ and $\Delta E_{int} = -W$ (B) $W = 0, Q > 0$ and $\Delta E_{int} = Q$
 (C) $W > 0, Q < 0$ and $\Delta E_{int} = 0$ (D) $Q = W = 0$ and $\Delta E_{int} = 0$
51. For an adiabatic process involving an ideal gas
 (A) $P^{\gamma-1} = T^{\gamma-1} = \text{constant}$ (B) $P^{1-\gamma} = T^{\gamma} = \text{constant}$
 (C) $PT^{\gamma-1} = \text{constant}$ (D) $P^{\gamma-1} T^{\gamma} = \text{constant}$
52. Figure shows four $P \rightarrow V$ diagrams, which of these curves represent isothermal and adiabatic processes ?



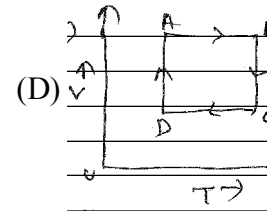
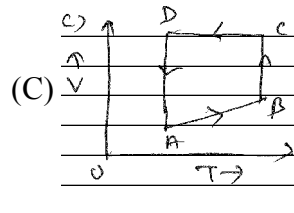
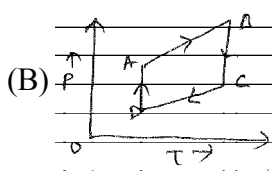
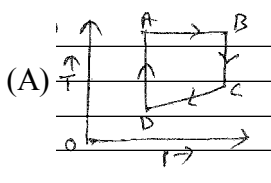
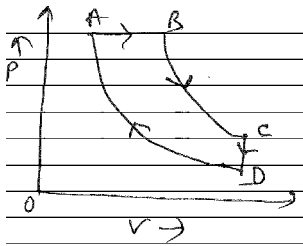
- (A) A and C (B) A and B (C) C and D (D) B and D
53. An ideal gas is taken through cyclic process as shown in the figure. The net work done by the gas is



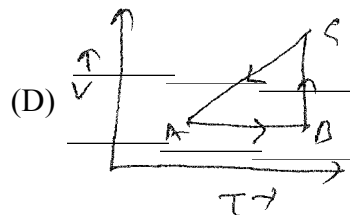
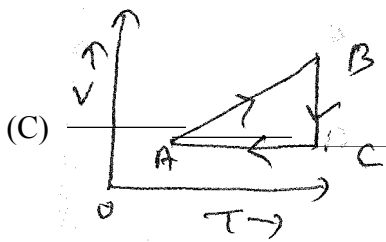
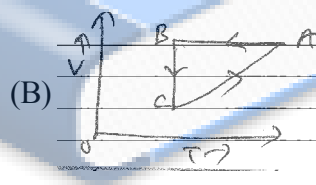
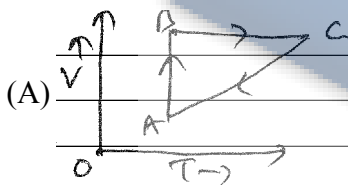
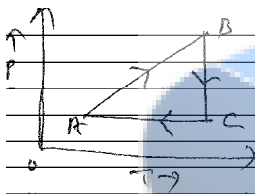
- (A) PV (B) $2PV$ (C) $3PV$ (D) zero
54. μ moles of gas expands from volume V_1 to V_2 at constant temperature T . The work done by the gas is

(A) $\mu RT \left(\frac{V_2}{V_1} \right)$ (B) $\mu RT \ln \left(\frac{V_2}{V_1} \right)$ (C) $\mu RT \left(\frac{V_2}{V_1} - 1 \right)$ (D) $\mu RT \ln \left(\frac{V_2}{V_1} + 1 \right)$

55. A Cyclic Process ABCD is Shown in the $P \rightarrow V$ diagram. which of the following curves represent the same Process ?



56. A Cyclic process is Shown in the $P \rightarrow T$ diagram. Which of the curve show the same process on a $V \rightarrow T$ diagram ?



57. One mole of an ideal gas $\frac{C_p}{C_v} = \gamma$ at absolute temperature T_1 is adiabatically compressed from an initial pressure P_1 to a final pressure P_2 The resulting temperature T_2 of the gas is given by.

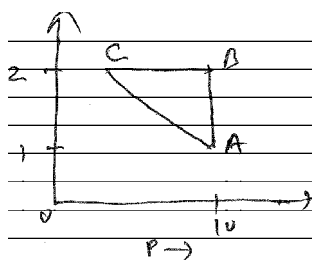
(A) $T_2 = T_1 \left(\frac{p_2}{p_1} \right)^{\frac{\gamma}{\gamma-1}}$

(B) $T_2 = T_1 \left(\frac{p_2}{p_1} \right)^{\gamma}$

(C) $T_2 = T_1 \left(\frac{p_2}{p_1} \right)^{\gamma}$

(D) $T_2 = T_1 \left(\frac{p_2}{p_1} \right)^{\gamma-1}$

58. 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 5J, the work done by the gas in the process $C \rightarrow A$ is



- (A) - 5 J (B) -10 J (C) -15 J (D) - 20 J
59. In an isothermal reversible expansion, if the volume of 96J of oxygen at 27°C is increased from 70 liter to 140 liter, then the work done by the gas will be
- (A) $300 R \log_e^{(2)}$ (B) $81 R \log_e^{(2)}$ (C) $200 R \log_{10}^2$ (D) $100 R \log_{10}^{(2)}$
60. For an isothermal expansion of a Perfect gas, the value of $\frac{\Delta P}{P}$ is equal to
- (A) $-\gamma^2 \frac{\Delta V}{V}$ (B) $-\gamma \frac{\Delta V}{V}$ (C) $-\gamma^2 \frac{\Delta V}{V}$ (D) $\gamma - \frac{\Delta V}{V}$
61. For an adiabatic expansion of a perfect gas, the value of $\frac{\Delta P}{P}$ is equal to
- (A) $-\sqrt{\gamma} \frac{\Delta v}{v}$ (B) $-\frac{\Delta v}{v}$ (C) $-\gamma^2 \frac{\Delta v}{v}$ (D) $-\gamma \frac{\Delta v}{v}$
62. If r denotes the ratio of adiabatic of two specific heats of a gas. Then what is the ratio of slope of an adiabatic and isothermal $P \rightarrow V$ curves at their point of intersection ?
- (A) $\frac{1}{\gamma}$ (B) $\gamma - 1$ (C) γ (D) $\gamma + 1$
63. Work done per mol in an isothermal ? change is
- (A) $RT \log_e \frac{v_2}{v_1}$ (B) $RT \log_{10} \frac{v_2}{v_1}$ (C) $RT \log_{10} \frac{v_1}{v_2}$ (D) $RT \log_e \frac{v_1}{v_2}$
64. The isothermal Bulk modulus of an ideal gas at pressure P is
- (A) vP (B) P (C) $\frac{P}{2}$ (D) $\frac{P}{v}$
65. The isothermal bulk modulus of a perfect gas at a normal Pressure is
- (A) $1.013 \times 10^6 \frac{\text{N}}{\text{m}^2}$ (B) $1.013 \times 10^{-11} \frac{\text{N}}{\text{m}^2}$ (C) $1.013 \times 10^5 \frac{\text{N}}{\text{m}^2}$ (D) $1.013 \times 10^{11} \frac{\text{N}}{\text{m}^2}$

66. An adiabatic Bulk modulus of an ideal gas at Pressure P is
- (A) γP (B) $\frac{P}{\gamma}$ (C) P (D) $\frac{P}{2}$
67. What is an adiabatic Bulk modulus of hydrogen gas at NTP? $\gamma = 1.4$
- (A) $1.4 \frac{N}{M^2}$ (B) $1.4 \times 10^5 \frac{N}{M^2}$ (C) $1 \times 10^{-8} \frac{N}{M^2}$ (D) $1 \times 10^5 \frac{N}{M^2}$
68. If a quantity of heat 1163.4 J is supplied to one mole of nitrogen gas, at room temperature at constant pressure, then the rise in temperature is $R = 8.31 \frac{J}{m.l.k}$
- (A) 28 K (B) 65 K (C) 54 K (D) 40 K
69. One mole of O_2 gas having a Volume equal to 22.4 liter at $0^\circ C$ and 1 atmospheric Pressure is Compressed isothermally so that its volume reduces to 11.2 litres. The work done in this Process is
- (A) 1672.4 J (B) -1728J (C) 1728J (D) -1572.4J
70. The Specific heat of a gas in an isothermal Process is
- (A) zero (B) Negative (C) Infinite (D) Remains
71. A Container that suits the occurrence of an isothermal process should be made of
- (A) Wood (B) Copper (C) glass (D) Cloth
72. A thermodynamic Process in which temperature T of the system remains constant through out Variable P and V may Change is called
- (A) Isothermal Process (B) Isochoric Process
(C) Isobaric Process (D) None of this
73. When 1g of water at $0^\circ C$ and $10^5 \frac{N}{m^2}$ Pressure is Converted into ice of Volume 1.091 cm^3 the external work done be J.
- (A) 0.0182 (B) -0.0091 (C) -0.0182 (D) 0.0091
74. The latent heat of Vaporisation of water is $2240 \frac{J}{g}$ If the work done in the Process of expansion of 1g is 168J. then increase in internal energy is J
- (A) 2072 (B) 2408 (C) 2240 (D) 1904
75. The Volume of an ideal gas is 1 liter and its Pressure is equal to 72 cm of Hg. The Volume of gas is made 900 cm^3 by compressing it isothermally. The pressure of the gas will be Hg column.
- (A) 4 cm (B) 6 cm (C) 7 cm (D) 8 cm

76. In adiabatic expansion
 (A) $\Delta u = 0$ (B) $\Delta u = \text{Positive}$ (C) $J_u = \text{Nagative}$ (D) $\Delta w = 0$
77. 1 mm^3 Of a gas is compressed at 1 atmospheric pressure and temperature 27°C to 627°C What is the final pressure under adiabatic condition. $r = 1.5$
 (A) $80 \times 10^5 \frac{\text{N}}{\text{m}^2}$ (B) $36 \times 10^5 \frac{\text{N}}{\text{m}^2}$ (C) $56 \times 10^5 \frac{\text{N}}{\text{m}^2}$ (D) $27 \times 10^5 \frac{\text{N}}{\text{m}^2}$
78. A monoatomic gas for it $\gamma = \frac{5}{3}$ is suddenly Compressed to $\frac{1}{8}$ of its original volume adiabatically then the final Pressure of gas is times its intial Pressure.
 (A) 8 (B) 32 (C) $\frac{24}{5}$ (D) $\frac{40}{3}$
79. The Pressure and density of a diatomic gas $\gamma = \frac{7}{5}$ Change adiabatically from (P, d) to (P', d') If $\frac{d'}{d} = 32$ then $\frac{P'}{P}$ Should be
 (A) 128 (B) $\frac{1}{128}$ (C) 32 (D) None of this
80. An ideal gas at 27°C is Compressed adiabatically, to $\frac{8}{27}$ of its original Volume. If $\gamma = \frac{5}{3}$, then the rise in temperature is
 (A) 225 k (B) 450 K (C) 375 K (D) 405 K
81. A diatomic gas intially at 18°C is Compressed adiabatically to one eight of its original volume. The temperature after Compression will be
 (A) 10°C (B) 668 K (C) 887°C (D) 144°C
82. Work done by 0.1 mole of a gas at 27°C to double its volume at constant Pressure is _____
 Cal. $R = 2 \frac{\text{Cal}}{\text{mol}^\circ\text{K}}$
 (A) 600 (B) 546 (C) 60 (D) 54
83. A gas expands 0.25m^3 at Constant Pressure $10^3 \frac{\text{N}}{\text{m}^2}$ the work done is
 (A) 250 J (B) 2.5 erg (C) 250 W (D) 250 N
84. In an isochoric Process $T_1 = 27^\circ\text{C}$ and $T_2 = 127^\circ\text{C}$ then $\frac{P_1}{P_2}$ will be equal to
 (A) $\frac{9}{59}$ (B) $\frac{2}{3}$ (C) $\frac{4}{3}$ (D) $\frac{3}{4}$
85. If the temperature of 1 mole of ideal gas is changed from 0°C to 100°C at constant pressure, then work done in the process is J. $R = 8.3 \frac{\text{J}}{\text{molk}}$
 (A) 8.3×10^{-3} (B) 8.3×10^2 (C) 8.3×10^{-2} (D) 8.3×10^3

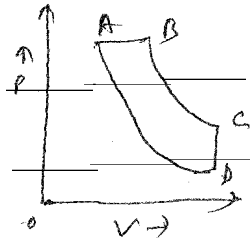
86. A mono atomic gas is supplied the heat Q very slowly keeping the pressure constant The work done by the gas.

- (A) $\frac{2}{5}Q$ (B) $\frac{2}{3}Q$ (C) $\frac{3}{5}Q$ (D) $\frac{1}{5}Q$

87. The Volume of air increases by 5% in an adiabatic expansion. The percentage decrease in its Pressure will be.

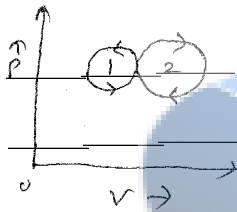
- (A) 5% (B) 6% (C) 7% (D) 8%

88. In $P \rightarrow V$ diagram given below, the isochoric, isothermal and isobaric path respectively are



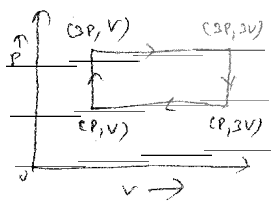
- (A) BA,AD,DC (B) DC,CB,BA (C) AB,BC,CD (D) CA,DA,AB

89. In the following indicators diagram, the net amount of work done will be



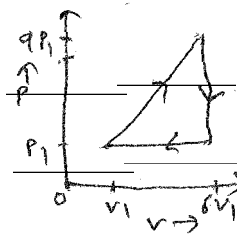
- (A) Negative (B) zero (C) Positive (D) Infinity

90. Work done in the given $P \rightarrow V$ diagram in the cyclic process is



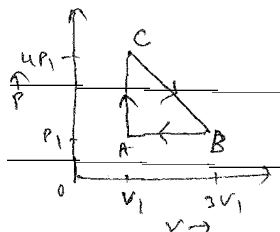
- (A) $4 PV$ (B) $3 PV$ (C) $2 PV$ (D) $\frac{PV}{2}$

91. In the Cyclic Process shown in the figure, the work done by the gas in one cycle

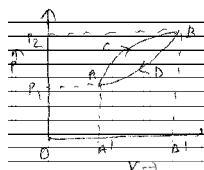


- (A) $40P_1V_1$ (B) $20P_1V_1$ (C) $10P_1V_1$ (D) $5P_1V_1$

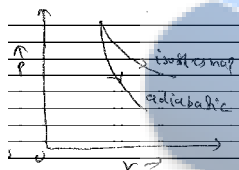
92. An ideal gas is taken V path ACBA as Shown in figure, The net work done in the whole cycle is



- (A) $-3P_1V_1$ (B) Zero (C) $5P_1V_1$ (D) $3P_1V_1$
93. A thermodynamic system is taken from state A to B along ACB and is brought back to A along BDA as Shown in the $P \rightarrow V$ diagram. The net work done during the complete cycle is given by the area.



- (A) $P_1 ACBP_2P_1$ (B) ACBDA (C) $ACBB^1A^1A$ (D) $ADBB^1A^1A$
94. Two identical samples of a gas are allowed to expand (i) isothermally (ii) adiabatically work done is



- (A) more in isothermal process (B) more is an adiabatic process
(C) equal in both process. (D) neither of them
95. What is the relationship Pressure and temperature for an ideal gas under going adiabatic Change.
(A) $PT^\gamma = \text{Const}$ (B) $PT^{-1+\gamma} = \text{Const}$ (C) $P^{1-\gamma}T^\gamma = \text{Const}$ (D) $P^{\gamma-1}T^\gamma = \text{Const}$

96. For adiabatic Process which relation is true mentioned below ? $\gamma = \frac{C_p}{C_v}$
(A) $p^\gamma V = \text{Const}$ (B) $T^\gamma V = \text{Const}$ (C) $TV^\gamma = \text{Const}$ (D) $TV^{\gamma-1} = \text{Const}$

97. For adiabatic Process which one is wrong statement?
(A) $dQ = 0$ (B) entropy is not constant
(C) $du = -dw$ (D) $Q = \text{constant}$

98. Air is filled in a motor tube at 27°C and at a Pressure of a atmosphere. The tube suddenly bursts. Then what is the temperature of air. given r of air = 1.5
(A) 150 K (B) 150°C (C) 75 K (D) 27.5°C

99. If ν is the ratio of Specific heats and R is the universal gas constant, then the molar Specific heat at constant Volume C_v is given by

- (A) $\frac{\nu R}{\nu - 1}$ (B) νR (C) $\frac{R}{\nu - 1}$ (D) $\frac{(\nu - 1)R}{r}$

100. A Carnot engine operating between temperature T_1 and T_2 has efficiency 0.4, when T_2 lowered by 50K, its efficiency increases to 0.5. Then T_1 and T_2 are respectively.
- (A) 300 K and 100 K (B) 400 K and 200 K
(C) 600 K and 400 K (D) 400 K and 300 K
101. A monoatomic gas is used in a Carnot engine as the working substance, If during the adiabatic expansion part of the cycle the volume of the gas increases from V to $8V_1$ the efficiency of the engine is ..
- (A) 60% (B) 50% (C) 75% (D) 25%
102. A System undergoes a Cyclic Process in which it absorbs Q_1 heat and gives out Q_2 heat. The efficiency of the Process is n and the work done is W . Which formula is wrong ?
- (A) $W = Q_1 - Q_2$ (B) $n = \frac{Q_2}{Q_1}$ (C) $n = \frac{W}{Q_1}$ (D) $n = 1 - \frac{Q_2}{Q_1}$
103. A Carnot engine whose sink is at a temperature of 300K has an efficiency of 40% By space should the temperature of the source be increase the efficiency to 60%
- (A) 275 K (B) 325 K (C) 300 K (D) 250 K
104. An ideal gas heat engine is operating between 227°C and 127°C . It absorbs 10^4 J of heat at the higher temperature. The amount of heat converted into work is J.
- (A) 2000 (B) 4000 (C) 5600 (D) 8000
105. Efficiency of a Carnot engine is 50%, when temperature of outlet is 500K. in order to increase efficiency up to 60% keeping temperature of intake the same what is temperature of outlet.
- (A) 200 K (B) 400 K (C) 600 K (D) 800 K
106. A Carnot engine takes 3×10^6 cal of heat from a reservoir at 627°C , and gives to a sink at 27°C . The work done by the engine is
- (A) $4.2 \times 10^6\text{ J}$ (B) $16.8 \times 10^6\text{ J}$ (C) $8.4 \times 10^6\text{ J}$ (D) Zero
107. For which combination of working temperatures the efficiency of Carnot engine is highest.
- (A) 80 K, 60 K (B) 100 K, 80 K (C) 60 K, 40 K (D) 40 K, 20 K
108. An ideal heat engine working between temperature T_1 and T_2 has an efficiency n . The new efficiency if both the source and sink temperature are doubled, will be
- (A) n (B) $2n$ (C) $3n$ (D) $\frac{n}{2}$
109. An ideal refrigerator has a freezer at a temperature of -13°C , The coefficient of performance of the engine is 5. The temperature of the air to which heat is rejected will be.
- (A) 325°C (B) 39°C (C) 325 K (D) 320°C
110. An engine is supposed to operate between two reservoirs at temperature 727°C and 227°C . The maximum possible efficiency of such an engine is
- (A) $\frac{3}{4}$ (B) $\frac{1}{4}$ (C) $\frac{1}{2}$ (D) 1

111. A car not engine Convertsm one sixth of the heat input into work. When the temperature of the sink is reduces by 62°C the efficiency of the engine is doubled. The temperature of the source and sink are
 (A) 80°C , 37°C (B) 95°C , 28°C (C) 90°C , 37°C (D) 99°C , 37°C
112. Car not engine working between 300 K and 600 K has work output of 800J per cycle. What is amount of heat energy supplied to the engine from source per cycle
 (A) $1600 \frac{\text{J}}{\text{cycle}}$ (B) $2000 \frac{\text{J}}{\text{cycle}}$ (C) $1000 \frac{\text{J}}{\text{cycle}}$ (D) $1800 \frac{\text{J}}{\text{cycle}}$
113. What is the value of sink temperature when efficiency of engine is 100%
 (A) 300 K (B) 273 K (C) 0 K (D) 400 K
114. A car not engine having a efficiency of $n = \frac{1}{10}$ as heat engine is used as a refrigerators. if the work done on the system is 10J. What is the amount of energy absorbed from the reservoir at lowes temperature !
 (A) 1 J (B) 90 J (C) 99 J (D) 100 J
115. The temperature of sink of car not engine is 27°C . Efficiency of engine is 25% Then find the temperature of source.
 (A) 227°C (B) 327°C (C) 27°C (D) 127°C
116. The efficiency of car not's engine operating between reservoirs, maintained at temperature 27°C and -123°C is
 (A) 0.5 (B) 0.4 (C) 0.6 (D) 0.25
117. If a heat engine absorbs 50KJ heat from a heat source and has efficiency of 40%, then the heat released by it in heat sink is
 (A) 40 KJ (B) 30 KJ (C) 20 J (D) 20 KJ
118. The efficiency of heat engine is 30% If it gives 30KJ heat to the heat sink, than it should have absorbed KJ heat from heat source.
 (A) 42.8 (B) 39 (C) 29 (D) 9
119. If a heat engine absorbs 2KJ heat from a heat source and release 1.5KJ heat into cold reservoir, then its efficiency is
 (A) 0.5% (B) 75% (C) 25% (D) 50%
120. If the doors of a refrigerators is kept open, then which of the following is true?
 (A) Room is cooled (B) Room is eithers cooled or heated
 (C) Room is neither cooled nor heated (D) Room is heated

Assertion-Reason

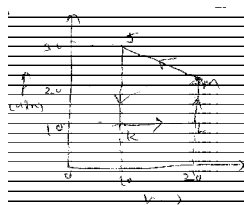
Instructions :-

Read the assertion and reason carefully to mark the correct option out of the options given below.

- (A) If both assertion and reason are true and the reason is the correct explanation of the assertion.
(B) If both assertion and reason are true but reason is not be correct explanation of assertion.
(C) If assertion is true but reason is false.
(D) If the assertion and reason both are false.
121. Assertion : The melting point of ice decreases with increase of Pressure
Reason : Ice contracts on melting.
(A) C (B) B (C) A (D) D
122. Assertion : Fahrenheit is the smallest unit measuring temperature.
Reason : Fahrenheit was the first temperature scale used for measuring temperature.
(A) A (B) C (C) B (D) D
123. Assertion : A beakes is completely, filled with water at 4°C . It will overflow, both when heated or cooled.
Reason : These is expansion of water below 4°C
(A) A (B) B (C) C (D) D
124. Assertion : The total translation kinetic energy of all the molecules of a given mass of an ideal gas is 1-5 times the product of its Pressure and its volume.
Reason : The molecules of a gas collide with each other and velocities of the molecules change due to the collision
(A) D (B) C (C) A (D) B
125. Assertion : The car not is useful in understanding the performace of heat engine
Reason : The car not cycle provides a way of determining the maximum possible efficiency achivable with reservoirs of given temperatures.
(A) A (B) B (C) C (D) D

Match column

126. Heat given to process is positive, match the following column I with the corresponding option of column I₁



Column-i

- (A) JK
(B) KL
(C) LM
(D) MJ

Column-ii

- (p) $\Delta W > 0$
(q) $\Delta Q < 0$
(r) $\Delta W < 0$
(s) $\Delta Q > 0$

- (A) A-p, B-q, C-r, D-s
(B) A-q, B-p, C-s, D-r

- (C) A-r, B-s, C-p, D-q
(D) A-s, B-r, C-q, D-p

127. In Column I different Process is given match corresponding option of column I₁

Column - I

- (A) adiabatic process
(B) Isobaric process
(C) Isochroic process
(D) Isothermal process

Column - II

- (p) $\Delta p = 0$
(a) $\Delta u = 0$
(r) $\Delta Q = 0$
(s) $\Delta W = 0$

- (A) A-p, B-s, C-r, D-q
(B) A-s, B-q, C-p, D-r

- (C) A-r, B-p, C-s, D-q
(D) A-q, B-r, C-q, D-p

Comprehehsion Type

In a container of negligible heat capacity, 200g ice at 0°C and 100g steam at 100°C are added to 200g of water that has temperature 55°C . Assume no heat is lost to the surroundings and the pressure in the container is constant 1 atm.

128. What is the final temperature the System ?
 (A) 72°C (B) 48°C (C) 100°C (D) 94°C
129. At the final temperature, mass of the total water present in the system is
 (A) 493.6g (B) 483.3g (C) 472.6 g (D) 500 g
130. Amount of the Sm left in the system, is equal to
 (A) 16-7 g (B) 8-4 g (C) 12 g (D) 0 g

KEY NOTE

1	B	26	A	51	B	76	C	101	C
2	A	27	C	52	C	77	D	102	B
3	D	28	D	53	A	78	B	103	D
4	D	29	A	54	B	79	A	104	A
5	A	30	D	55	C	80	C	105	B
6	D	31	B	56	D	81	B	106	C
7	C	32	C	57	B	82	C	107	D
8	C	33	D	58	A	83	A	108	A
9	B	34	B	59	C	84	D	109	B
10	A	35	C	60	D	85	B	110	C
11	C	36	A	61	D	86	A	111	D
12	D	37	B	62	C	87	C	112	A
13	A	38	A	63	A	88	D	113	C
14	C	39	C	64	B	89	C	114	B
15	B	40	D	65	C	90	A	115	D
16	D	41	B	66	A	91	B	116	A
17	C	42	A	67	B	92	D	117	B
18	A	43	D	68	D	93	B	118	A
19	B	44	C	69	D	94	A	119	C
20	D	45	C	70	C	95	C	120	D
21	C	46	B	71	B	96	D	121	C
22	D	47	A	72	A	97	B	122	B
23	D	48	D	73	D	98	A	123	A
24	A	49	A	74	A	99	C	124	D
25	B	50	D	75	D	100	D	125	A
								126	B
								127	C
								128	C
								129	B
								130	A

HINT

1. $\frac{\Delta C}{100^\circ} = \frac{\Delta F}{180^\circ} \therefore \frac{25}{100} = \frac{\Delta F}{180}$

2. $T = 273.15 + t^\circ\text{C}$

$0 = 273.15 + t^\circ\text{C}$

3. equal

4. 4°C

$$\frac{c}{5} = \frac{F-32}{9}$$

5. $\frac{c}{5} = \frac{F-32}{9}$

$y = mx + c$ with comparison $m = \frac{5}{9}$

6. $\frac{c}{5} = \frac{f-32}{9}$

7. $\frac{F-32}{9} = \frac{K-273}{5}$

8. $\frac{\Delta T_c}{100} = \frac{\Delta T_F}{180}$

$$\frac{\Delta T_c}{100} \quad \Delta T_c = 40^\circ\text{C}$$

9. $\frac{\Delta T}{T} = \frac{1}{2} 2\Delta\theta = \frac{1}{2} \times 2 \times 10^{-6} \times 10 = 10^{-5}$

$$\% \text{ in increase} = \frac{\Delta T}{T} \times 100 = 10^{-5} \times 100 = 1 \times 10^{-3} \%$$

10. $W = p\Delta V$

11. $C_p = \left(\frac{f}{2} + 1\right)R, C_v = \frac{f}{2}R$

$$\frac{C_p}{C_v} = \frac{\left(\frac{f}{2} + 1\right)R}{\frac{f}{2}R} = \frac{\frac{f}{2} + 1}{\frac{f}{2}} = \frac{f+2}{f} = 1 + \frac{2}{f}$$



12. $W = \text{area inside the closed curve}$

$$= \pi \frac{1}{2} (p_2 - p_1) \times \frac{1}{2} (V_2 - V_1) \quad [\text{Treat Circle as an ellipse}]$$

13.
$$\Delta u = \frac{P(V_2 - V_1)}{\gamma - 1} = \frac{PV}{\gamma - 1}$$

14.
$$u \propto T \quad \frac{u_1}{u_2} = \frac{T_1}{T_2} \quad \therefore \frac{u_1 - u_2}{u_2} = \frac{T_1 - T_2}{T_2}$$

15. For all gases $C_p - C_v = R$

16.
$$\Delta Q = \Delta u + \Delta w$$

17.
$$\frac{C_p}{C_v} = \gamma = 1 + \frac{2}{f}$$

18. $W = \text{area enclosed by } P \rightarrow V \text{ graph}$

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

19. $W = \text{area under the } P \rightarrow V \text{ curve}$

Smallest for curve 1, largest for curve 3

20. monoatomic gas $C_v = \frac{3}{2}R$, diatomic gas $C_v = \frac{5}{2}R$,

$$\text{one mole of each gas is mixed } C_v(\text{mix}) = \frac{1}{2} \left[\frac{3}{2}R + \frac{5}{2}R \right] = 2R$$

$$C_p(\text{mix}) = R + C_v \\ = R + 2R = 3R$$

$$r = \frac{C_p(\text{mix})}{C_v(\text{mix})} = \frac{3R}{2R} = 1.5$$

21. $dQ = du + dw$

22.
$$C_v(\text{mix}) = \frac{n_1 C_{v1} + n_2 C_{v2}}{n_1 + n_2}$$

23.
$$\Delta U = \gamma C_v \Delta T$$

24.
$$u = \frac{nfRt}{2}$$

n = number of mole of the gas

f = number of degree of freedom

$$U_{\text{Total}} = U_{O_2} + U_{AB}$$

25. TV = [an adiabatic process]

$$T_1 V_1 = T_2 V_2$$

$$\frac{T_1}{T_2} = \left(\frac{V_2}{V_1} \right)$$

26. W = area under the curve and volume axis on the P → V diagram.

27. dQ = du + dw

28. energy required in heating water = ms Δθ

from coil, $\frac{\text{energy available}}{S} = \text{Power of coil} - \text{Power lost}$

For 840 J time required = 1S

$$\therefore 4.2 \times 10^4 J = ?$$

$$t = \frac{4.2 \times 10^5}{840} 500 J = 8.33 \text{ min} = 8 \text{ min } 20 \text{ s}$$

29. $C_p = \frac{(\Delta Q) P}{r \Delta T}$

$$c_v = c_p - R$$

$$= 7 - 2 = 5 \frac{\text{cal}}{\text{mol K}}$$

30. $\frac{\Delta u}{\Delta Q} = \frac{1}{\gamma} = \frac{5}{7}$

$$\Delta u = n C_v \Delta T$$

$$\Delta Q = n C_p \Delta T$$

$$\frac{\Delta u}{\Delta Q} = \frac{C_v}{C_p} = \frac{1}{\gamma} = \frac{5}{7}$$

31. For A (ΔQ) P = r C_p (ΔT) A (Isobaric process)

For B (ΔQ) V = C_v (ΔT) B (Isochroic process)

$$(\Delta Q)_p = (\Delta Q)_v$$

32. Decrease in K.E = increase in I.E $\frac{1}{2} m v^2 = \gamma C_v \Delta T$

33. Rate of heat produced = (Viscous force F) × (Velocity V)

$$\frac{dQ}{dt} = 6\pi n r v^2 \left[\frac{2}{g} \frac{(\rho - \rho_0) r^2 g}{n} \right]^2$$

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

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

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

$\Delta u = \Delta u - 62.25$ (adiabatic process)

$u = 62.25 \times 10^3 \text{ J}$

$$C_v = \frac{\Delta u}{n\Delta t}$$

for monoatomic gas $C_v = \frac{3}{2}R = \frac{3}{2} \times 8 - 3 = 12.45 \frac{\text{J}}{\text{mol K}}$

40. If CP_1 and CV_1 is a molar specific heat $CP_1 - CV_1 = R$ ----- (i)

$C_p = \frac{C_{p1}}{32}, C_v = \frac{C_{v1}}{32}$ [molar mass of $O_2 = 32$]

From equation (i) $32C_p - 32C_v = R$

$$C_p - C_v = \frac{R}{32}$$

41. (i) $Q = w + \Delta u$

(ii) $Q = W + \Delta u$ (Δu equal)

42. $u = \frac{5}{2} \ell RT$

$PV = \ell RT$ [diatomic gas]

$$u = \frac{5}{2} PV$$

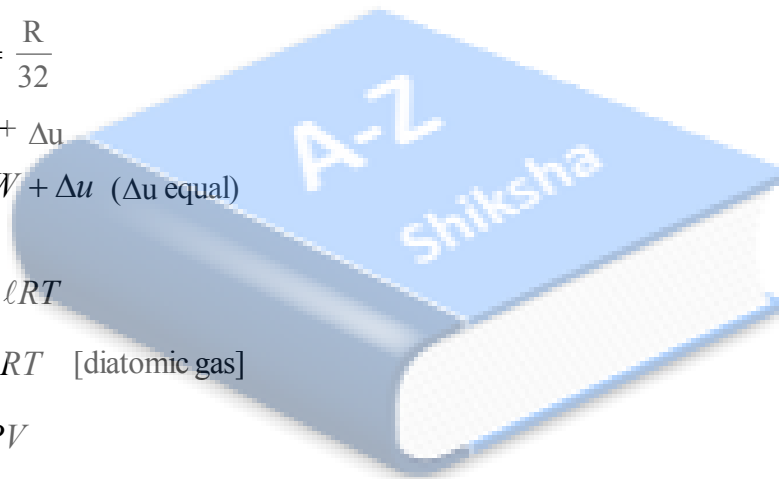
$$V = \frac{\text{mass}}{\text{density}} = \frac{1}{5} \text{ m}^3$$

43. $\Delta u = mc\Delta T$

44. given $P \propto T^5$

an adiabatic process $P \propto T^{\frac{\phi}{\phi-1}}$

45. $\therefore \frac{\phi}{\phi-1} = 5$



46. $W = P_{atm} V$

$$= P_{atm} \frac{V}{2} \quad (i)$$

Initially $P_{atm} V = nRT \therefore V = \frac{nRT}{P_{at}}$

49. (i) Volume Constant $\therefore W = P\Delta V = 0$

(ii) Pressure Constant $W = P \int_v^{2v} dv = PV$

50. For free expansion $Q = W = 0$

$T = \text{const} \quad \mu \therefore \Delta E_{int} = 0$

51. $PV^\gamma = \text{Const}$ (an adiabatic process)

$$PV = nRt \therefore V = \frac{RT}{P}$$

52. For both slope is negative & Slope of adiabatic curve is more

53. $W = \text{area of } P \rightarrow V \text{ diagram}$

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

54. AB \rightarrow Constant P, increasing V, increasing T

BC \rightarrow Constant T, increasing V, decreasing P

CD \rightarrow Constant V, decreasing P, decreasing T

DA \rightarrow Constant T, decreasing V, increasing P

Also BC is at higher temperature than AD

57. $PV^\gamma = \text{const}$ [adiabatic process]

ideal gas $PV = RT$ ($m = 1$) $\therefore V = \frac{RT}{P}$

$$\therefore P \left(\frac{RT}{P} \right)^\gamma = \text{const} \quad \frac{T^\gamma}{P^{(\gamma-1)}} = \text{const}$$

58. $Q = W_{AB} + W_{BC} + W_{CA}$

$$5 = 10 + 0 + W_{CA}$$

59. $W = RT \log_e \frac{V_2}{V_1} = 2.3 \left(\frac{M}{M_0} \right) RT \log_{10} \frac{V_2}{V_1}$

60. $PV = \text{constant}$ (isothermal Process)

$$P\Delta V - V\Delta P = 0 \quad \therefore \frac{\Delta P}{P} = \frac{-\Delta V}{V}$$

61. $PV^\gamma = \text{Constant}$ (an adiabatic Process)

$$Pr v^{\gamma-1} \Delta V + V^\gamma \Delta P = 0$$

$$62. \frac{\left(\frac{\Delta P}{P}\right)_{\text{adiabatic}}}{\left(\frac{\Delta P}{P}\right)_{\text{isothermal}}} = \frac{-\gamma \frac{\Delta V}{V}}{-\frac{\Delta V}{V}} = \gamma$$

$$63. W = mRT \ln \frac{V_2}{V_1} = R T \log_e \frac{V_2}{V_1}$$

64. isothermal bulk modulus $B = P$

$$64. \text{ isothermal bulk modulus } B = P = 1.013 \times 10^5 \frac{\text{N}}{\text{m}^2}$$

66. an adiabatic bulk modulus $B = \gamma P$

$$67. \text{ an adiabatic bulk modulus } B = 1.4 \times 1 \times 10^5 = 1.4 \times 10^5 \frac{\text{N}}{\text{m}^2}$$

$$68. dQ = \mu C_p dT$$

$$69. W = -\mu RT \log e \text{ (isothermally compressed)}$$

$$70. C_{\text{iso}} = \frac{\Delta Q}{m\Delta T} = \infty \text{ [} \Delta T = 0, \text{ isothermal process]}$$

71. For isothermal process base is conductivity

$$73. W = P \Delta V \text{ (isothermal process)}$$

$$74. \Delta u = \Delta Q - \Delta w$$

$$74. P_1 V_2 = P_2 V_2 \text{ (isothermal process)}$$

$$76. \Delta Q = \Delta u + \Delta W$$

$$0 = \Delta u + \Delta W$$

$$77. \frac{T^\gamma}{P^{\gamma-1}} = \text{Constant [adiabatic change]}$$

$$\left(\frac{P_2}{P_1}\right)^{\frac{1}{2}} = \left(\frac{T_2}{T_1}\right)^{\frac{3}{2}}$$

$$\left(\frac{P_2}{10^5}\right)^{\frac{1}{2}} = \left(\frac{900}{300}\right)^{\frac{3}{2}}$$

78. $PV^{\gamma} = \text{Constant}$ (adiabatic compressed)

$$\frac{P_2}{P_1} = \left(\frac{V_1}{V_2} \right)^{\gamma}$$

79. $PV^{\gamma} = \text{Constant}$ (adiabatic process)

$$80. \frac{T_2}{T_1} = \left(\frac{V_1}{V_2} \right)^{\gamma-1}$$

81. $TV^{\gamma-1} = \text{Constant}$

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

$$82. \frac{V_1}{V_2} = \frac{T_1}{T_2} \quad \therefore \frac{V}{2V} = \frac{300}{T_2} \quad \therefore T_2 = 600\text{K}$$

$$W = P \Delta V$$

$$83. W = P \Delta V = 10^3 \times 0.25 = 250 \text{ J}$$

84. $P \propto T$ (constant volume)

$$\frac{P_1}{P_2} = \frac{T_1}{T_2}$$

$$85. W = P \Delta V$$

$$86. \Delta W = (\Delta Q)_P - \Delta u$$

$$= (\Delta Q)_P - (\Delta Q)_V$$

$$= (\Delta Q)_P - \left[1 - \frac{(\Delta Q)_V}{(\Delta Q)_P} \right]$$

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

$$87. \frac{dp}{p} \times 100 = -\gamma \frac{dv}{V} \times 100 \quad (\text{adiabatic expansion})$$

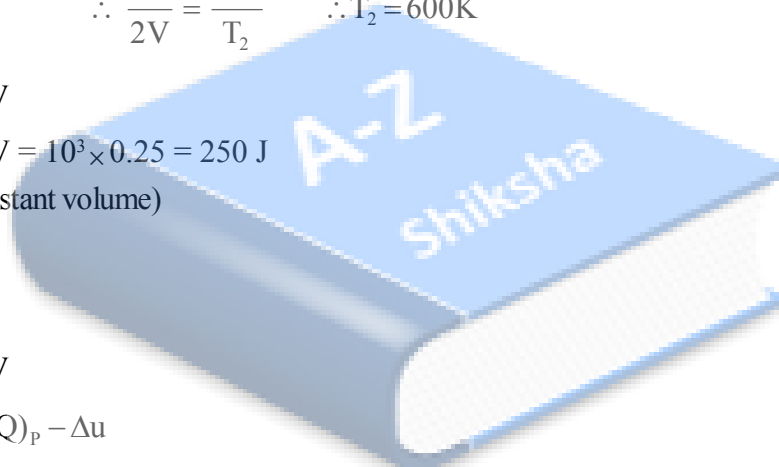
89. cyclic process 1 negative work

 2 net positive

90. $W \rightarrow$ area of closed $P \rightarrow V$ diagram

$$= (3V - V)(3P - P) = 4PV$$

91. $W =$ area under curve



92. $W = \text{area under curve}$
 94. $\therefore (\text{Area})_{\text{iso}} > (\text{Area})_{\text{adi}} \quad W_{\text{iso}} > w_{\text{adi}}$
 95. $PV^\gamma = \text{Con}$
 96. $PV^\gamma = \text{Con}$
 97. $\Delta Q = 0, Q = \text{const}, du = -dw$ (adiabatic process)

$$98. \frac{T_2}{T_1} = \left(\frac{P_2}{P_1} \right)^{\frac{\gamma-1}{\gamma}}$$

$$99. \frac{CP}{CV} = \gamma \quad \therefore CP = \gamma CV$$

$$\text{but } CP - CV = R$$

$$100. n = 1 - \frac{T_2}{T_1} \quad 100.$$

$$101. T_1 V_1^{\gamma-1} = T_2 V_2^{\gamma-1} \quad (\text{adiabatic process})$$

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

$$103. n = 1 - \frac{T_2}{T_1}$$

$$104. n = 1 - \frac{T_2}{T_1} = \frac{1-400}{500} = \frac{1}{5}$$

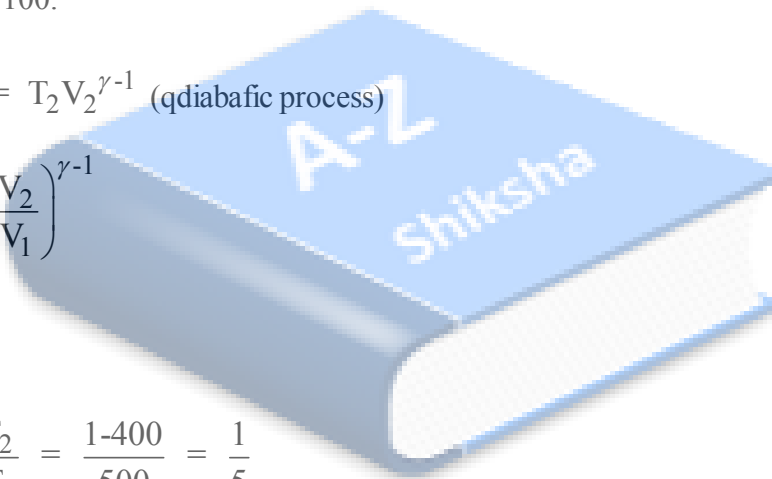
$$W = n Q_1$$

$$105. n = 1 - \frac{T_2}{T_1}, \frac{T_2}{T_1} \text{ Should be minimum}$$

$$106. (i) n = 1 - \frac{T_2}{T_1}$$

$$(ii) n^1 = 1 - \frac{2T_2}{2T_1} = 1 - \frac{T_2}{T_1} = n$$

$$109. \alpha = \frac{T_2}{T_1 - T_2}$$



$$110. \quad n = 1 - \frac{T_2}{T_1} = 1 - \frac{500}{1000} = \frac{1}{2}$$

$$111. \quad (i) \quad n = 1 - \frac{T_2}{T_1} = \frac{W}{Q_1} = \frac{1}{6} \quad \therefore \quad n = \frac{1}{6} \quad - (1)$$

$$(ii) \quad n^1 = 1 - \frac{T_2 - 62}{T_1}$$

$$= 1 - \frac{T_2}{T_1} + \frac{62}{T_1}$$

$$= n + \frac{62}{T_1} \quad - (2)$$

$$\text{Now, } n^1 = 2n$$

$$112. \quad n = 1 - \frac{T_2}{T_1} = \frac{W}{Q_1} \quad \therefore \quad Q = \left(\frac{T_1}{T_1 - T_2} \right)^w$$

$$113. \quad n = 1 - \frac{T_2}{T_1}$$

$$114. \quad n = 1 - \frac{T_2}{T_1}$$

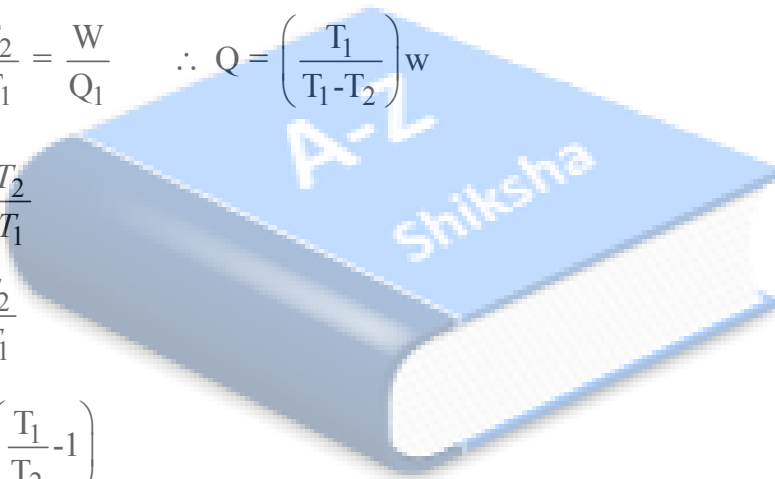
$$W = Q_2 \left(\frac{T_1}{T_2} - 1 \right)$$

$$115. \quad n = 1 - \frac{T_2}{T_1}$$

$$116. \quad n = 1 - \frac{T_2}{T_1}$$

$$117. \quad n = 1 - \frac{Q_2}{Q_1}$$

$$118. \quad n = 1 - \frac{Q_2}{Q_1}$$



$$119. \quad n = 1 - \frac{Q_2}{Q_1}$$

121. with rise in pressure melting point of ice decreases. Also ice contracts on melting.

122. celcius scale was the first temperature scale and Fahrenheit is the smallest unit measuring.

123. Water has maximum density at 4°C on heating above 4°C or cooling below 4°C density of water decreases and its volume increases, therefore, water overflows in the both cases.

$$124. \quad \frac{1}{2} m (v^2) = \frac{3}{2} RT$$

125. car not cycle has maximum efficiency

126. (a) isochoric process $\Delta w = 0 \quad \therefore \Delta Q = \Delta u$

$P \propto T$ P decrease, T also decrease $\therefore \Delta u$ negative $\therefore \Delta Q < 0$

(b) isobasic process, volume increase $\therefore \Delta W > 0$

(c) isochoric process $\Delta W = 0 \quad \Delta Q = \Delta u$

$P \propto T$, P increase $\therefore T$ increase $\therefore \Delta Q > 0$

(d) Volume decrease $\Delta W < 0$

127. adiabatic process $\Delta Q = 0$

Isobasic process $P = \text{const} \quad \therefore \Delta P = 0$

Isochroic process $V = \text{const} \quad \therefore \Delta W = 0$

Isothermal process $T = \text{const} \quad \therefore \Delta u = 0$

128 to 130.

head rewuired by ice and water to go up

to $100^\circ\text{C} = m_1 L + m_1 s_w \Delta T + m_w s_w \Delta T$

$= 200 \times 80 + 200 \times 1 \times 100 + 200 \times 1 \times 45$

$= 16,000 + 20,000 + 9,000$

$= 45,000 \text{ cal}$

$=$ give by m_s mass of steam

$= m_s L$

$m_s = \frac{45,000}{540} = 83.3 \text{ g}$ convert into waters of 100°C

Total water $= 200 + 200 + 83.3$

$= 483.3 \text{ g}$

steam left $= 100 - 83.3 = 16.79$