



**MADE EASY**

India's Best Institute for IES, GATE & PSUs

**Test Centres:** Delhi, Noida, Hyderabad, Bhopal, Jaipur, Lucknow, Bhubaneswar, Indore, Pune, Kolkata, Patna

**UPPSC AE 2019**  
ASSISTANT ENGINEER

**MECHANICAL  
ENGINEERING**

**Test 2**

**Part Syllabus Test**

Thermodynamics

**ANSWER KEY**

1. (d)	11. (b)	21. (a)	31. (a)	41. (b)
2. (c)	12. (b)	22. (b)	32. (c)	42. (b)
3. (c)	13. (a)	23. (c)	33. (a)	43. (c)
4. (b)	14. (b)	24. (c)	34. (b)	44. (c)
5. (c)	15. (d)	25. (b)	35. (c)	45. (b)
6. (b)	16. (c)	26. (c)	36. (c)	46. (a)
7. (c)	17. (a)	27. (b)	37. (b)	47. (c)
8. (b)	18. (d)	28. (b)	38. (c)	48. (c)
9. (c)	19. (b)	29. (d)	39. (a)	49. (a)
10. (a)	20. (c)	30. (d)	40. (a)	50. (c)

## DETAILED EXPLANATIONS

1. (d)

For ideal gas  $z = 1$   
 For a non ideal gas  $z < 1, z > 1$   
 but  $z \neq 1$

3. (c)

According to 1st law of thermodynamics,

$$\oint \delta Q = \oint \delta W = Q_{1-2} + Q_{2-1} = W_{1-2} + W_{2-1}$$

$$20 - 10 = 40 + W_{2-1}$$

$$10 = 40 + W_{2-1} \Rightarrow W_{2-1} = -30 \text{ kJ}$$

4. (b)

Net heat intraction,  $\oint \delta Q = \oint \delta W = \text{Area enclosed in } p-v \text{ diagram}$

$$= \pi R^2 = \pi (1) \text{ m}^3 (1) \text{ MPa}$$

$$= \pi \times 1 \text{ m}^3 \times 1000 \text{ kPa} = 3141 \text{ kJ}$$

8. (b)

$$(dS)_{\text{system}} = mc_p \ln\left(\frac{T_2}{T_1}\right) = 4 \times 1 \times \ln\left(\frac{400}{800}\right) = -2.7726 \text{ kJ/K}$$

$$(dS)_{\text{surrounding}} = \frac{mc_p(T_1 - T_2)}{T_o} = \frac{4 \times 1 \times 400}{300} = 5.33 \text{ kJ/K}$$

$$(dS)_{\text{universe}} = -2.7726 + 5.33 = 2.5601 \text{ kJ/kg}$$

$$\text{Unavailable energy} = T_o dS = 300 \times 2.5601 = 768.22 \text{ kJ}$$

$$\text{Heat transferred} = mc dT = 4 \times 1 \times (800 - 400) = 1600 \text{ kJ}$$

$$\text{Available energy} = 1600 - 768.22 = 831.78 \text{ kJ}$$

19. (b)

For maximum work, heat engine must be reversible,

$$(dS)_{\text{HE}} = 0$$

$$(dS)_1 + (dS)_2 = 0$$

Let  $m, c_p$  are the mass and specific heat of body respectively.

$$mc_p \ln\left(\frac{T_f}{T_1}\right) + mc_p \ln\left(\frac{T_f}{T_2}\right) = 0$$

$$\Rightarrow \ln\left(\frac{T_f^2}{T_1 \times T_2}\right) = 0$$

$$\Rightarrow \frac{T_f^2}{T_1 \times T_2} = 1$$

$$\Rightarrow T_f = \sqrt{T_1 T_2}$$

23. (c)

On applying steady flow energy equation for a nozzle,

$$h_1 + \frac{V_1^2}{2000} = h_2 + \frac{V_2^2}{2000} \quad (W = 0, Q = 0, \Delta PE = 0)$$

$$\begin{aligned} V_2 &= \sqrt{2000(h_1 - h_2)} = \sqrt{2000 \times 45} \\ &= \sqrt{90000} = 300 \text{ m/s} \end{aligned}$$

24. (c)

$$\begin{aligned} (\text{COP})_{\text{HP}} &= 1 + (\text{COP})_{\text{RE}} \\ &= 1 + 3.4 = 4.4 \end{aligned}$$

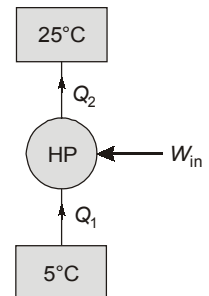
25. (b)

Given:  $m = 0.2 \text{ kg}$ ,  $c_w = 4.2 \text{ kJ/kgK}$ , LH of vaporization =  $2256 \text{ kJ/kg}$ 

$$\text{Rate of heat transfer to water, } \dot{Q} = \frac{m \times LH}{t} = \frac{0.2 \times 2256}{10} = 45.12 \text{ kJ/min}$$

28. (b)

$$\begin{aligned} \dot{Q}_2 &= 18000 \text{ kJ/hr} = \frac{18000}{3600} = 5 \text{ kW} \\ \dot{W}_{\text{in}} &= 1.9 \text{ kW} \\ (\text{COP})_{\text{HP}} &= \frac{\dot{Q}_2}{\dot{W}_{\text{in}}} = \frac{5}{1.9} = 2.63 \end{aligned}$$



30. (d)



$$\text{On balancing hydrogen, } 2b = 14 \Rightarrow b = 7$$

$$\text{On balancing carbon, } a = 6$$

$$\begin{aligned} \text{On balancing oxygen, } 0.21 \times 2x &= 2a + b \\ &= 2 \times 6 + 7 = 19 \end{aligned}$$

$$x = \frac{19}{0.42} = 45.238 \text{ mole}$$

$$\begin{aligned} \text{On balancing nitrogen, } 0.79 \times 2x &= 2c \\ c &= 3.5738 \end{aligned}$$

$$\text{So, Mole of fraction of CO}_2, y_{\text{CO}_2} = \frac{a}{a + b + c} = \frac{6}{6 + 7 + 3.5738} = 0.362 = 36.2\%$$

31. (a)

Intensive properties are independent of mass.

32. (c)

$$\begin{aligned}
 \text{Let} & R = R_0(1 + \alpha t) \\
 \text{as } R = 2.5, \text{ at } t = 0 & 2.5 = R_0(1 + \alpha \times 0) \\
 \Rightarrow & R_0 = 2.5 \\
 \text{Again, at } t = 100^\circ\text{C}, R = 5, & 5 = 2.5(1 + \alpha \times 100) \\
 \Rightarrow & \alpha = 0.01 \\
 \text{So when, } R = 9\Omega, & 9 = 2.5(1 + 0.01 \times t) \\
 \Rightarrow & t = 260^\circ\text{C}
 \end{aligned}$$

33. (a)

$$\begin{aligned}
 \text{As} & Tds = dh - vdP \\
 \text{So for constant pressure lines } dP = 0 & \\
 \Rightarrow & \left(\frac{dh}{ds}\right)_p = T = \text{Slope} = \text{always positive as 'T' is always positive for superheated region.}
 \end{aligned}$$

34. (b)

Let methane is burned with  $x$  moles of air,  
 $\text{CH}_4 + x[0.21 \text{ O}_2 + 0.79\text{N}_2] \rightarrow a\text{CO}_2 + b\text{H}_2\text{O} + c\text{O}_2 + d\text{N}_2$   
 $\therefore c = 1$

$$\begin{aligned}
 \text{On balancing hydrogen,} & b = 2 \\
 \text{On balancing carbon,} & a = 1 \\
 \text{On balancing oxygen,} & 0.21 \times 2x = 2a + b + 2 \\
 & = 2 \times 1 + 2 + 2 = 6 \\
 & x = \frac{6}{0.42} = 14.2857
 \end{aligned}$$

$$\text{So, } \frac{\dot{m}_{\text{air}}}{\dot{m}_{\text{fuel}}} = \frac{14.2857 \times 29}{1 \times (12 + 4 \times 1)} = 25.89$$

36. (c)

An isolated system is either of the following:

1. a physical system so far removed from other systems that it does not interact with them.
2. a thermodynamic system enclosed by rigid immovable walls through which neither matter nor energy can pass.

38. (c)

Given:  $c_p = 1 \text{ kJ/kgK}$ ,  $c_v = 0.75 \text{ kJ/kgK}$ ,  $T = 27^\circ\text{C} = (27 + 273) \text{ K} = 300 \text{ K}$ ,  $p = 1 \text{ bar} = 100 \text{ kPa}$

Gas constant:  $R = c_p - c_v = 1 - 0.75 = 0.25 \text{ kJ/kgK}$

Applying equation of state in term of density,

$$\begin{aligned}
 p & = \rho R T \\
 100 & = \rho \times 0.25 \times 300 \\
 1 & = 0.75 \rho
 \end{aligned}$$

$$\text{or } \rho = \frac{1}{0.75} = \frac{10}{75} = 1.33 \text{ kg/m}^3$$

39. (a)

We know that,

$$F = 1.8C + 32 \quad (F = \text{Temp. in Fahrenheit, } C = \text{Temp. in Celsius})$$

If

$$F = 2C$$

∴

$$1.8C + 32 = 2C$$

or

$$0.2C = 32$$

or

$$C = \frac{32}{0.2} = 160^\circ\text{C}$$

41. (b)

1st law of thermodynamic for process,

$$\delta Q = dU + \delta W$$

For adiabatic process,

$$\delta W = -dU$$

Adiabatic work is equal to change in internal energy. The internal energy is point function. Thus, adiabatic is also point function.

42. (b)

For an ideal gas,

$$s_2 - s_1 = c_v \ln \frac{T_2}{T_1} + R \ln \frac{V_2}{V_1}$$

$$\frac{V_2}{V_1} = \left( \frac{T_1}{T_2} \right)^{\frac{1}{n-1}} \quad (\text{for polytropic process})$$

$$s_2 - s_1 = c_v \ln \frac{T_2}{T_1} + R \ln \left( \frac{T_1}{T_2} \right)^{\frac{1}{n-1}}$$

$$s_2 - s_1 = \left( \frac{R}{\gamma-1} - \frac{R}{n-1} \right) \ln \frac{T_2}{T_1} \quad \left( c_v = \frac{R}{\gamma-1} \right)$$

$$= \left( \frac{R}{\gamma-1} - \frac{R}{n-1} \right) R \ln \frac{T_2}{T_1} = \left\{ \frac{n-1-\gamma+1}{(\gamma-1)(n-1)} \right\} R \ln \frac{T_2}{T_1}$$

$$s_2 - s_1 = \frac{(n-\gamma)R}{(\gamma-1)(n-1)} \ln \frac{T_2}{T_1} = \frac{(n-\gamma)R}{(\gamma-1)(n-1)} \times \ln \frac{T_2}{T_1}$$

$$(S_2 - S_1) = m(s_2 - s_1) = \left( \frac{n-\gamma}{n-1} \right) m c_v \log_e \frac{T_2}{T_1} \quad \left( c_v = \frac{R}{\gamma-1} \right)$$

44. (c)

Given data:

$$V_1 = 1 \text{ m}^3$$

$$p_1 = 5 \text{ bar} = 5 \times 10^5 \text{ Pa}$$

$$V_2 = 2 \text{ m}^3$$

$$p_2 = 5 \text{ bar} = 5 \times 10^5 \text{ Pa}$$

Paddle work done on the system,

$$W_{\text{Paddle}} = 200000 \text{ Nm} = 2 \times 10^5 \text{ Nm}$$

Work done by the system,

$$W_{1-2} = p(V_2 - V_1) = 5 \times 10^5(2 - 1) = 5 \times 10^5 \text{ Nm}$$

Net work done by the system,

$$W_{\text{net}} = W_{1-2} - W_{\text{Paddle}} = 5 \times 10^5 - 2 \times 10^5 = 3 \times 10^5 \text{ Nm}$$

46. (a)

At initial state

Given:  $m_1 = 10 \text{ kg}$ ,  $T_1 = 300 \text{ K}$ ,  $V_1 = 1 \text{ m}^3$

At final state

Given:  $m_2 = ?$ ,  $T_2 = 500 \text{ K}$ ,  $P_2 = P_1$ ,  $V_2 = V_1 = 1 \text{ m}^3$  ( $\because$  Rigid tank)

$$P_1 V_1 = m_1 R T_1$$

$$P_2 V_2 = m_2 R T_2 \quad (P_2 = P_1 \text{ and } V_1 = V_2)$$

$$m_1 T_1 = m_2 T_2$$

$$10 \times 300 = m_2 \times 500$$

or

$$m_2 = 6 \text{ kg}$$

$$\text{Mass of air escaped} = m_1 - m_2 = 10 - 6 = 4 \text{ kg}$$

47. (c)

$$\text{Thermal reservoir, } ds = \frac{Q}{T}$$

48. (c)

Heat required to convert 1 kg of ice to 1 kg of water = L.H. = 335 kJ/kg

Heat given by water at 25°C =  $mc_w \times \Delta T = 1 \times 4.2 \times 25 = 105 \text{ kJ/kg} < 335$

Hence equilibrium temperature will be 0°C, as at equilibrium mixture of ice and water will be present.

49. (a)

$$T_1 = (327 + 273)\text{K} = 600 \text{ K}$$

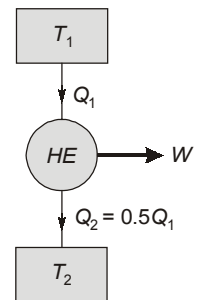
$$T_2 = (27 + 273)\text{K} = 300 \text{ K}$$

Carnot efficiency:

$$\begin{aligned} \eta_{\text{Carnot}} &= 1 - \frac{T_2}{T_1} \\ &= 1 - \frac{300}{600} = 1 - 0.5 = 0.5 = 50\% \end{aligned}$$

Actual efficiency:

$$\eta = 1 - \frac{Q_2}{Q_1} = 1 - \frac{0.5Q_1}{Q_1} = 1 - 0.5 = 0.5 = 50\% = \eta_{\text{Carnot}}$$



○○○○