ROLL NO: MS NAME:

PHY202 Jan-Aug 2019: Midsem 1 Dated February 4, 2019: Inst: Dipanjan Chakraborty

- Time: 60 minutes
- Max Marks: 40
- Attempt all questions. No aids (Books/Notes/Gadgets).

Question	1	2	3	4	T
Marks					
					5

1. Each of the following question has 1 marks for the correct tick mark and the rest for justification.

(a): The differential $dz = 2xydx + (x^2 + 2y)dy$ is not an exact differential

YES NQ

Justification

$$\frac{\partial A}{\partial Y} = 2x \qquad \frac{\partial B}{\partial x} = 2x \qquad -0$$

$$dz = d(x^2y + y^2) \qquad -C$$

(b): For an ideal gas, the internal energy per unit volume is $P/(\gamma - 1)$, where P is the pressure and γ is the ratio of specific heats.

Justification

$$U = \frac{3}{2} nRT$$

$$Q = \frac{5}{2} nR$$

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(c): For n moles of an ideal gas, initially confined in volume V and at temperature T is expanded to a volume αV (i) by a reversible isothermal expansion (ii) removing a partition and allowing a free expansion. The change in entropy is the same in the two cases.

Justification

Entropy is a state function. Change in]
Entropy is
$$\Delta S = nR \ln \alpha$$
.

(b): An ideal monoatomic gas in three dimensions is initially at a temperature T. The temperature is changed to 4T. The root-mean square velocity $v_{\rm rms} = \sqrt{\langle \vec{v}^2 \rangle}$ changes by a factor 3.

Justification

$$V_{rms}^{\perp} = \sqrt{\langle \vec{V}^2 \rangle} = \sqrt{\frac{3 \, \text{kgT}}{m}} \quad \text{(i)} \quad V_{rms}^2 = \sqrt{\frac{12 \, \text{kgT}}{m}} = 2 \sqrt{\frac{3 \, \text{kgT}}{m}} \quad \text{(i)}$$

$$\frac{V_{\text{rms}}^2}{V_{\text{rms}}^4} = 2$$

Ideal monoatomic Gas. & Specific heats constant?

2. Consider a hydrostatic system with fixed particle number. The system undergoes a change in state going from a state-1 (P_1, V_1) to a state-2 (P_2, V_2) by an isothermal expansion. Now consider another process in which we first go from (P_1, V_1) to (P_2, V_1) and then to (P_2, V_2) . Estimate the entropy change in going from state 1 to state 2 along each of these paths. What do you conclude?

[Marks=12]

For isothermal expansion for
$$(P_1,V_1)$$
 to (P_2,V_2)

$$\Delta S = \pi R \int \frac{dV}{V} = \pi R \ln V_2/V_1 - \mathbb{C}$$
Change in entropy T' (P_1,V_1) to (P_2,V_1)

$$\Delta S = Gp \int \frac{dT}{T} = Gp \ln \frac{T'}{T_p} - \mathbb{C}$$
Using in entropy for (P_2,V_1) to (P_2,V_2)

$$\Delta S = Gp \int \frac{dT}{T} = Gp \ln \frac{T_2}{T_1} - \mathbb{C}$$

$$\Delta S = Gp \int \frac{dT}{T} = Gp \ln \frac{T_2}{T_1} + Gp \ln \frac{T_2}{T_1}$$

$$Gp - Gr = \pi R \qquad \Delta S = Gr \ln \frac{T'}{T_1} + Gr \ln \frac{T}{T_1} + \pi R \ln \frac{T}{T_1}$$

$$Gp - Gr = \pi R \qquad \Delta S = (Gr + \pi R) \ln \frac{T'}{T_1} + Gr \ln \frac{T}{T_1} + \pi R \ln \frac{T}{T_1}$$

$$= \pi R \ln \frac{T}{T_1}, = \pi R \ln \frac{R}{R}$$

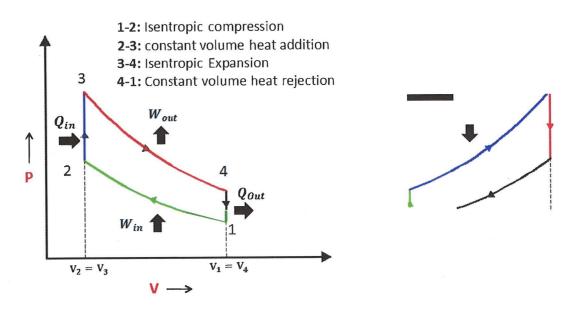
$$\Delta S = \pi R \ln V_2$$

$$\Delta S = \pi R \ln V_2$$

$$\Delta S = \pi R \ln V_2$$

Entropy change is a state function and it's change depends only on the initial as final points. (1)

3. Consider an ideal heat engine operated in a cycle using two adiabatic processes and two isochoric processes. The cycle in the p-V diagram is shown below: The heat engine is operated with an ideal gas as a working substance. Calculate



the efficiency of the heat engine and show that this efficiency is less than a Carnot engine working between the same hot and cold reservoirs.

[Marks=8+4]

$$Q_{in} = C_{V} (T_{3} - T_{2}) \qquad \text{and} \qquad Q_{out} = C_{V} (T_{4} - T_{7}) - 2$$

$$\therefore \eta = \frac{N}{Q_{in}} = N \text{ Nhere } N = N_{in} - N_{out}$$

$$= 1 - \frac{Q_{out}}{Q_{in}} = 1 - \frac{(T_{V} - T_{I})}{(T_{3} - T_{L})} - (4)$$

$$= \frac{1}{3} V_{2}^{Y-I} = T_{V} V_{I}^{Y-I}$$

$$= T_{2} V_{2}^{Y-I} = T_{1} V_{I}^{Y-I}$$

$$= T_{2} V_{2}^{Y-I} = T_{1} V_{I}^{Y-I}$$

$$\therefore \eta = 1 - \frac{1}{\chi^{Y-I}} = 1 - \chi$$

$$\text{Nhere } \chi = \frac{V_{I}}{V_{2}}$$

: nc > n - 1

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