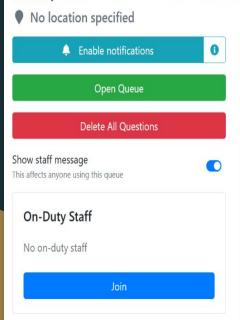
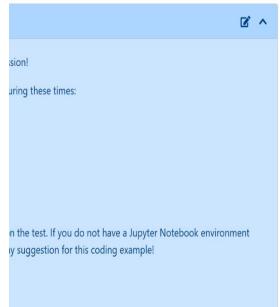
C.A.R.E. PHYS 213 Quiz 2 Review Session



CARE/CARE PHYS 213 Exam Review Session







This queue is closed. Check back later!

Units for the Exam

- Kinetic Theory of Ideal Gases
- Quasistatic Processes
- Thermodynamic Cycles
- Gibbs Free Energy

Ideal gas and Equipartition

- Ideal Gas: Approximation of particles as points with no interactions:

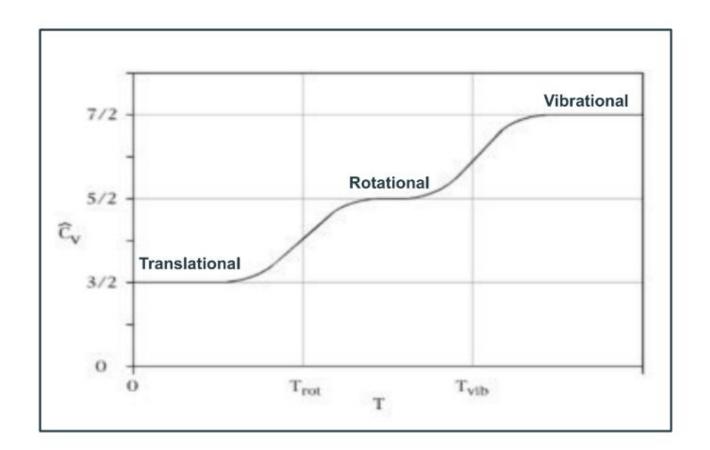
Follows ideal gas law:
$$pV = NkT = nRT$$

- **Equipartition:** each degree of freedom contributes $\frac{1}{2}kT$ of energy
 - Internal energy in each particle:

$$U = (N_{DOF}/2)kT$$

Molar heat capacity:

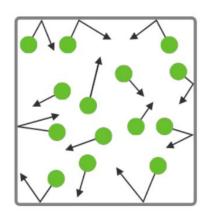
$$c_{\rm M} = (N_{\rm DOF}/2)kN_{\rm A}$$



Root-Mean-Square Velocity

- v_{rms} : Average (translational) velocity of gas particles
- Translational Kinetic Energy: $KE_{translational} = 1/2 \text{ m} (v_{rms})^2$
- Relationship to temperature:

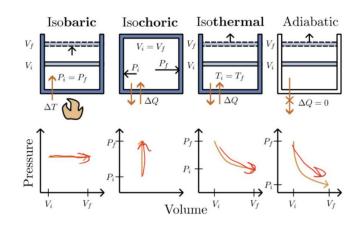
$$\frac{1}{2}mv_{rms}^{2} = \frac{3}{2}kT$$



- This only applies to ONE PARTICLE
- Notice: this does <u>NOT</u> depend on the number of DOFs; it's <u>ALWAYS</u> (3/2)kT
 - \circ Why? Translational KE only depends on the translational modes motion (there are only 3 translational modes: v_x , v_y , v_y)

Thermodynamic Processes

- <u>Isochoric</u> or Isovolumetric
 - Constant <u>VOLUME</u>
- <u>Isobaric</u>
 - Constant <u>PRESSURE</u>



- Isothermal
 - Constant <u>TEMPERATURE</u>, <u>REVERSIBLE</u>

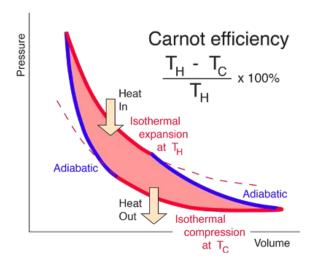
$$(\Delta S_{total} = 0, \Delta U = 0)$$

- Adiabatic
 - Constant <u>HEAT</u> (dQ = 0), <u>REVERSIBLE</u>

$$(\Delta S_{total} = 0, \Delta Q = 0)$$

Reversible Processes

- <u>Isothermal</u> + <u>Adiabatic</u> processes are <u>REVERSIBLE</u>
 - $\circ \quad \Delta S_{total} = 0 \quad \text{(no change in entropy)}$
 - For <u>isothermal</u> processes:
 - \blacksquare PV = constant
 - For <u>adiabatic</u> processes:
 - \blacksquare PV $^{\gamma}$ = constant
 - $\gamma = (2/N_{DOF}) + 1$ (given on equation sheet)



 Assume we have a gas undergoing an adiabatic process, determine the work done given the following parameters:

$$\circ$$
 V_i = 10 m³, p_i = 10 kPa

$$V_f = 4 \text{ m}^3, N_{DOF} = 3$$

1. Calculate γ

2. Find pV^{γ}

3. Calculate W

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$$W = \int_{V_i}^{V_f} p dV = \int_{V_i}^{V_f} \frac{C}{V_i^{\gamma}} dV = \left[C \frac{V_f \left(-\frac{5}{3} + 1 \right)}{-\frac{5}{3} + 1} \right] - \left[C \frac{V_i \left(-\frac{5}{3} + 1 \right)}{-\frac{5}{3} + 1} \right]$$

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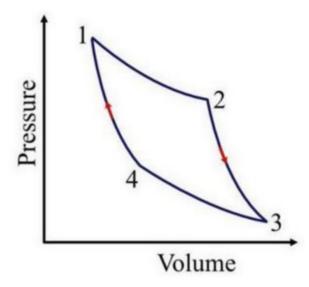
$$W = \left[(464158.88) \frac{4^{\left(-\frac{5}{3}+1\right)}}{-\frac{5}{3}+1} \right] - \left[(464158.88) \frac{10^{\left(-\frac{5}{3}+1\right)}}{-\frac{5}{3}+1} \right] \approx -126.30 \, kJ$$

p-V Diagrams

Used to visualize thermodynamic cycles

- Area enclosed in the curve is equal to the work per cycle
 - Clockwise direction: work is positive (engine did work)
 - Counterclockwise direction: work is negative (work done on engine)

$$W_{\mathrm{by}} = \int_{V_i}^{V_f} p \ \mathrm{d}V$$



Heat Engines

- Cycles of Thermodynamic processes are used to make engines, heat pumps, and refrigerators

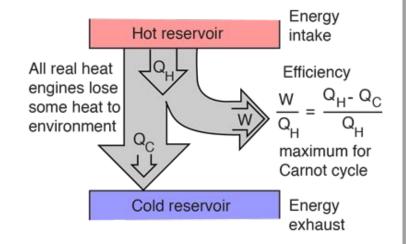
Efficiency of engines:
$$\epsilon = \frac{W_{by}}{Q_H} \le 1 - \frac{T_C}{T_H}$$

COP of pumps and refrigerators:

O Heat pump:
$$COP = \frac{Q_H}{W_{on}} \le \frac{1}{1 - \frac{T_C}{T_H}}$$

O Refrigerator:
$$COP = \frac{Q_C}{W_{on}} \le \frac{1}{\frac{T_H}{T_C} - 1}$$

Efficiency/COP can be thought of as "what you get out" divided by "what you put in."



Example Problem: Engine/Heat Pump COP

Given a hot reservoir at a temperature of $T_h=373K$ and a cold reservoir at a temperature of $T_c=293K$, calculate the maximum efficiency ϵ of an engine and the maximum COP of a heat pump between the two reservoirs.

Heat Engine Efficiency:	Heat Pump COP:

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Heat Engine Efficiency:

$$\epsilon = 1 - rac{T_c}{T_h} pprox 0.21$$

Heat Pump COP:

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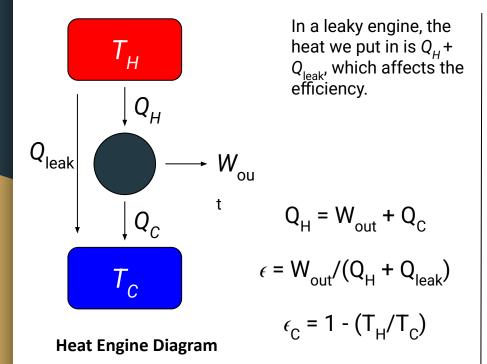
Heat Engine Efficiency:

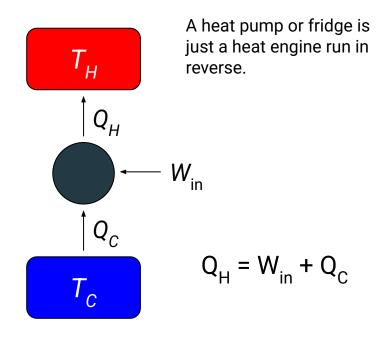
$$\epsilon = 1 - rac{T_c}{T_h} pprox 0.21$$

Heat Pump COP:

$$COP = rac{1}{1 - T_c/T_h} pprox 4.7$$

Engine, Pump, and Refrigerator Diagrams





Heat Pump/Refrigerator Diagram

Gibbs Free Energy

• Useful when temperature and pressure are fixed

$$G = U - T_{env}S + pV$$

- Minimizing Gibbs of a system will maximize total (system + environment) Entropy
 - As a system approaches equilibrium, free energy will decrease to a minimum

- Fundamental Thermodynamic Relation in Equilibrium:
 - $\circ \quad TdS = dU + pdV \mu dN$
 - \circ $\mu = (dG/dN) \rightarrow \mu N = G$ (at fixed temperature and pressure)
 - \circ Equilibrium favors lowest μ

Good luck!

Feel free to ask any questions you may have.

You got this!

