CAE 208 / MMAE 320: Thermodynamics Fall 2023

October 3, 2023 Energy analysis of closed systems (1)

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ANNOUNCEMENTS

Announcements

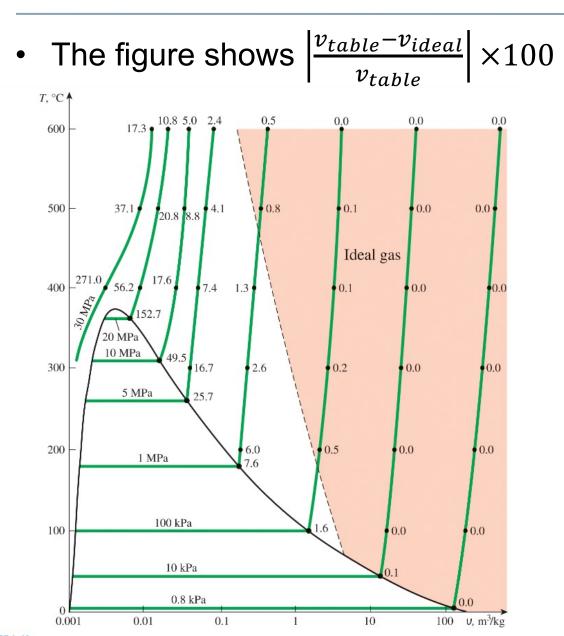
- The solution to Assignment 4 is posted
 Please submit one file only
 Always look at the solutions posted
- Assignment 5 will be posted next Tuesday (Time to focus on the exam)
- Midterm is scheduled for October 10
- Today's lecture will be in the exam (Content in our upcoming Thursday lecture will not be included in the exam)
- A problem-solving lecture was posted on Blackboard

Announcements

- For the first midterm exam:
 - □ The first part of the exam is closed book (You are allowed to have one page for your cheat sheet to include equations)
 - □ The second part of the exam is open book (only a thermodynamics book or thermodynamics property tables)

CLASS ACTIVITY

COMPRESSIBILITY FACTOR – A MEASURE OF OF DEVIATION FROM IDEAL-GAS BEHAVIOR



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 Z factor for all gases is approximately the same at the same reduced temperature and pressure due to the principle of corresponding states

$$v = \frac{RT}{P}$$

$$T_R = \frac{T}{T_{Cr}}$$

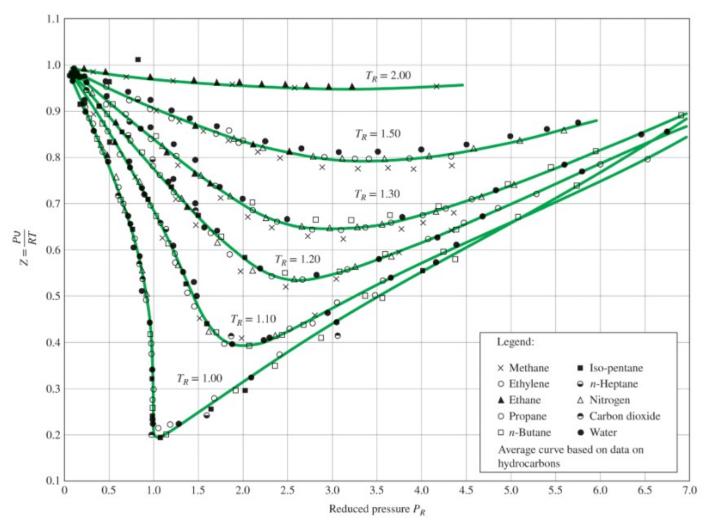
 Z factor for all gases is approximately the same at the same reduced temperature and pressure due to the principle of corresponding states

Pv = ZRT

$$v = \frac{ZRT}{P}$$

$$Z = \frac{v_{actual}}{v_{idea}}$$

- We can define a "generalized compressibility chart"
- Let's look at a few observations:



CLASS ACTIVITY

 Determine specific volume of refrigeratnt-134a at 1 MPa and 50 °C using (a) the ideal-gas equation of state and (b) the generalized compressibility chart. Compare the values obtained to the actual value of 0.021796 m³/kg and determine the error involved in each case.

• Solution (a):

TABLE A-1							
Molar mass, gas constant, and critic							
Substance	Formula Molary	ar mass, <i>M</i> kg/kmol	Gas constant, <i>R</i> kJ/kg · K*		Critical-point properties		
	Formula Molar	nass, <i>w</i> kg/knioi	Gas constant, A KJ/Kg · K		Temperature, K	Pressure, MPa	Volume, m ³ /l
Propane	C_3H_8	44.097	0.1885		370	4.26	0.1998
Propylene	C_3H_6	42.081	0.1976		365	4.62	0.1810
Sulfur dioxide	SO ₂	64.063	0.1298		430.7	7.88	0.1217
Tetrafluoroethane (R-134a)	CF_3CH_2F	102.03	0.08149		374.2	4.059	0.1993
Trichlorofluoromethane (R-11)	CCl ₃ F	137.37	0.06052		471.2	4.38	0.2478
Water	H_2O	18.015	0.4615		647.1	22.06	0.0560
Xenon	Xe	131.30	0.06332		289.8	5.88	0.1186

$$v = \frac{RT}{P} = \frac{\left(0.0815 \frac{kJ}{kg.K}\right)(50 + 273.15 K)}{1000 \, kPa} = 0.026325 \frac{m^3}{kg}$$

$$Error = \frac{0.026325 - 0.021796}{0.021796} = 0.208$$

• Solution (b):

$$P_R = \frac{P}{P_{cr}} = \frac{1 MPa}{4.059 MPa} = 0.246$$

Z = 0.84

$$T_R = \frac{T}{T_{cr}} = \frac{323 \ K}{374.2 \ K} = 0.863$$

$$v_{actual} = Zv_{ideal} = (0.84) \left(0.026325 \frac{m^3}{kg} \right) = 0.022113 \frac{m^3}{kg}$$

$$Error = \frac{0.022113 - 0.021796}{0.021796} \sim 0.02$$

CLASS ACTIVITY

- Determine the specific volume of refrigerant-134a vapor at 0.9 MPa and 70°C based on
 - a) The ideal-gas equation
 - b) The generalized compressibility chart
 - c) Data from tables. Also, determine the error involved in the first two cases.

• Solution (a):

TABLE A-1

Molar mass, gas constant, and critical-point properties

Substance	Formula Mola	r mass, <i>M</i> kg/kmol	Gas constant, <i>R</i> kJ/kg · K*		Critical-point properties			
Substance	Formula Mola	i mass, w kg/kmoi	Gas constant, A KJ/Kg · K		Temperature, K	Pressure, MPa	Volume, m ³ /kmol	
Propane	C_3H_8	44.097	0.1885		370	4.26	0.1998	
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Xenon	Xe	131.30	0.06332		289.8	5.88	0.1186	

$$R = 0.08149 \frac{kJ}{kg - K}$$

 $T_{cr} = 374.2 K$

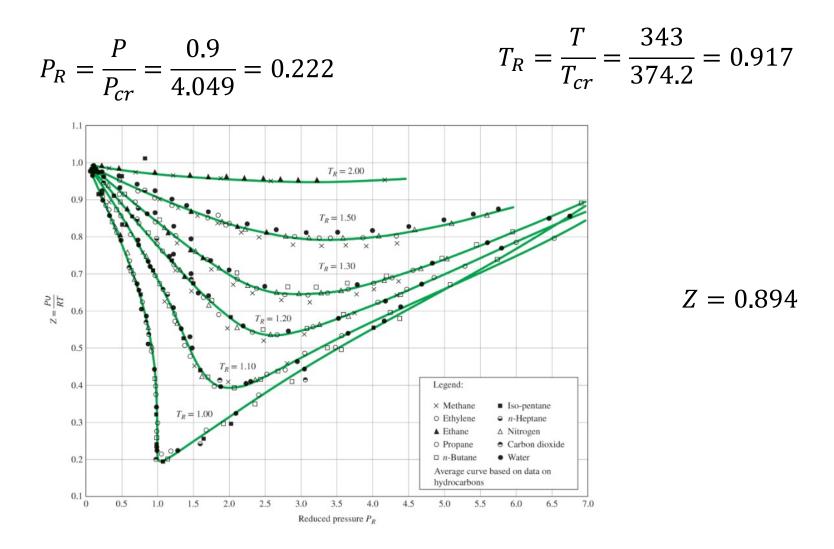
 $P_{cr} = 4.049 MPa$

• Solution (a):

Pv = RT

$$v = \frac{RT}{P} = \frac{(0.08149 \frac{kJ}{kg - K})(273.15 + 70 K)}{0.9 \times 10^3 kPa} = 0.03105 \frac{m^3}{kg}$$

• Solution (b):



• Solution (b):

$$v = Zv_{ideal} = (0.894) \left(0.03105 \frac{m^3}{kg} \right) = 0.02776 \frac{m^3}{kg}$$

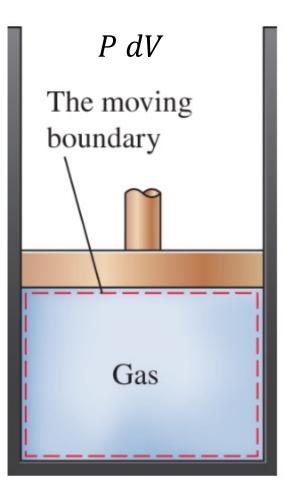
• Solution (c):

TABLE A-13										
Super	Superheated refrigerant-134a									
T °C	U m ³ /kg	<i>u</i> kJ/kg	h kJ/kg	s kJ/kg ∙ K	v m ³ /kg	u kJ/kg	<i>h</i> kJ/kg	s kJ/kg ∙ K		
	$P = 0.80 \text{ MPa} (T_{\text{sat}} = 31.31^{\circ}\text{C})$				$P = 0.90 \text{ MPa} (T_{\text{sat}} = 35.51^{\circ}\text{C})$					
Sat.	0.025645	246.82	267.34	0.9185	0.02268	36 248.82	269.25	0.9169		
40	0.027035	254.84	276.46	0.9481	0.02337	253.15	274.19	0.9328		
50	0.028547	263.87	286.71	0.9803	0.02480	9 262.46	284.79	0.9661		
60	0.029973	272.85	296.82	1.0111	0.02614	6 271.62	295.15	0.9977		
70	0.031340	281.83	306.90	1.0409	0.02741	3 280.74	305.41	1.0280		
80	0.032659	290.86	316.99	1.0699	0.02863	30 289.88	315.65	1.0574		
90	0.033941	299.97	327.12	1.0982	0.02980	06 299.08	325.90	1.0861		

QUIZ

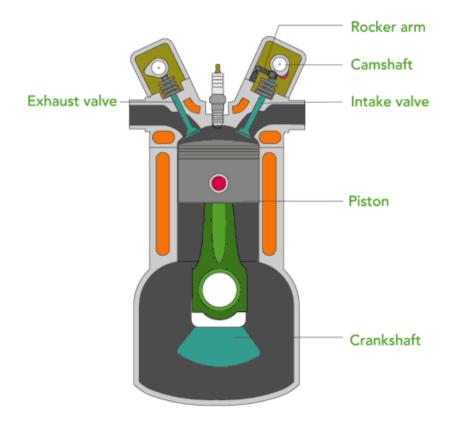
MOVING BOUNDARY WORK

 The expansion or compression work is often called moving boundary work or simply boundary work

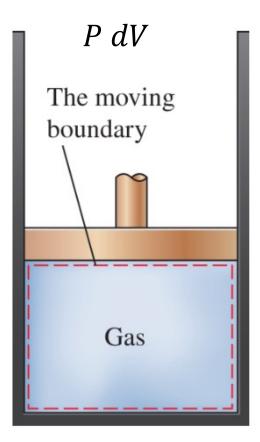


Think about internal combustion engines

 The expansion or compression work is often called moving boundary work or simply boundary work

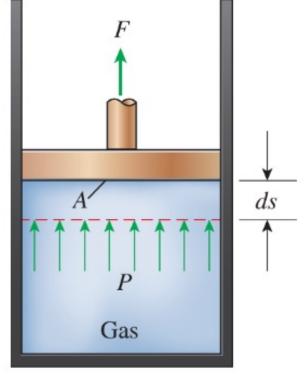


 The moving work associated with real engines or compressors cannot be determined exactly from a thermodynamic analysis alone. Why?



- Consider the gas enclosed in the cylinder-piston device
- The process is quasi equilibrium (ds)

$$\delta W_b = Fds = PAds = P \, dV$$

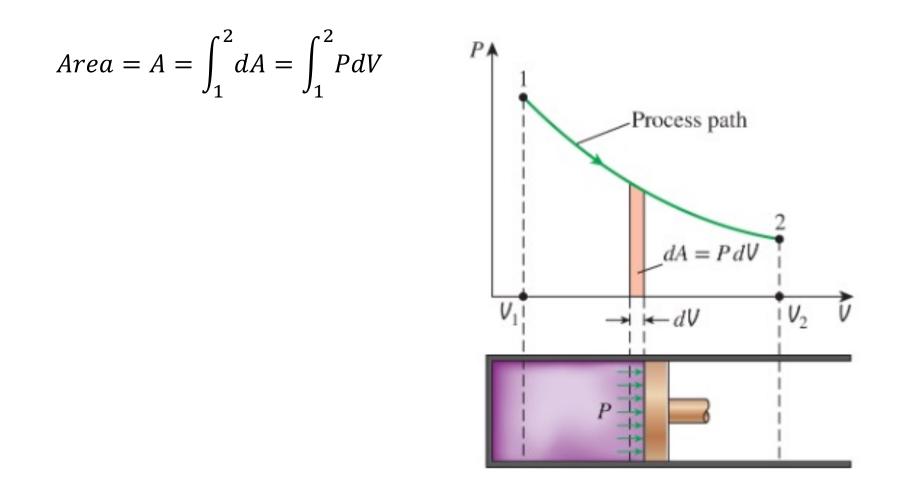


• Let's think about dV in terms of expansion and compression

 The total boundary work done during the entire process as the piston moves is obtained by adding all the differential works from the initial state to the final state

$$W_b = \int_1^2 P dV$$

• For a quasi-equilibrium expansion process, we can write:



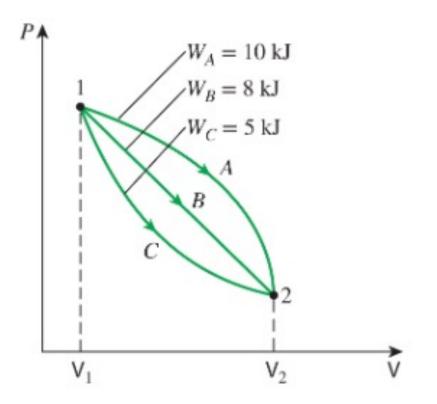
• Let's look at some basic integration

$$Area = A = \int_{1}^{2} dA = W_b = \int_{1}^{2} P dV$$

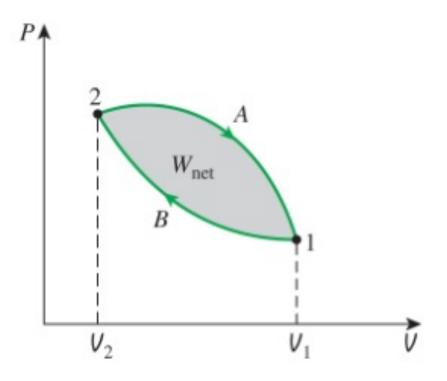
• Basically, we can say:

The area under the process curve on a P-V diagram is equal, in magnitude, to the work done during a quasiequilibrium expansion or compression process of a closed system (on the P-V diagram, in presents the boundary work done per unit mass)

 The boundary work done during a process depends on the path followed as well as the end states:



 The net work done during a cycle is the difference between the work done by the system and the work done on the system. Why?



• We can use the equation for nonquasi-equilibrium process since properties are defined for equilibrium states only

$$W_b = \int_1^2 P_i dV$$

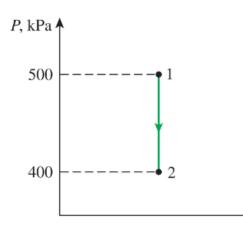
• We can use the equation for nonquasi-equilibrium process since properties are defined for equilibrium states only

$$W_{b} = W_{friction} + W_{atm} + W_{crank} = \int_{1}^{2} (F_{friction} + P_{atm}A + F_{crank}) dx$$

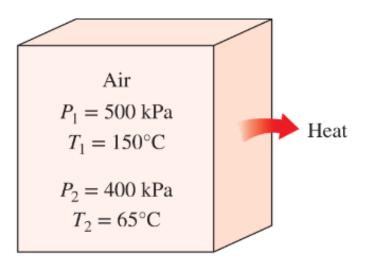
CLASS ACTIVITY

Class Activity

 A rigid tank contains air at 500 kPa and 150 °C. As a result of heat to the surroundings, the temperature and pressure inside the tank drop to 65 °C and 400 kPa, respectively.
 Determine the boundary work done during this process • Solution



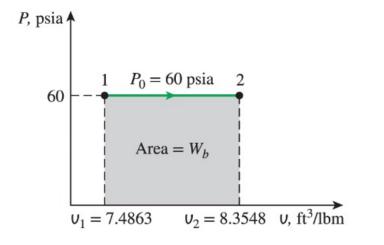
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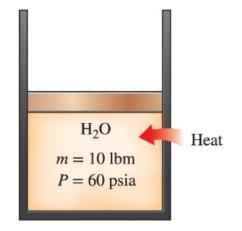


CLASS ACTIVITY

Class Activity

 A frictionless piston-cylinder device contains 10 lbm of steam at 60 psia and 320 °F. Heat is now transferred to the steam until the temperature reaches 400 °F. If the piston is not attached to a shaft and its mass is constant, determine work done by the steam during this process. • Solution:





$$W_b = \int_1^2 P dV = P_0 \int_1^2 dV = P_0 (V_2 - V_1)$$

 $W_b = mP_0(v_2 - v_1)$

• Solution:

$$\begin{cases} P_1 = 60 \ psia \\ T_1 = 320 \ ^\circ F \end{cases} \quad v_1 = 7.4863 \frac{ft^3}{lbm} \end{cases}$$

Use Table A-6

$$\begin{cases} P_2 = 60 \ psia \\ T_2 = 400 \ ^\circ F \end{cases} \quad v_2 = 8.3548 \frac{ft^3}{lbm} \end{cases}$$

$$W_b = mP_0(v_2 - v_1)$$

= (10 lbm)(8.3548 - 7.4863) $\left(\frac{ft^3}{lbm}\right) \left(\frac{1Btu}{5.404 \, psia - ft^3}\right) = 96.4 \, Btu$

Class Activity

Conversion Factors

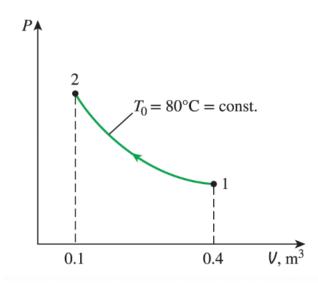
Dimension	Metric	English
Acceleration	$1 \text{ m/s}^2 = 100 \text{ cm/s}^2$	$1 \text{ m/s}^2 = 3.2808 \text{ ft/s}^2$ $1 \text{ ft/s}^2 = 0.3048 \text{ m/s}^2$
Area	$1 \text{ m}^2 = 104 \text{ cm}^2 = 10^6 \text{ mm}^2 = 10^{-6} \text{ km}^2$	$1 \text{ m}^2 = 1550 \text{ in.}^2 = 10.764 \text{ ft}^2$ $1 \text{ ft}^2 = 144 \text{ in.}^2 = 0.0929034 \text{ m}^2$
Density	$1 \text{ g/cm}^3 = 1 \text{ kg/L} = 1000 \text{ kg/m}^3$	1 g/cm ³ = 62.428 lbm/ft ³ = 0.036127 lbm/in. ³ 1 lbm/in. ³ = 1728 lbm/ft ³ 1 kg/m ³ = 0.062428 lbm/ft ³
Energy, Heat, Work, Internal Energy, Enthalpy	$1 kJ = 1000 J = 1000 N \cdot m = 1 kPa \cdot m^{3}$ $1 kJ/kg = 1000 m^{2}/s^{2}$ 1 kWh = 3600 kJ 1 Wh = 3600 J 1 cal = 4.1868 J 1 Cal = 4.1868 kJ	1 kJ = 0.94782 Btu 1 Btu = 1.055056 kJ = 5.40395 psia·ft ³ = 778.169 lbf·ft 1 Btu/lbm = 25.037 ft ² /s ² = 2.326 kJ/kg 1 kJ/kg = 0.430 Btu/lbm 1 kWh = 3412.14 Btu 1 therm = 10^5 Btu = 1.055×10^5 kJ (natural gas)

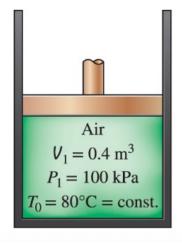
CLASS ACTIVITY

Class Activity

A piston-cylinder device initially contains 0.4 m³ of air at 100 kPa and 80 °C. The air is now compressed to 0.1 m³ in such a way that the temperature inside the cylinder remains constant. Determine the work done during this process

• Solution:

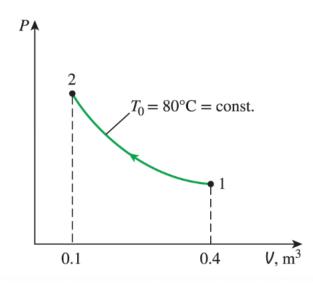




$$PV = mRT_0 \rightarrow P = \frac{C}{V}$$

$$W_{b} = \int_{1}^{2} P dV = \int_{1}^{2} (\frac{C}{V}) dV = C \int_{1}^{2} (\frac{dV}{V})$$

• Solution:

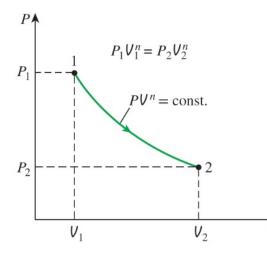


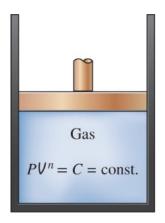
$$W_b = C \times Ln(V) \Big|_{V_1}^{V_2} = C[Ln(V_2) - Ln(V_1)] = P_1 V_1 \times Ln(\frac{V_2}{V_1})$$

$$W_b = (100 \ kPa)(0.4 \ m^3) \left(\ln\left(\frac{0.1}{0.4}\right) \right) \left(\frac{1kJ}{1 \ kPa. \ m^2}\right) = -55.5 \ kJ$$

 During expansion or compression processes of gases and volume are often related in PVⁿ = C which is known as a Polytropic process

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$$P = CV^{-n}$$

• Polytropic process

 $P = CV^{-n}$

$$W_b = \int_1^2 P \, dV = \int_1^2 (CV^{-n}) dV = \frac{C((V^{-n+1} - V^{-n+1}))}{(-n+1)} = \frac{P_2 V_2 - P_1 V_1}{1 - n}$$

• Polytropic process

 $C = P_1 V_1^n = P_2 V_2^n$

$$W_b = \frac{mR(T_2 - T_1)}{1 - n} \quad n \neq 1 \ (kJ)$$

• For the case of n =1

$$W_{b} = \int_{1}^{2} P dV = \int_{1}^{2} CV^{-1} dV = PV \times Ln(\frac{V_{2}}{V_{1}})$$

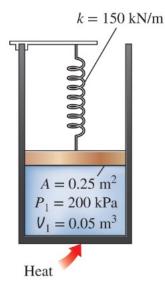
CLASS ACTIVITY

Class Activity

- A piston-cylinder device contains 0.05 m³ of a gas initially at 200 kPa. At this state, a linear spring that has a spring constant of 150 kN/m is touching the piston but exerting no force on it. Now heat is transferred to the gas, causing the piston to rise and to compress the spring until the volume inside the cylinder doubles. If the cross-sectional area of the piston is 0.25 m², determine
 - a) The final pressure inside the cylinder
 - b) The total work done by the gas
 - c) The fraction of this work done against the spring to compress it

Class Activity

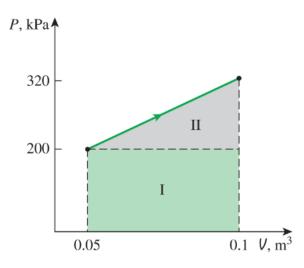
• Solution



$$V_2 = 2V_1 = 2(0.05 m^3) = 0.1 m^3$$

$$x = \frac{\Delta V}{A} = \frac{(0.1 - 0.05)m^3}{0.25 m^2} = 0.2 m$$

• Solution

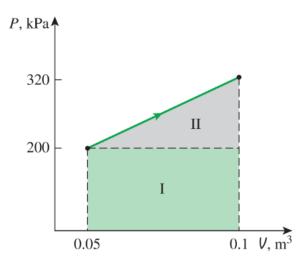


$$F = kx = \left(150\frac{kN}{m}\right)(0.2\ m) = 30\ kN$$

$$P = \frac{F}{A} = \frac{30 \ kN}{0.25 \ m^2} = 120 \ kN$$

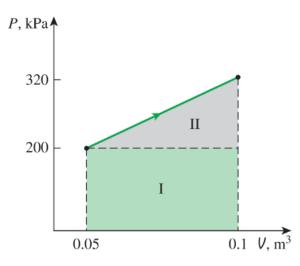
 $200 + 120 = 320 \ kN$

• Solution (b)



$$W = area = \frac{(200 + 320)kPa}{2} [0.1^2 - 0.05^2] \left(\frac{1kJ}{1 \ kPa. \ m^2}\right) = 13 \ kJ$$

• Solution (c)



$$W_{spring} = \frac{1}{2} [320 - 200] kPa(0.05 \ m^3) = 3 \ kJ$$

$$W_{spring} = \frac{1}{2}k(x_2^2 - x_1^2) = \frac{1}{2}(150)(0.2^2 - 0^2) = 3 kJ$$

• Moving boundary work under different processes

Process	Moving boundary work
Constant volume	0
Constant pressure	$P_0(V_2-V_1)$
Isothermal	$P_{1}V_{1} \times Ln(\frac{V_{2}}{V_{1}})$ $P_{1}V_{1} \times Ln(\frac{P_{1}}{P_{2}})$ $mRT_{o} \times Ln(\frac{V_{2}}{V_{1}})$
Polytropic	$\frac{\frac{P_2V_2 - P_1V_1}{1 - n}}{\frac{mR(T_2 - T_1)}{1 - n}}$

ENERGY BALANCE FOR CLOSED SYSTEMS

 The net change (increase or decrease) in the total energy of the system during a process is equal to the difference between the total energy entering and the total energy leaving the system during the process

(Total energy entering the system) – (Total energy leaving the system) = (Change in the total energy of the sysem)

$$E_{in} - E_{out} = \Delta E_{system}$$

This is known as the energy balance

• Energy change of a system ΔE_{system}

Energy change = Energy at final state - Energy at initial state

$$\Delta E_{system} = E_{final} - E_{initial} = E_2 - E_1$$

Energy is a property, and the value of a property does not change unless the state of the system changes

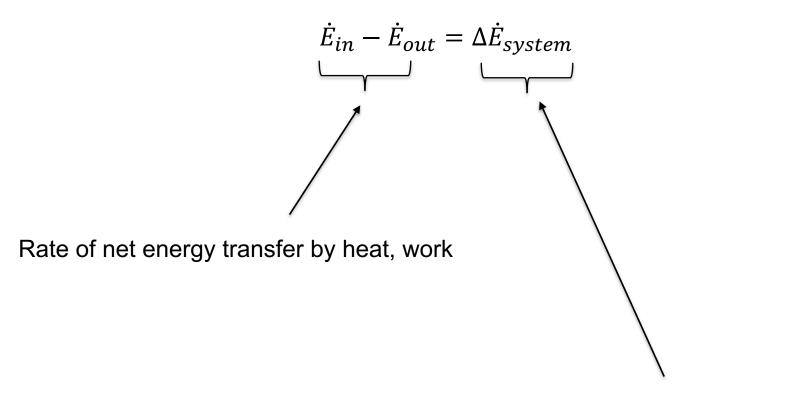
• We can sum the heat, work, and mass, and the heat transfer:

$$E_{in} - E_{out} = (Q_{in} - Q_{out}) + (W_{in} - W_{out}) = \Delta E_{system}$$

Net energy transfer by heat, work

Change in internal, kinetic, potential, ..., energies

 We can sum the heat, work, and mass, and the heat transfer in the rate form:



Rate of change in internal, kinetic, potential, ..., energies

 The energy balance can be expressed on a per unit mass basis as

 $e_{in} - e_{out} = \Delta e_{system}$

• For constant rates, we can write:

$$Q = \dot{Q} \Delta t$$

 $W = \dot{W} \Delta t$

$$E = \left(\frac{dE}{dt}\right)\Delta t$$

 For a closed system undergoing a cycle, the initial and final states are identical:

V

• We can write:

