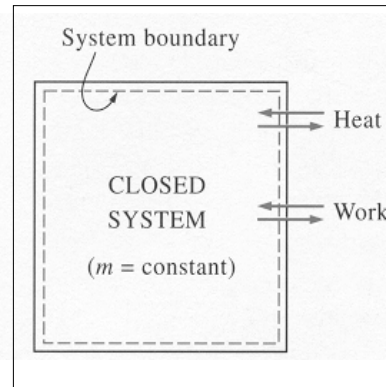


## First Law of Thermodynamics

- During an interaction between a system and its surroundings, the amount of energy gained by the system must be exactly equal to the amount of energy lost by the surroundings.
- Energy can cross the boundary of a closed system in two distinct forms: **heat** and **work**.



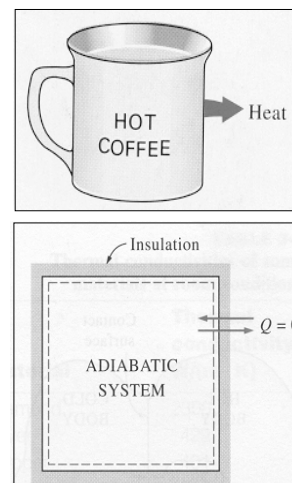
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## Heat Transfer

- Heat is a form of energy that is transferred between a system and its surroundings by virtue of a temperature difference.
- A process during which there is no heat transfer is called an adiabatic process.
- An adiabatic process should not be confused with an isothermal process.



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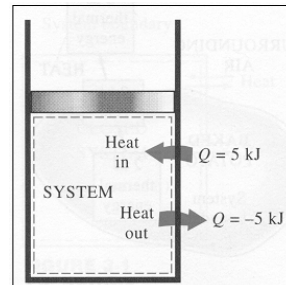
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## Heat: Units & Direction

- Heat has energy units, kJ (or Btu).
- The amount of heat transferred during a process between two states is denoted by  $Q_{12}$  or just  $Q$ .
- Heat transfer per unit mass of a system is denoted  $q$  as:

$$q = \frac{Q}{m} \quad \text{kJ/kg}$$

- Heat is a directional quantity. Heat transfer **to** a system is **positive**, and heat transfer **from** a system is **negative**.



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## Modes of Heat Transfer

- Heat can be transferred in three different ways: **conduction**, **convection**, and **radiation**.
- All modes of heat transfer require the existence of a temperature difference.
- All modes of heat transfer are from the high-temperature medium to a lower temperature one.

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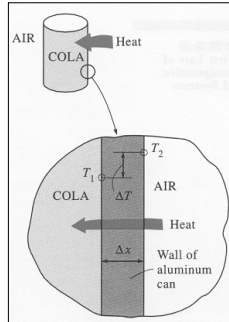
# Conduction Heat Transfer

## Fourier's Law:

$$\dot{Q}_{cond} = kA \frac{\Delta T}{\Delta x}$$

or

$$\dot{Q}_{cond} = -kA \frac{dT}{dx} \quad (\text{W})$$



## Thermal Conductivity

Material	Thermal conductivity, W/(m · K)
Diamond	2300
Silver	429
Copper	401
Gold	317
Glass fiber	0.043
Air (g)	0.026
Urethane, rigid foam	0.026

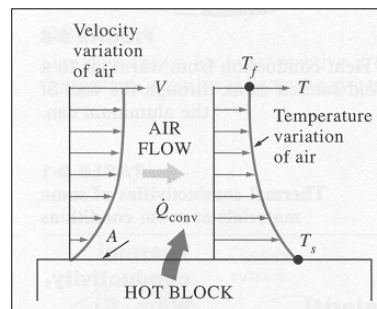
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# Convection Heat Transfer

- Convection is the mode of energy transfer between a solid surface and the adjacent liquid or gas which is in motion.
- It involves the combined effects of conduction and fluid motion.
- The faster the fluid motion, the greater the convection heat transfer.



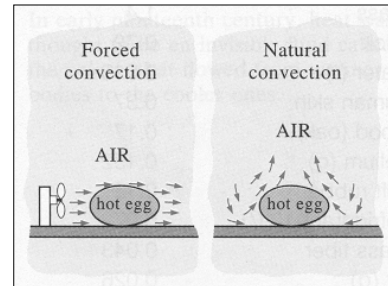
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## Forced vs. Natural Convection

- Convection is called **forced convection** if the fluid is forced to flow by external means such as a fan, or a pump,
- Convection is called **free or natural convection** if the fluid motion is caused by buoyancy forces that are induced by density differences.



$$\dot{Q}_{conv} = hA(T_s - T_f)$$

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## Heat Transfer Coefficient, h

$$\dot{Q}_{conv} = hA(T_s - T_f)$$

- h is a function of surface geometry, the nature of fluid motion, the properties of the fluid, and the bulk fluid velocity.
- Typical values of h, in W/(m<sup>2</sup> .K):
  - 2-25 free convection of gases,
  - 50-1000 free convection of liquids,
  - 25-250 forced convection of gases,
  - 50-20,000 forced convection of liquids,
  - 2500-100,000 convection in boiling and condensation processes.

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## Radiation Heat Transfer

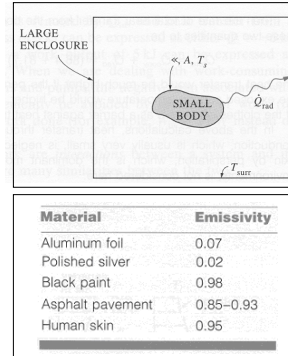
- Radiation is the energy emitted by matter in the form of electromagnetic waves (or photons) as a result of the changes in the electronic configurations of the atoms or molecules.

### Stefan-Boltzmann Law:

$$\dot{Q}_{emit,max} = \sigma A T_s^4$$

$$\dot{Q}_{rad} = \epsilon \sigma A (T_s^4 - T_{surr}^4)$$

- $\sigma = 5.67 \times 10^{-8} \text{ W}/(\text{m}^2 \cdot \text{K}^4)$
- $0 \leq \epsilon \leq 1$
- Temperatures are **absolute!**



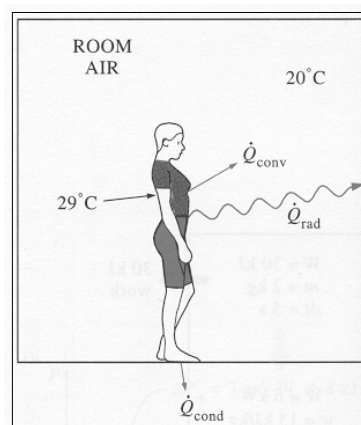
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## Example

- Given:
  - $A = 1.9 \text{ m}^2$
  - $h = 6 \text{ W}/\text{m}^2 \cdot \text{C}$
- Calculate total rate of heat transfer.



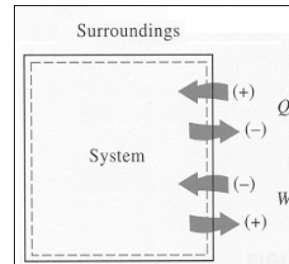
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## Work

- Work is an energy interaction which is not caused by a temperature difference.
- Work is associated with a force acting through a distance (rising piston, rotating shaft, electric wire crossing the system boundaries).
- Work done by a system is positive, and work done on a system is negative.



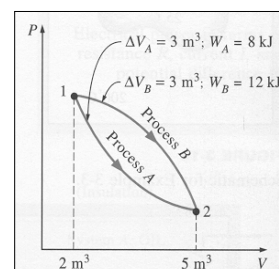
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## Similarities

- Both heat and work are boundary phenomena.
- Systems possess energy, but not heat or work. That is, heat and work are transfer phenomena.
- Both are associated with a process, not a state.
- **Both are path functions.**



$$\int_1^2 dV = V_2 - V_1 = \Delta V$$

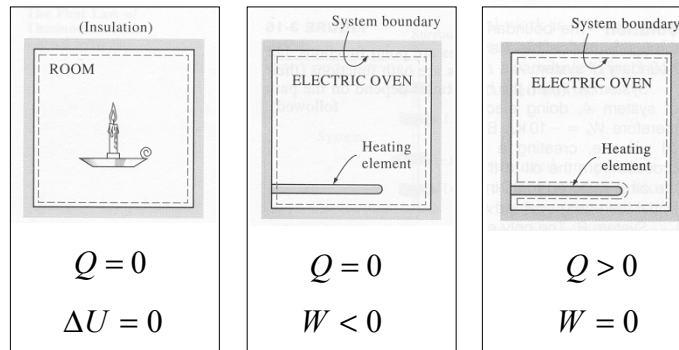
$$\int_1^2 \delta W = W_{12} \quad \text{NOT} \quad \Delta W$$

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## Examples



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## Electrical Work

- In an electric field, electrons in a wire move under the effect of electromotive forces, doing work.

$$W_e = V N \quad (\text{kJ}) \quad \dot{W}_e = V I \quad (\text{kW})$$

$$W_e = \int_1^2 V I dt \quad (\text{KJ})$$

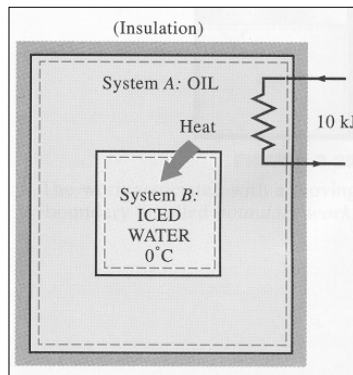
$$W_e = V I \Delta t \quad (\text{KJ})$$

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## Example



- Discuss the heat and work interactions for system A, B, and the combined system.

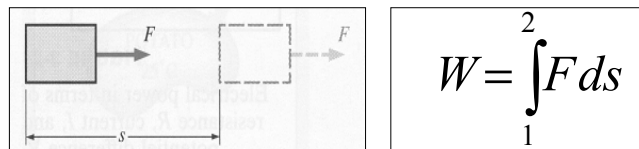
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## Mechanical Work

- Work is force times displacement.



### **Requirements for a work interaction:**

- (1) There must be a force acting on the boundary, and
- (2) The boundary must move.

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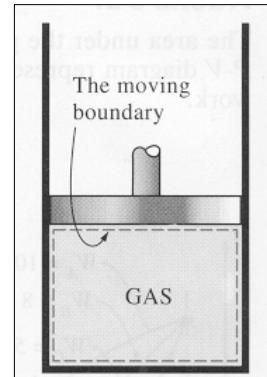
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## Moving Boundary Work

- It is associated with the expansion or compression of a gas in a piston-cylinder device,
- It is also called **boundary work** or  **$PdV$  work**,
- Moving boundary work is the primary form of work involved in automobile engines.



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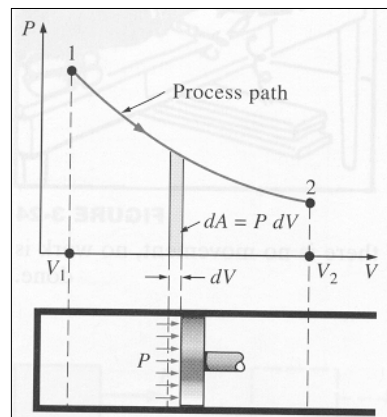
## Boundary Work

(Quasi-Equilibrium Process)

$$\delta W_b = F ds = P A ds = P dV$$

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

$$\text{area} = A = \int_1^2 dA = \int_1^2 P dV$$



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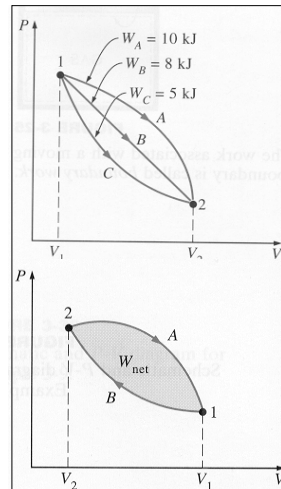
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## Boundary Work

(Quasi-Equilibrium Process)

- 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.

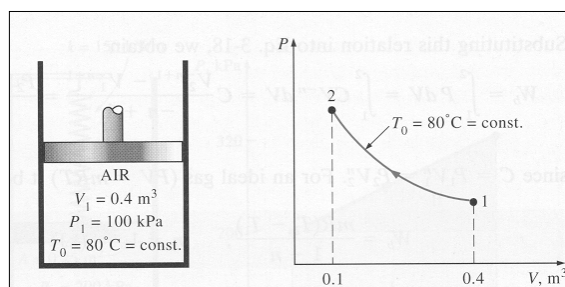


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## Example



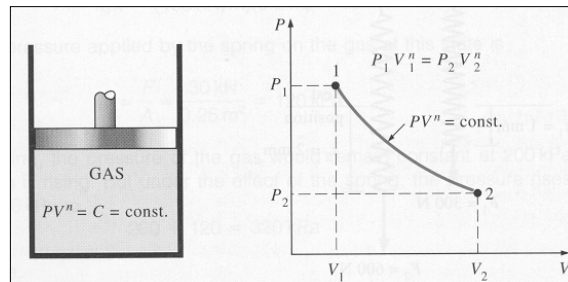
- Determine the work done during this process.

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## Polytropic Process



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

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

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## Spring Work

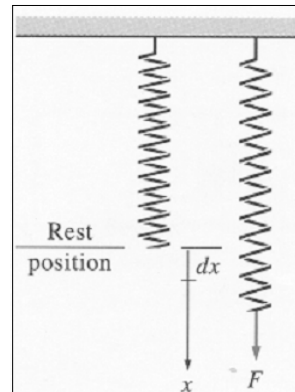
$$\delta W_{\text{spring}} = F dx \quad (\text{kJ})$$

- For linear elastic springs:

$$F = k_s x \quad (\text{kN})$$

$k_s$  = Spring constant (kN/m)

$$W_{\text{spring}} = \frac{1}{2} k_s (x_2^2 - x_1^2)$$



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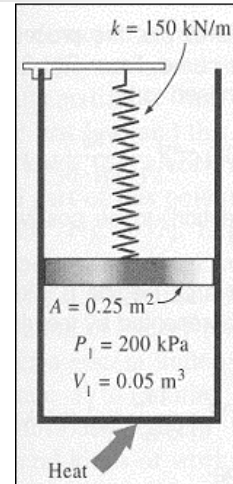
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## Example

### Determine:

- Final pressure inside the cylinder,
- Total work done by the gas, and
- Fraction of this work done against the spring to compress it.

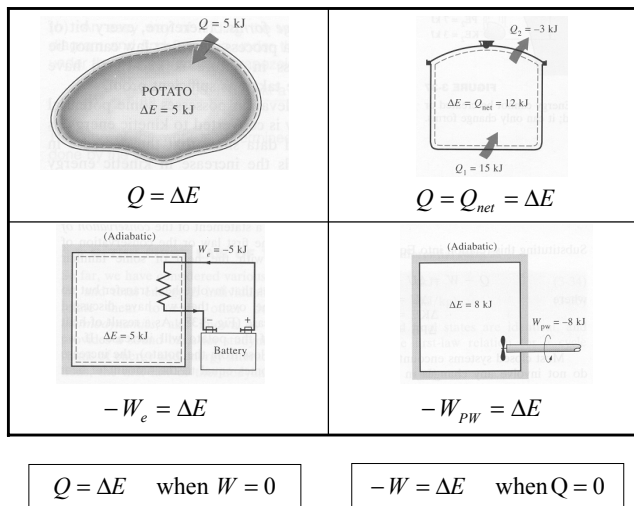


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## First Law of Thermodynamics



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## First Law of Thermodynamics

$$\left[ \begin{array}{c} \text{Net energy transfer to (or from)} \\ \text{the system as heat and work} \end{array} \right] = \left[ \begin{array}{c} \text{Net increase (or decrease) in the} \\ \text{total energy of the system} \end{array} \right]$$

$$Q - W = \Delta E \quad (\text{kJ})$$

Where:

$$Q = \sum Q_{in} - \sum Q_{out}$$

$$W = \sum W_{out} - \sum W_{in}$$

$$\Delta E = E_2 - E_1$$

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## First Law of Thermodynamics

$$\Delta E = \Delta U + \Delta KE + \Delta PE$$

$$Q - W = \Delta U + \Delta KE + \Delta PE$$

$$\text{Where } \begin{cases} \Delta U = m(u_2 - u_1) \\ \Delta KE = 1/2 m(v_2^2 - v_1^2) \\ \Delta PE = mg(z_2 - z_1) \end{cases}$$

For **Stationary Systems**:

$$Q - W = \Delta U$$

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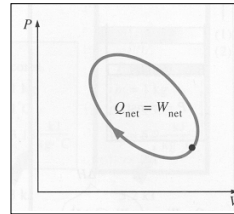
## Other Forms of the First Law

Per unit mass:  $q - w = e$  (kJ/kg)

In rate form:  $\dot{Q} - \dot{W} = \frac{dE}{dt}$  (kW)

For **cyclic processes**:

$$Q - W = 0 \quad (\text{kJ})$$



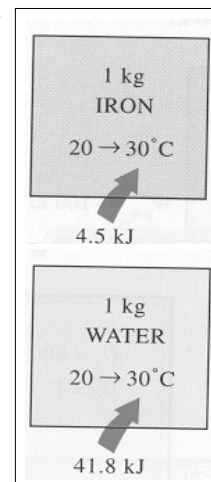
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## Specific Heats

- **Specific heat** is defined as the energy required to raise the temperature of a unit mass of a substance by one degree.
- Specific heat depends on how the process is executed.
- Specific heat at constant pressure  $C_p$  is always greater than specific heat at constant volume  $C_v$ .



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## Specific Heats

For a stationary closed system undergoing a constant-volume process:

$$\delta q - \delta w_{other} = du \quad C_v dT = du$$

$$C_v = \left( \frac{\partial u}{\partial T} \right)_v$$

$$C_p = \left( \frac{\partial h}{\partial T} \right)_p$$

- Note that  $C_v$  and  $C_p$  are thermodynamic properties. They are both state functions.

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## U, h, and C for Ideal Gases

Joule's Experiment:

$$u = u(T)$$

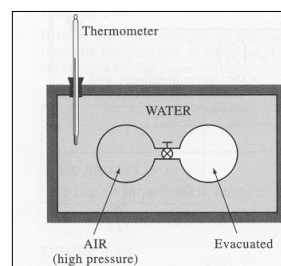
$$h = u + Pv$$

$$Pv = RT \rightarrow h = u + RT$$

$$h = h(T)$$

$$du = C_v(T) dT$$

$$dh = C_p(T) dT$$



$$\Delta u = u_2 - u_1 = \int_1^2 C_v(T) dT$$

$$\Delta h = h_2 - h_1 = \int_1^2 C_p(T) dT$$

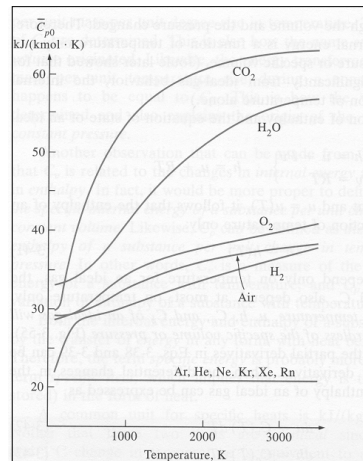
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## Specific Heats of Ideal Gases

- Specific heats of gases with complex molecules are higher and increase with temperature.
- Variation of specific heats with temperature may be approximated as linear over small temperature intervals.
- Ideal-gas specific heats of monatomic gases remain constant over the entire temperature range.



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## Specific Heats of Ideal Gases

$$h = u + RT \Rightarrow dh = du + R dT$$

$$C_p = C_v + R \quad (\text{kJ/kg.K})$$

On a molar basis:

$$\bar{C}_p = \bar{C}_v + R_u \quad (\text{kJ/kmol.K})$$

Specific Heat Ratio:  $k = \frac{C_p}{C_v} \begin{cases} k = 1.667 & \text{Monatomic Gases (Ar)} \\ k = 1.4 & \text{Diatomic Gases (Air)} \end{cases}$

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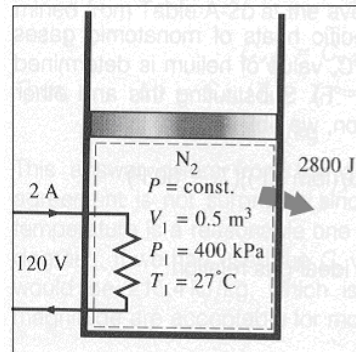
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## Example

- Determine the final temperature of the nitrogen.



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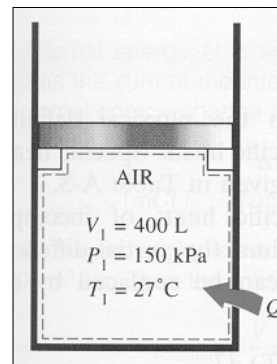
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## Example

Determine:

- Final temperature,
  - Work done by the air,
  - Total heat added.
- 350 kPa pressure is required to move the piston.



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## u, h, and C of Solids and Liquids

- Solids and liquids are essentially incompressible.

$$C_p = C_v = C \quad (\text{Solids \& Liquids})$$

$$du = C_v dT = C(T) dT \Rightarrow \Delta u = u_2 - u_1 = \int_1^2 C(T) dT$$

$$h_2 - h_1 = (u_2 - u_1) + v(P_2 - P_1) \Rightarrow \Delta h = \Delta u + v\Delta P$$

- The term ( $v\Delta P$ ) is often small compared with the first term ( $\Delta u$ ) and can be neglected without significant loss in accuracy.

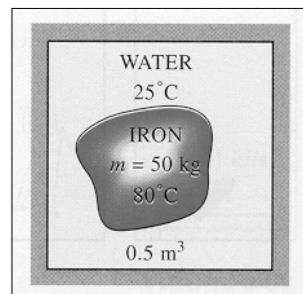
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## Example

- Determine the temperature when thermal equilibrium is reached.



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