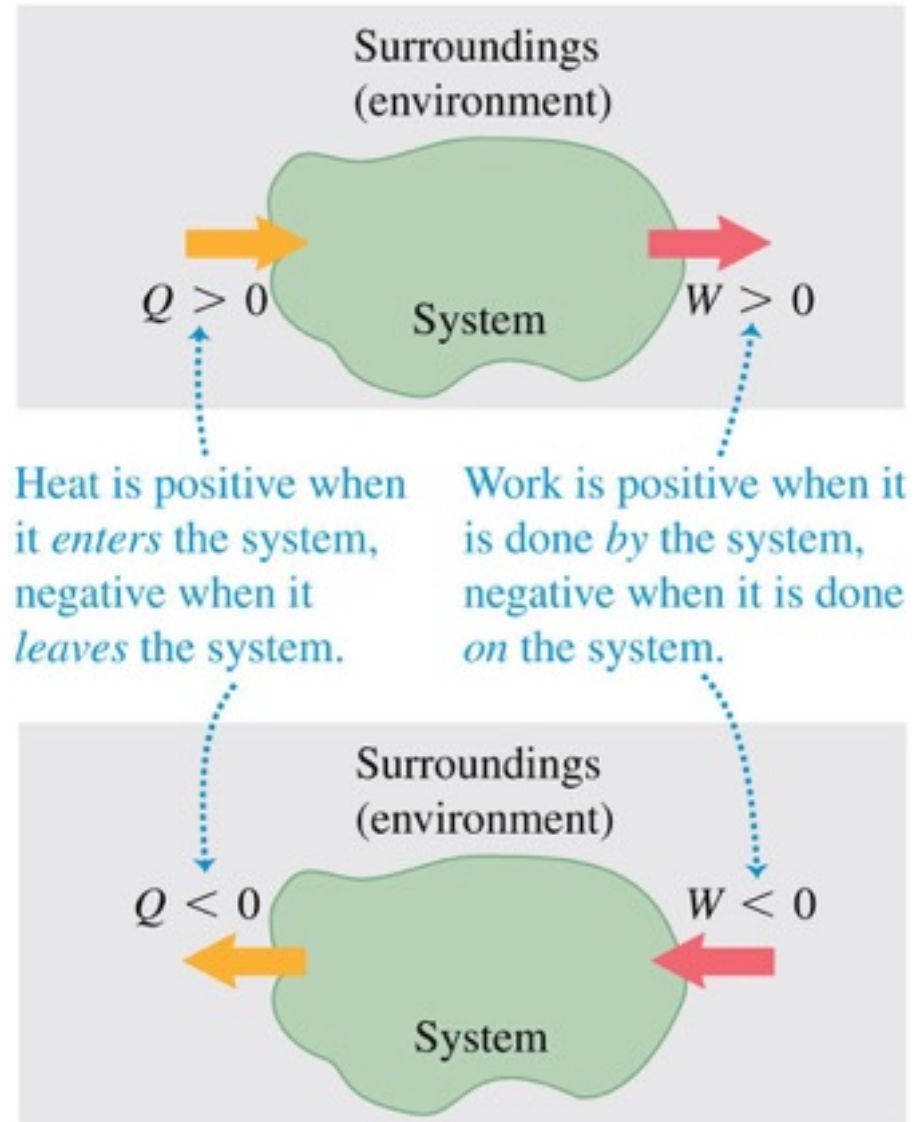


# Lecture 5

PHYC 161 Fall 2016

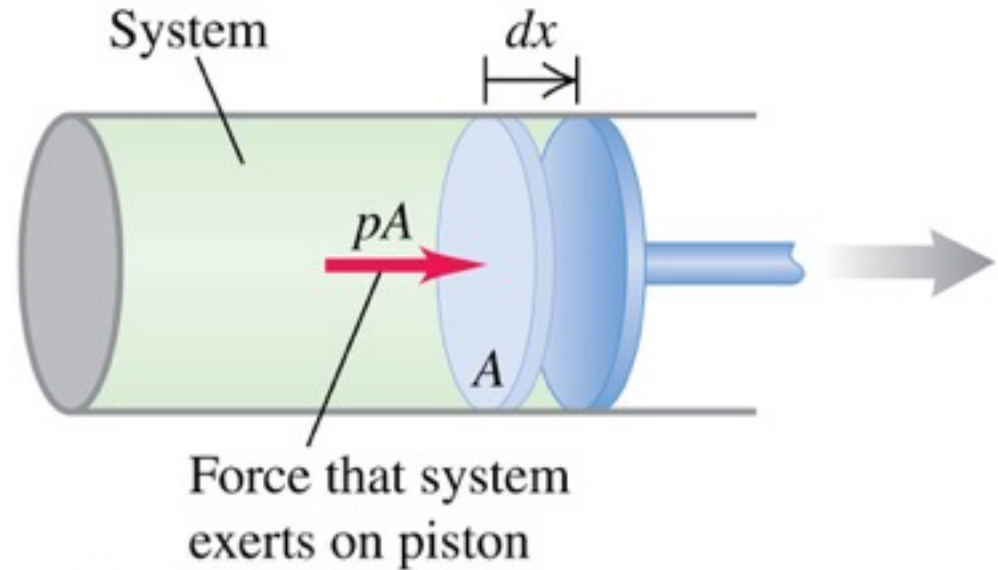
# Ch. 19 First Law of Thermodynamics

- In a *thermodynamic process*, changes occur in the state of the system.
- Careful of signs!
- $Q$  is positive when heat flows *into* a system.
- $W$  is the work done *by* the system, so it is positive for expansion.



# Work done during volume changes

- The infinitesimal work done by the system during the small expansion  $dx$  is  $dW = pA dx$ .
- In a finite change of volume from  $V_1$  to  $V_2$ :



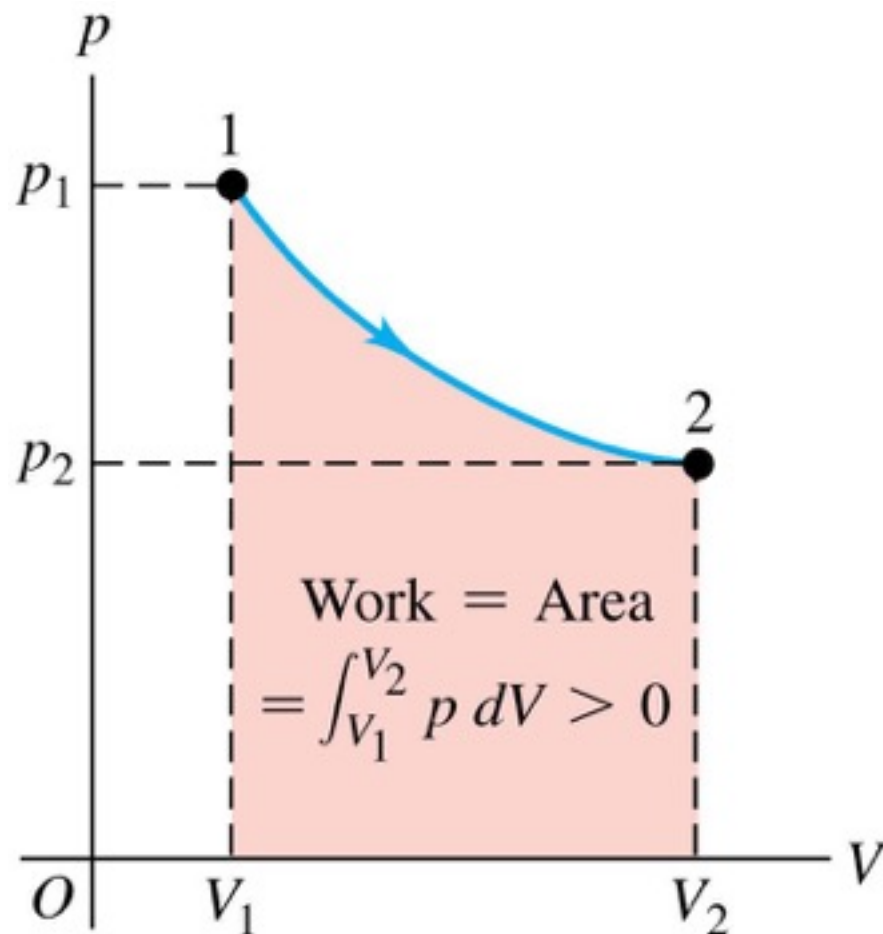
Work done in a volume change

$$W = \int_{V_1}^{V_2} p dV$$

Upper limit = final volume  
Integral of the pressure with respect to volume  
Lower limit = initial volume

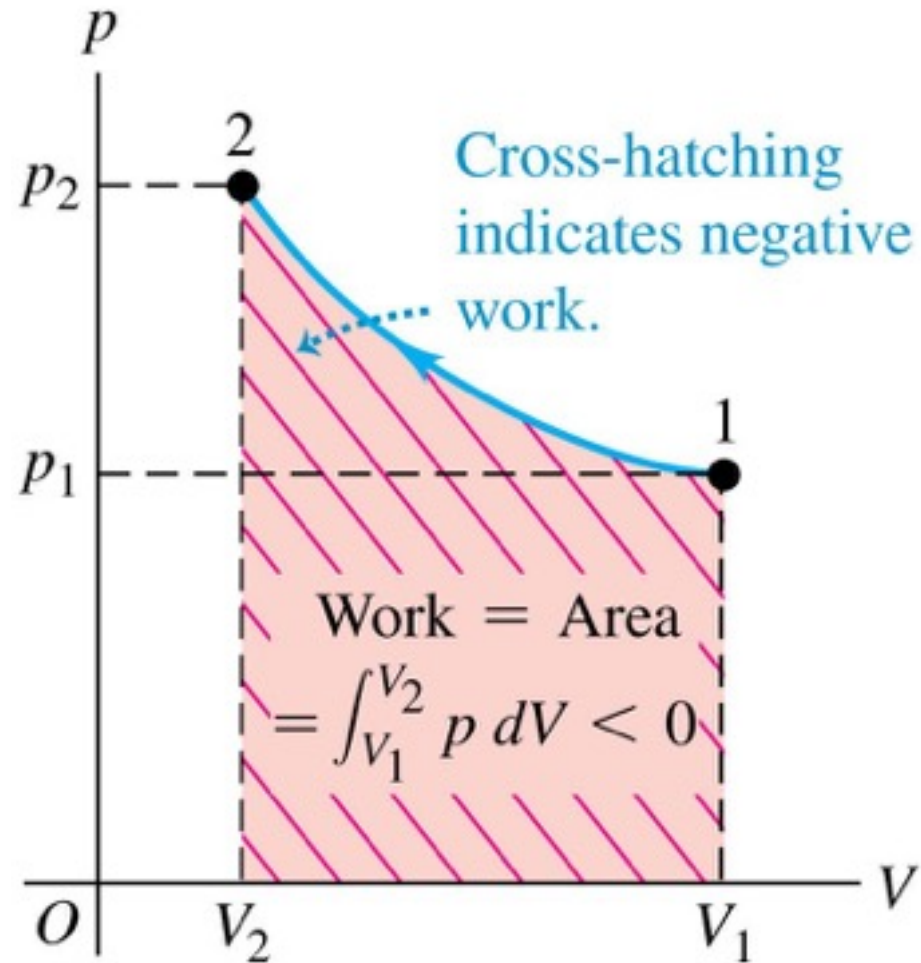
# Work on a $pV$ -diagram

- The work done equals the area under the curve on a  $pV$ -diagram.
- Shown in the graph is a system undergoing an **expansion** with varying pressure.



# Work on a $pV$ -diagram

- Shown in the graph is a system undergoing a **compression** with varying pressure.
- In this case the work is *negative*.



# First law of thermodynamics

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- The change in the internal energy  $U$  of a system is equal to the heat added minus the work done by the system:

First law of thermodynamics:

Internal energy change of thermodynamic system

$$\Delta U = Q - W$$

Heat added to system  $\rightarrow$   $\leftarrow$  Work done by system

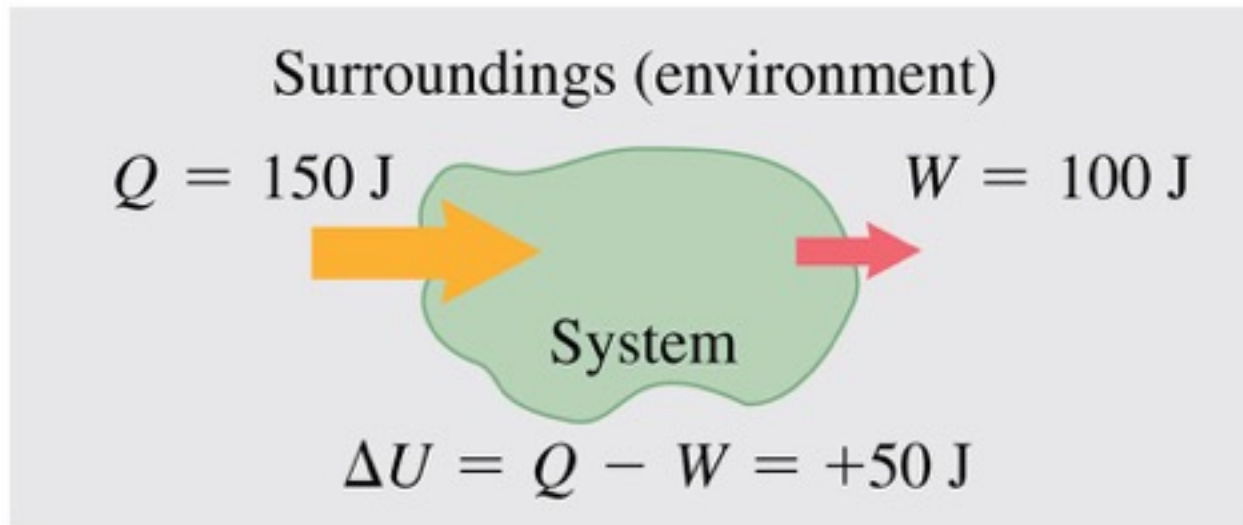
The diagram shows the equation  $\Delta U = Q - W$  centered on a light yellow background. Above the equation, the text "Internal energy change of thermodynamic system" is written in blue. Below the equation, "Heat added to system" is written in blue with a blue dotted arrow pointing from the text to the  $Q$  term. To the right, "Work done by system" is written in blue with a blue dotted arrow pointing from the  $W$  term to the text.

- The first law of thermodynamics is just a generalization of the conservation of energy.
- Both  $Q$  and  $W$  depend on the path chosen between states, but  $\Delta U$  is *independent of the path*.
- If the changes are infinitesimal, we write the first law as  $dU = dQ - dW$ .

# First law of thermodynamics

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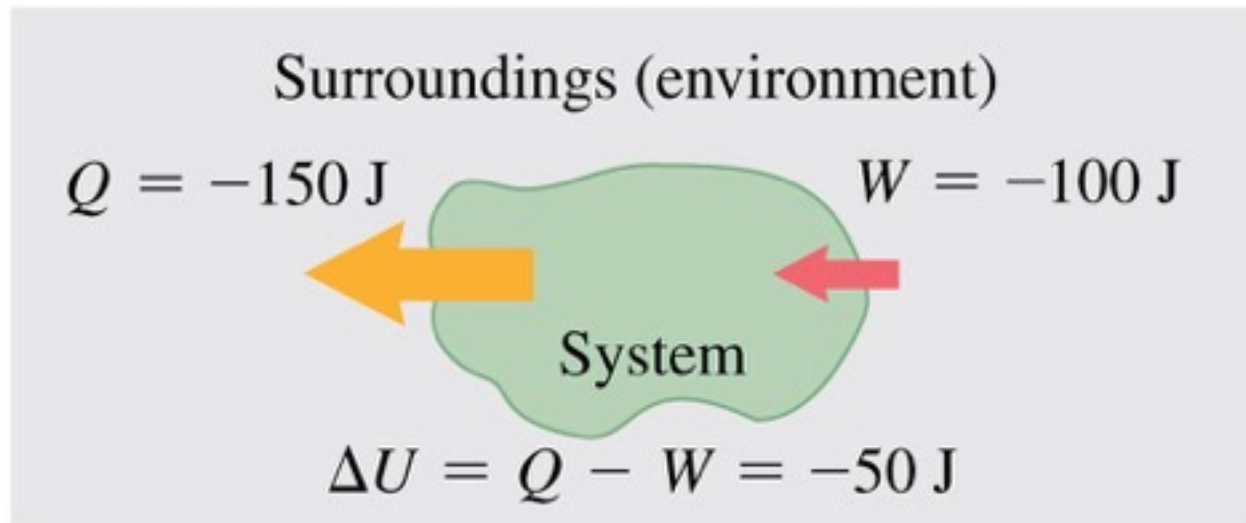
- In a thermodynamic process, the internal energy  $U$  of a system may *increase*.
- In the system shown below, more heat is added to the system than the system does work.
- So the internal energy of the system increases.



# First law of thermodynamics

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- In a thermodynamic process, the internal energy  $U$  of a system may *decrease*.
- In the system shown below, more heat flows out of the system than work is done.
- So the internal energy of the system decreases.

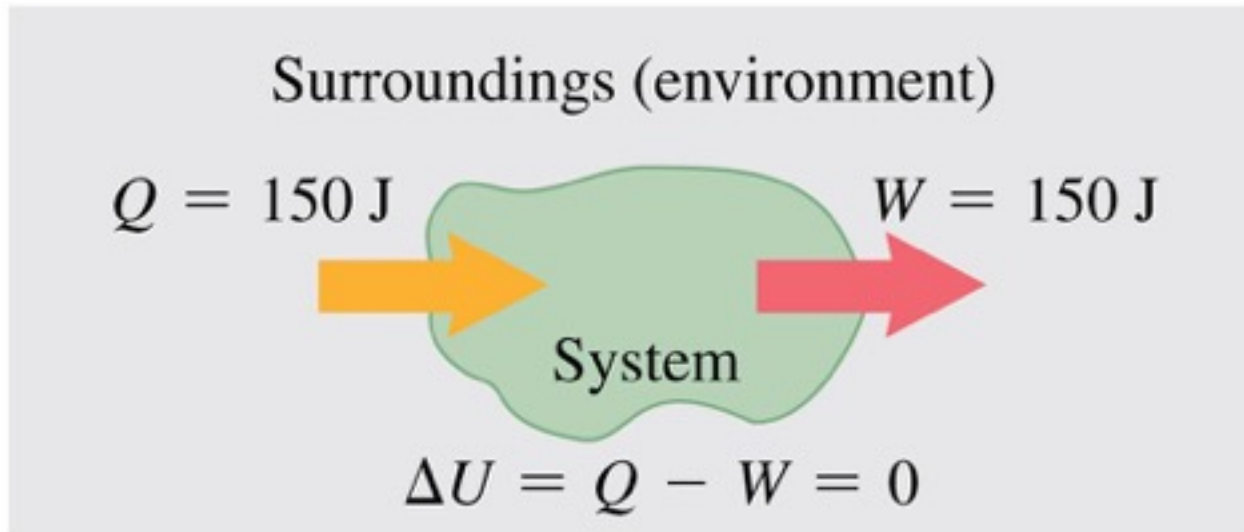




# First law of thermodynamics

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- In a thermodynamic process, the internal energy  $U$  of a system may *remain the same*.
- In the system shown below, the heat added to the system equals the work done by the system.
- So the internal energy of the system is unchanged.



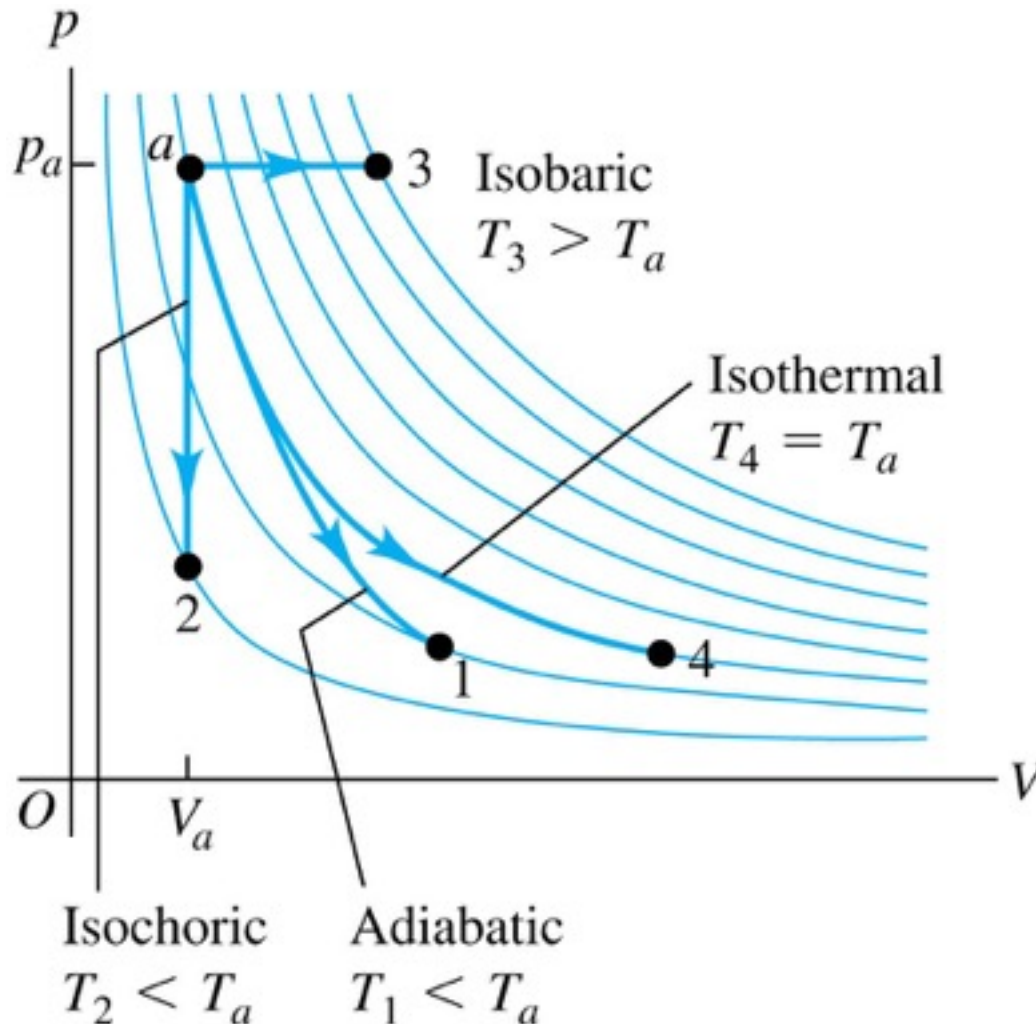
# Four kinds of thermodynamic processes

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- There are four specific kinds of thermodynamic processes that occur often in practical situations:
  - **Adiabatic:** No heat is transferred into or out of the system, so  $Q = 0$ . Also,  $U_2 - U_1 = -W$ .
  - **Isochoric:** The volume remains constant, so  $W = 0$ .
  - **Isobaric:** The pressure remains constant, so  $W = p(V_2 - V_1)$ .
  - **Isothermal:** The temperature remains constant.

# The four processes on a $pV$ -diagram

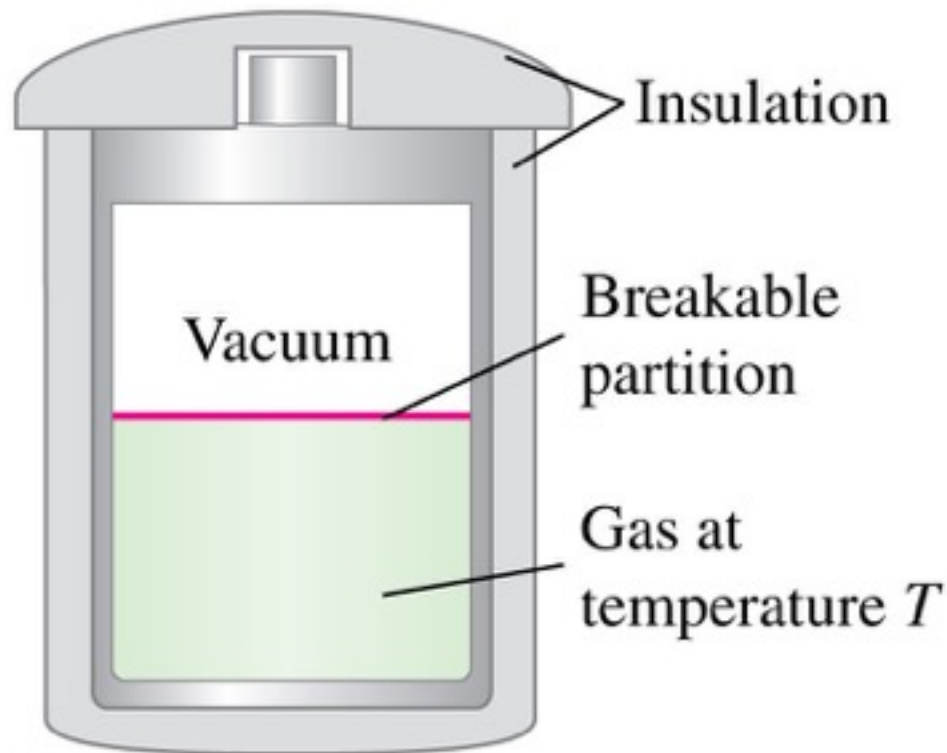
- Shown are the paths on a  $pV$ -diagram for all four different processes for a constant amount of an ideal gas, all starting at state  $a$ .



# Internal energy of an ideal gas

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- The internal energy of an ideal gas depends *only* on its *temperature*, not on its pressure or volume.
- The temperature of an ideal gas does *not* change during a free expansion.

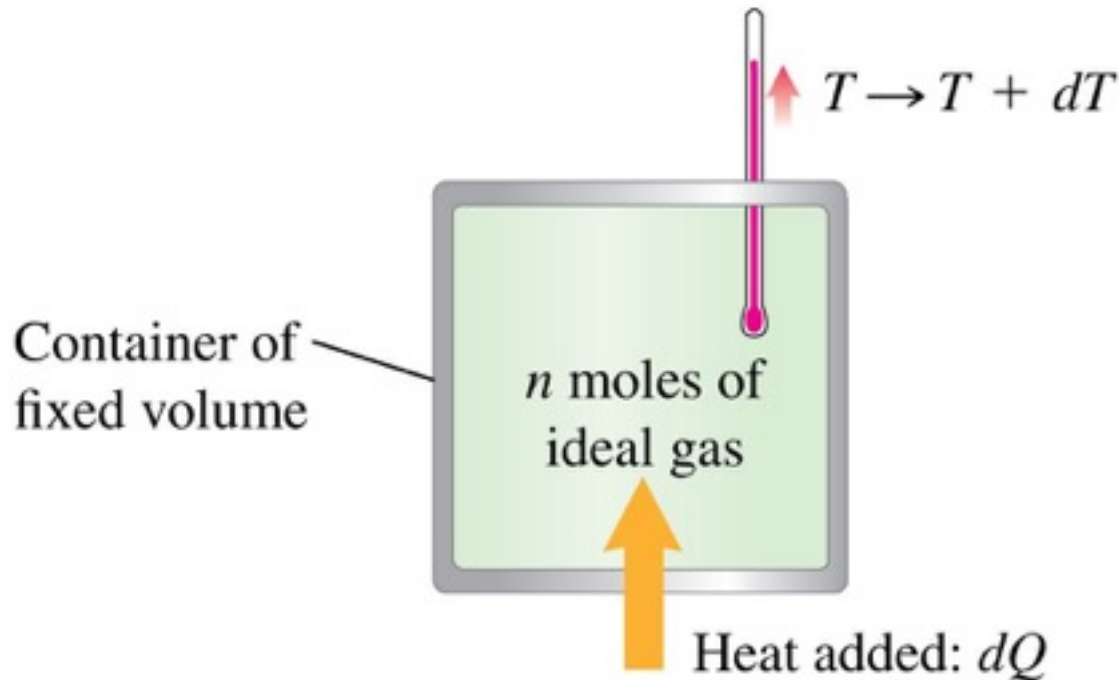


# Heat capacities of an ideal gas

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- $C_V$  is the molar heat capacity at constant volume.
- To measure  $C_V$ , we raise the temperature of an ideal gas in a rigid container with constant volume, ignoring its thermal expansion.

$$\text{Constant volume: } dQ = nC_V dT$$

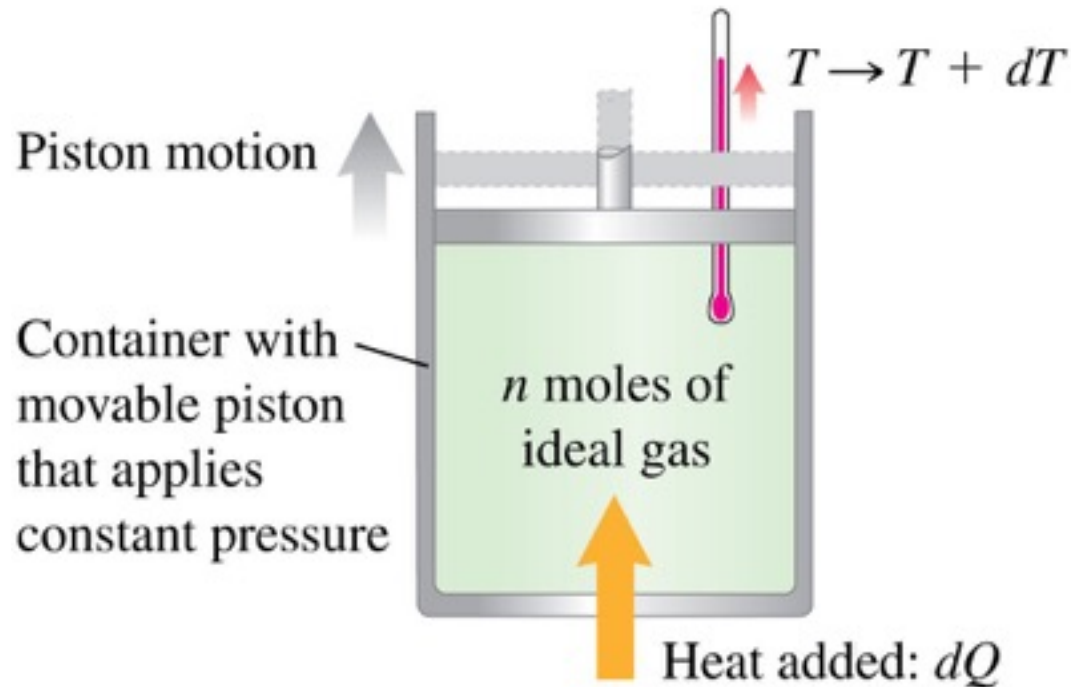


# Heat capacities of an ideal gas

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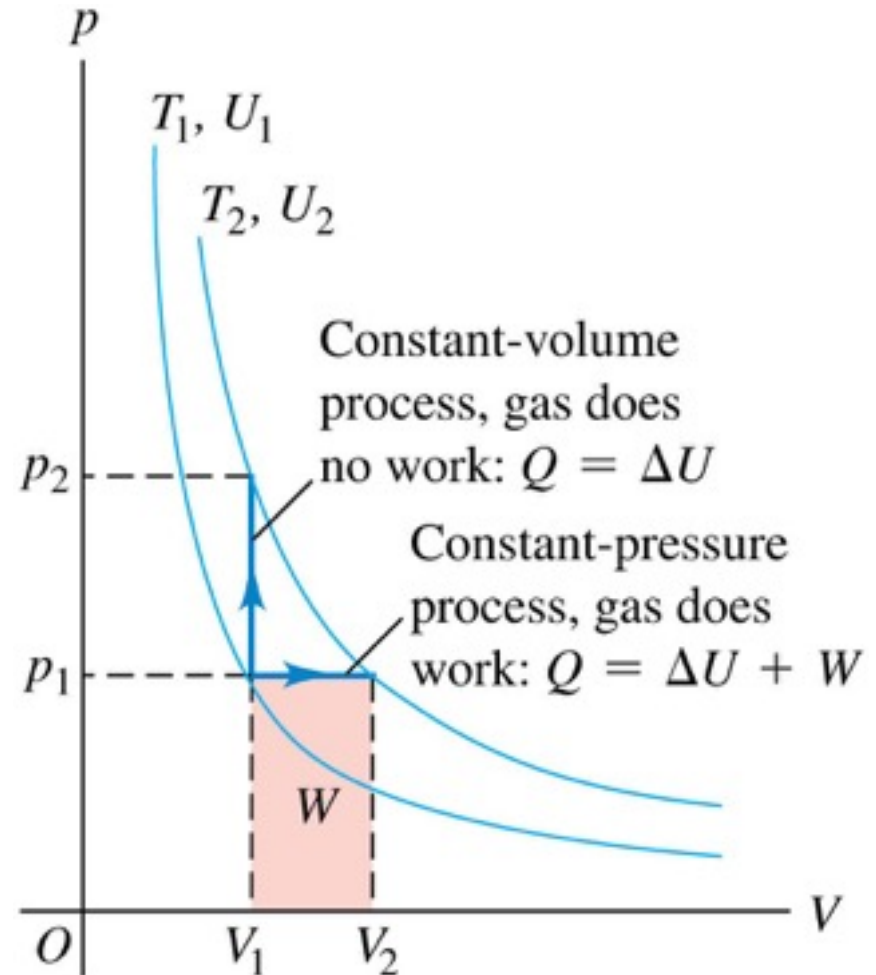
- $C_p$  is the molar heat capacity at constant pressure.
- To measure  $C_p$ , we let the gas expand just enough to keep the pressure constant as the temperature rises.

$$\text{Constant pressure: } dQ = nC_p dT$$



## Relating $C_p$ and $C_V$ for an ideal gas

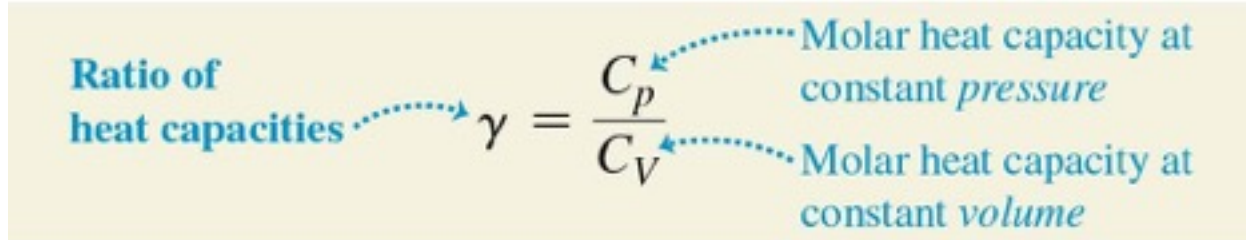
- To produce the same temperature change, more heat is required at constant pressure than at constant volume since  $\Delta U$  is the same in both cases.
- This means that  $C_p > C_V$ .
- $C_p = C_V + R$ .
- $R$  is the gas constant  
 $R = 8.314 \text{ J/mol} \cdot \text{K}$ .



# The ratio of heat capacities

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- The *ratio of heat capacities* is:



Ratio of heat capacities  $\gamma = \frac{C_p}{C_v}$

Molar heat capacity at constant *pressure*

Molar heat capacity at constant *volume*

- For monatomic ideal gases,

$$\gamma = 1.67.$$

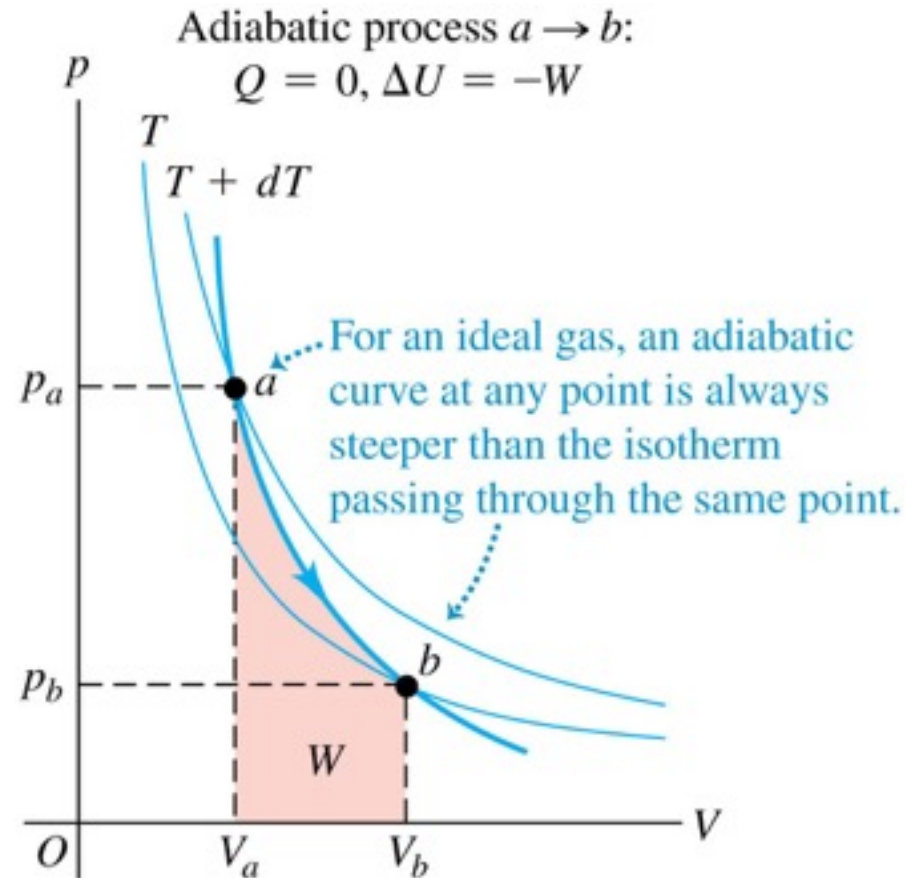
- For diatomic ideal gases,

$$\gamma = 1.40.$$



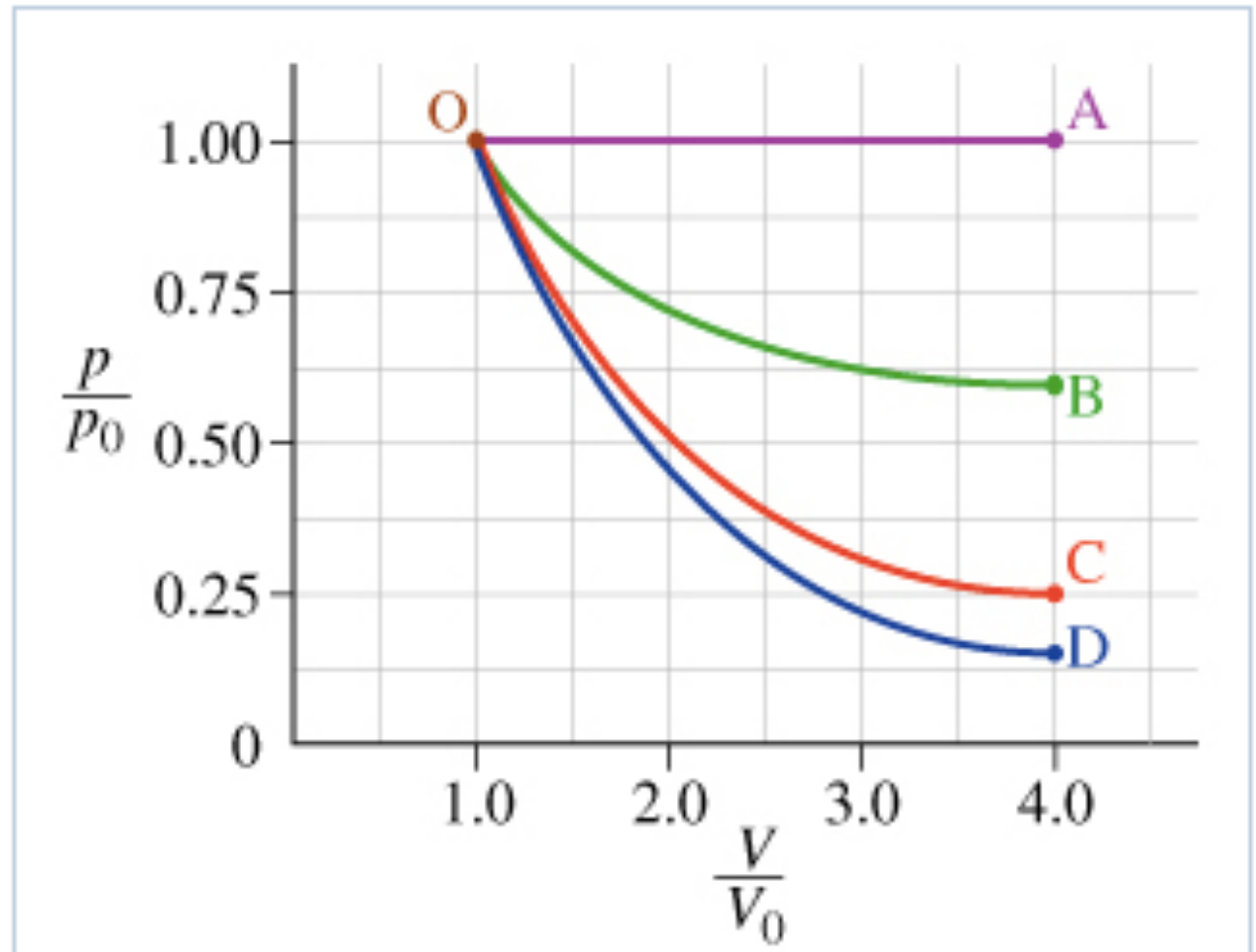
# Adiabatic processes for an ideal gas

- In an adiabatic process, no heat is transferred in or out of the gas, so  $Q = 0$ .
- Shown is a  $pV$ -diagram for an adiabatic expansion.
- As the gas expands, it does positive work  $W$  on its environment, so its internal energy decreases, and its temperature drops.



- Note that an adiabatic curve at any point is always steeper than an isotherm at that point.

# Expansion of a gas



- *Adiabatic*: No heat is added or removed during the expansion.
- *Isobaric*: The pressure remains constant during the expansion.
- *Isothermal*: The temperature remains constant during the expansion.

# Clicker question

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- You have 10 moles of a monatomic gas, with an initial volume  $V_i$ . You then compress the gas to half the initial volume in two ways:
- A. ISOTHERMAL compression
- B. ADIABATIC compression
  
- Q: In which process, A or B, is the final pressure of the gas higher?