



ATMOS 5130

Lecture 9

- Enthalpy
- Conservation Property
- The Second Law and Its Consequences
  - Entropy

# CLASS Presentation

- Form group of 2 students
- Present ~20 minute presentation (~ 10 minute each person)
- Focus on topics found within “In Practice” section of your textbook.
- Your research should go beyond the textbook.
- Be Creative! Props

# Examples of material to present

- Hygrometer
- Wind Chill
- Radiosonde
- Skew-T diagram
- Psychrometer
- Heat Index
- Aneroid Barometer
- Altimeter
- Subsidence inversion

# Suggestions for effective display

- **Keep it simple!** Fancy designs or color shifts make important material hard to read. Less is more.
- Use at least a 24-point font so everyone in the room can read your material.
- Try to limit the material to eight lines per slide, and keep the number of words to a minimum. **Keep it simple.**
- Limit the tables to four rows/columns for readability. Sacrifice content for legibility. Large tables can be displayed more effectively as a graph.
- Use easily read fonts.
- Don't fill up the slide - the peripheral material may not make it onto the display screen.
- **Identify the journal when you give references.**
- Finally, always preview your presentation. You will look foolish if symbols that looked OK in a WORD document didn't translate into anything readable in POWERPOINT.

# First Law of Thermodynamics

- Special Case of the Law of Conservation of Energy
- Energy stored by the parcel is its **internal energy**
  - Pressure volume work done by the system = reduction in internal energy + heat supplied by the environment
  - Pressure volume work done on the system = increase in internal energy + heat transferred to the environment

$$\delta w = -du + \delta q$$

Where  $\delta w$  = increment of **work** (per unit mass)  
 $du$  = change in the **internal energy** (per unit mass)  
 $\delta q$  = increment of **heat energy** (per unit mass)

Review

Recall:  $\delta w \equiv p d\alpha$

$$\delta q = -du + p d\alpha$$

# First Law of Thermodynamics

Review

$$\delta q = c_v dT + p d\alpha$$

$$\delta q = c_p dT - \alpha dp$$

# Enthalpy – Sensible Heat

- Accounts for both the gain in internal energy of the parcel and the mechanical work done in displacing the surrounding air as it expands

# Enthalpy – Sensible Heat

Specific Enthalpy – intensive units

$$h \equiv u + p\alpha$$

Internal energy      Mechanical work

$$dh \equiv du + d(p\alpha) = du + \alpha dp + p d\alpha$$

$$\delta q = du + p d\alpha$$

Substitute in 1<sup>st</sup> law of Thermo

$$dh - \alpha dp = du + p d\alpha$$

$$\delta q = dh - \alpha dp$$

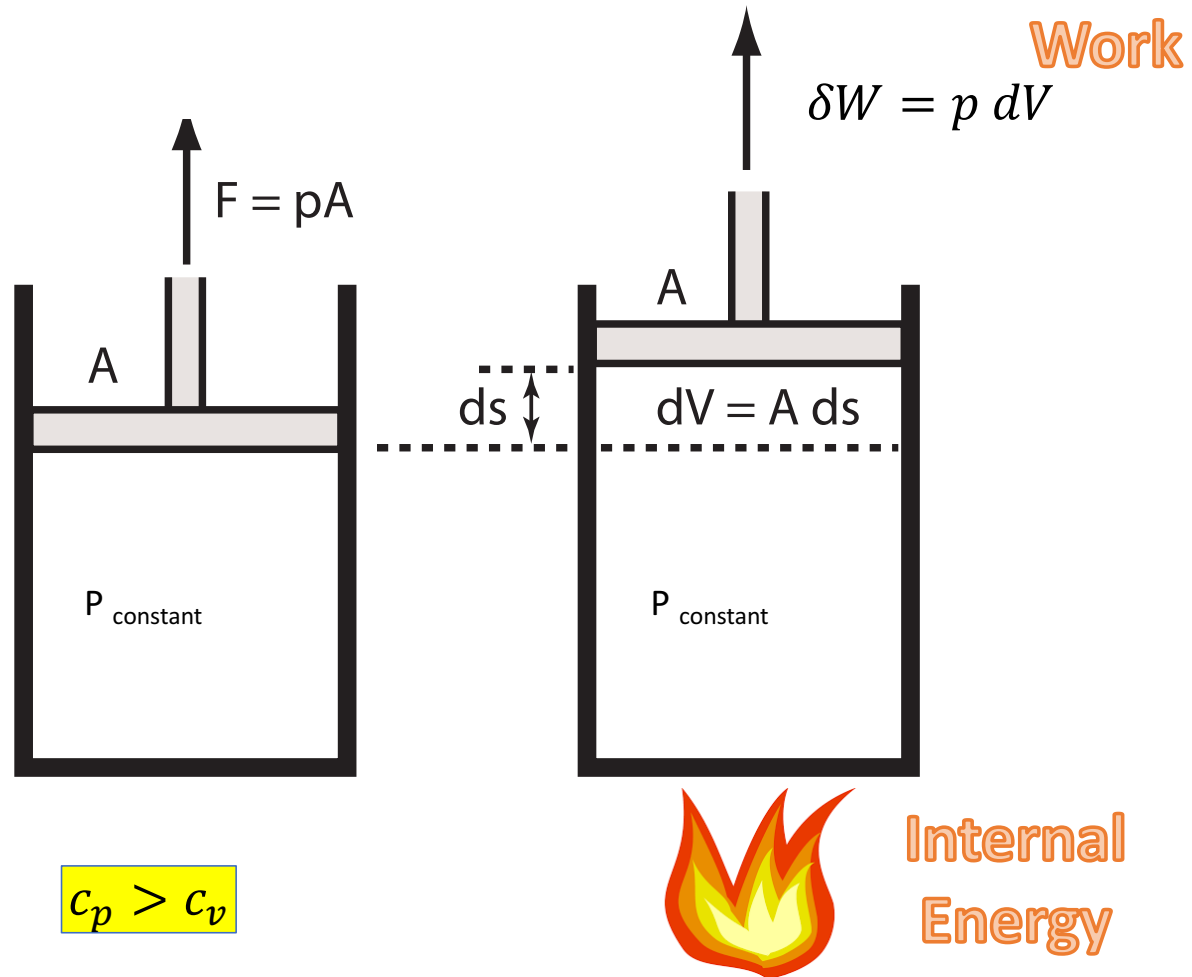
$$\delta q = c_p dT - \alpha dp$$

Compare to 1<sup>st</sup> law form

$$dh \equiv c_p dT$$



Heat Capacity at Constant Pressure = Isobaric case



# Special Cases of the First Law

$$\delta q = c_v dT + p d\alpha$$
$$\delta q = c_p dT - \alpha dp$$

Review

Isobaric Process: ( $dp = 0$ )

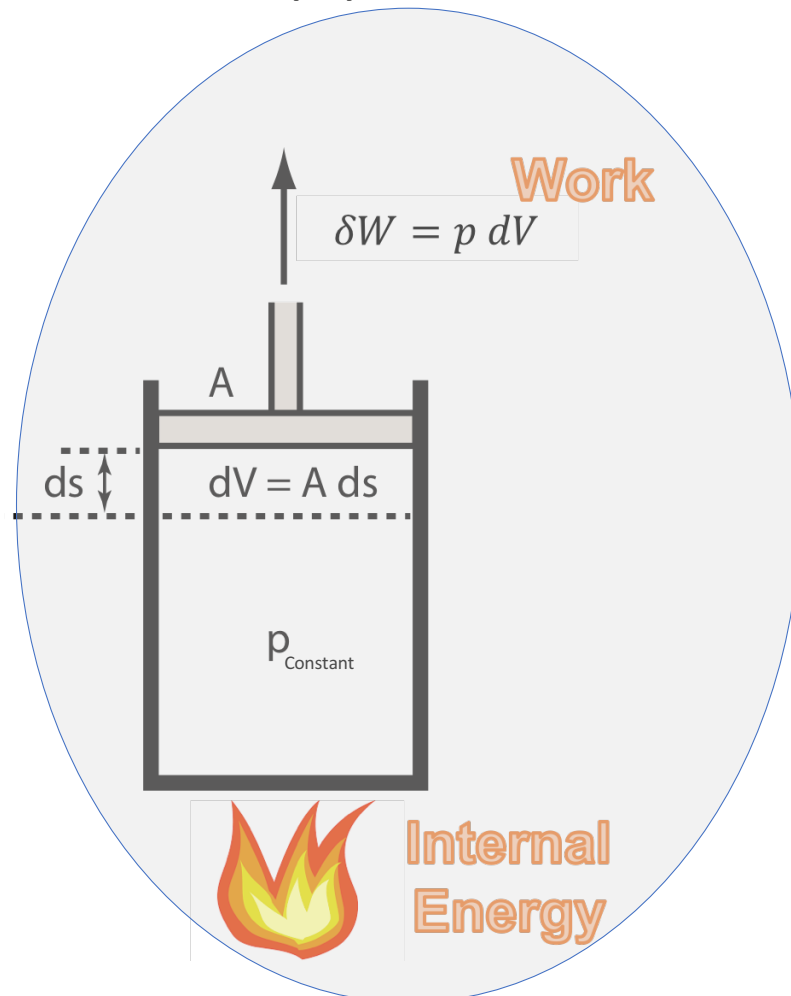
$$\delta q = c_p dT$$

Isothermal Process

Isochoric Process

Adiabatic Process

# Enthalpy – Sensible Heat



$$dh \equiv c_p dT$$

Gain in heat at constant pressure =  
Gain in Enthalpy

# Enthalpy – Sensible Heat

- Accounts for both the gain in internal energy of the parcel and the mechanical work done in displacing the surrounding air as it expands

Specific Enthalpy – intensive units

$$h \equiv u + p\alpha$$

Internal energy

Mechanical work: “work required to make room for the parcel by displacing the surrounding atmosphere.”

# Enthalpy – Sensible Heat

Specific Enthalpy – intensive units

$$h \equiv u + p\alpha$$

Internal energy      Mechanical work

$$dh \equiv du + d(p\alpha) = du + \alpha dp + p d\alpha$$

$$\delta q = du + p d\alpha$$

Substitute in 1<sup>st</sup> law of Thermo  
 $dh - \alpha dp = du + p d\alpha$

$$\delta q = dh - \alpha dp$$

New form of 1<sup>st</sup> law in terms of enthalpy

# Dry Static Energy Conservation

$$\delta q = dh - \alpha dp$$

Substitute in the  
Hydrostatic relationship

$$dp = -\rho g dz$$

Assumes the  
environmental temp  
~ temp of parcel

$$\delta q = dh - \alpha (-\rho g dz)$$

Recall  $\alpha = \frac{1}{\rho}$  and  $d\phi = g dz$

$$\delta q = dh - \alpha (-\rho g dz)$$

$$\delta q = dh + d\phi = d(h + \phi)$$

$$\underline{h + \phi}$$

Dry Static  
Energy

Conserved in  
dry adiabatic  
situation

## Real World – Generalization to an irreversible process

What happens when the environmental temperature is not equal to parcel temperature?

- Not in Equilibrium – Positive or Negative Buoyancy
- Forces acting on the parcel as it is displaced, gains/loses kinetic energy (i.e. it accelerates).

$$\delta q = dh + d\phi + e_k$$

- Dissipates kinetic energy – stops accelerating
- That kinetic energy must heat the parcel or environment. How much to parcel and how much to environment? We don't know.

$$\delta q \geq dh + d\phi$$

# First Law applied to cyclic process

Recall:

$$q_{net} = w_{net} = \oint \delta q = \oint c_v dT + \oint p d\alpha \neq 0$$

Thus, heat is not a state variable!



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Thus, heat is not a state variable!

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## Specific Entropy

**NEW STATE VARIABLE**

$$\oint \frac{1}{T} \delta q = \oint \frac{1}{T} c_v dT + \oint \frac{1}{T} p d\alpha$$

$$\oint \frac{1}{T} \delta q = c_v \oint \frac{dT}{T} + R \oint \frac{1}{\alpha} d\alpha$$

Now all in state variables

$$\oint \frac{1}{T} \delta q = 0 \equiv ds$$

Dry Adiabats  
Isentropes

Lines of Constant Entropy

Potential  
Temperature

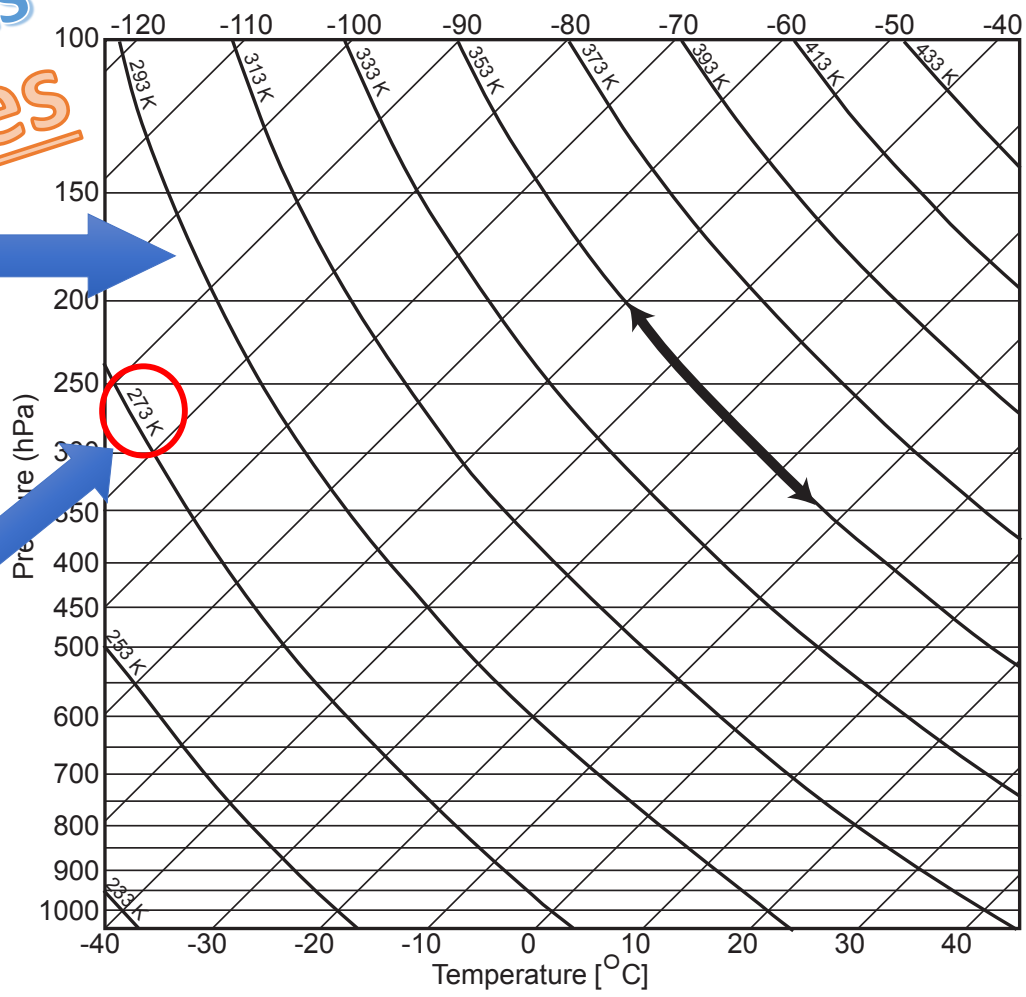


Fig. 5.3

First Law of Thermodynamics  
in term of Entropy

$$ds = c_v d \ln T + R d \ln \alpha$$
$$ds = c_p d \ln T + R d \ln p$$

$$\delta q = c_v dT + p d\alpha$$

$$\delta q = c_p dT - \alpha dp$$

Entropy = Energy dispersal

- Unconstrained energy spontaneously tends to disperse
- For example, energy spontaneously flows from hot to cold

# Second Law of Thermodynamics

- Within any isolated system that is not at equilibrium, the net effect of any active process is to always to increase the total entropy of the system.
- A state of equilibrium is reached when the total entropy of the system has achieved its maximum possible value. At this point, no further evolution of system state variables is possible

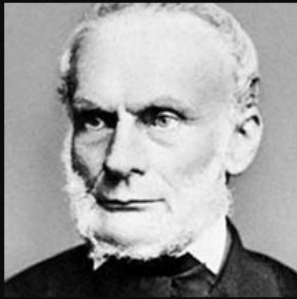
The increase of disorder or entropy is what distinguishes the past from the future, giving a direction to time.

— *Stephen Hawking* —

# Second Law of Thermodynamics

- Thermal equilibrium in a two body system is reached when both bodies have the same temperature

Rudolf Clausius



The entropy of an isolated system not in equilibrium will tend to increase over time, approaching a maximum value at equilibrium.

AZ QUOTES

## In Class problem

Calculate the change in air pressure if the specific entropy decreases by  $0.05 \text{ J g}^{-1}\text{K}^{-1}$  and the air temperature decreases by 5%.

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Calculate the change in air pressure if the specific entropy decreases by  $0.05 \text{ J g}^{-1}\text{K}^{-1}$  and the air temperature decreases by 5%.

$$ds = c_p d \ln T + R d \ln p$$

$$\ln \frac{p_f}{p_i} = \frac{c_p}{R} \ln\left(\frac{0.95T_i}{T_i}\right) - \frac{ds}{R}$$

$$\ln \frac{p_f}{p_i} = \frac{1005}{287} \ln\left(\frac{0.95T_i}{T_i}\right) - \frac{-50}{287}$$

$$p_f = 0.994 p_i$$

Pressure will decrease by a small amount of  $\sim 0.6\%$