

Specific heat and its features in ideal & real gases

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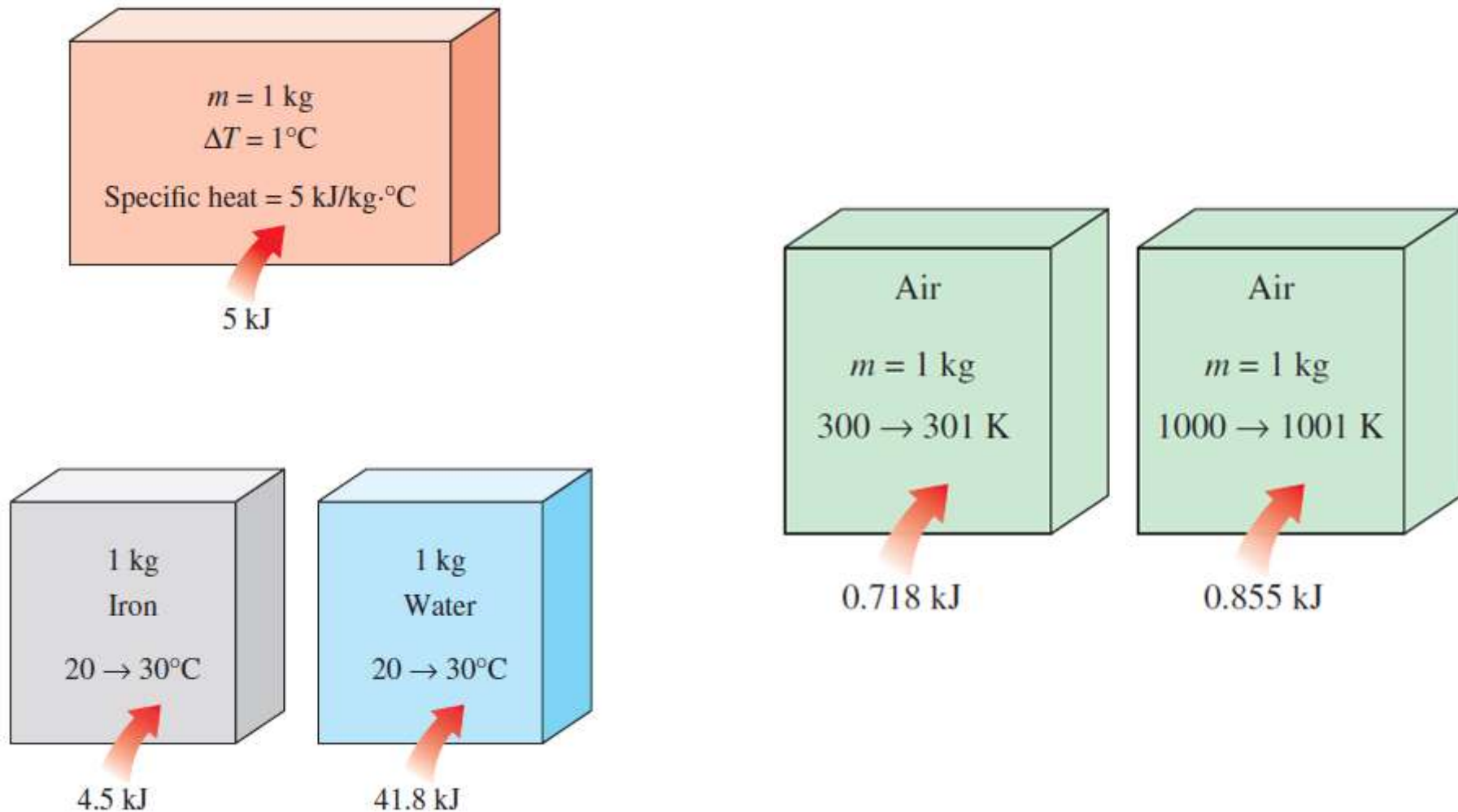
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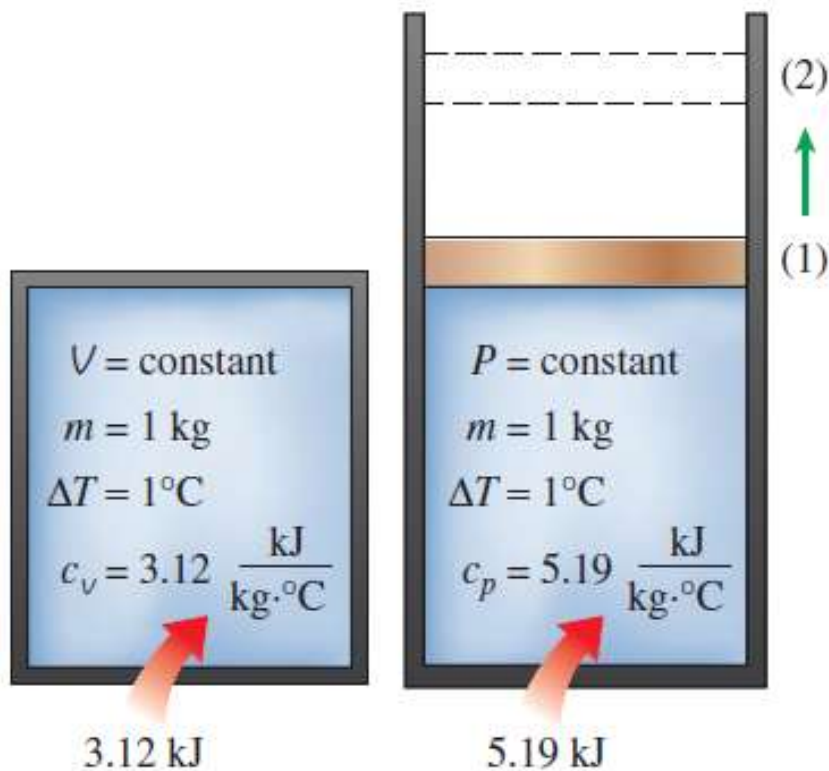
Cyclic processes, Enthalpy & Boundary work for $P=0$

- Work is a path fxn \rightarrow Net work from cyclic processes!
- Energy for processes at constant P: $H=U+PV$; $Q - W_{\text{other}} = \Delta H$
- Process at $P=0$

Specific heat is a material property



Specific heat at constant volume and pressure



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

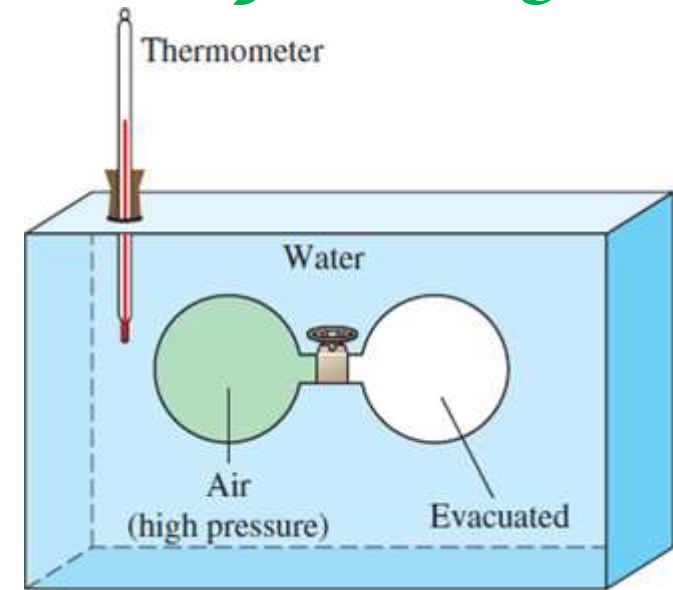
= the change in internal energy
with temperature at
constant volume

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

= the change in enthalpy with
temperature at constant
pressure

Joule's experiment & Specific heat of ideal gas

- No boundary work, P & V varying; $Q=0$
- T remains constants & hence, $U(T)$
- Ideal gas: $H=U(T)+PV=U(T)+RT=H(T)$
- Of course U is $U(T)$ -Degrees of freedom of ideal gas!

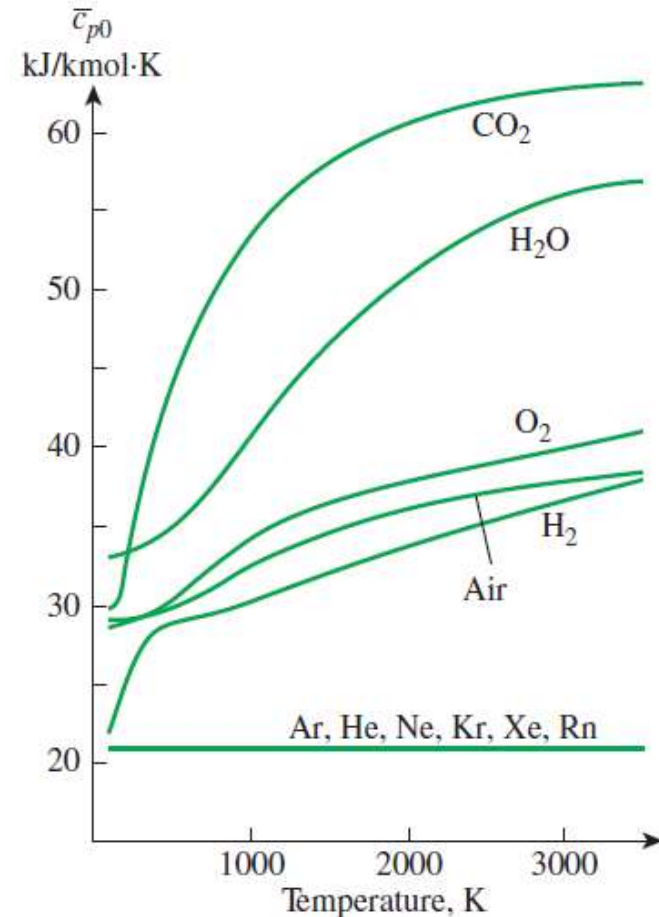


$$\Delta u = u_2 - u_1 = \int_1^2 c_v(T) dT \qquad \Delta h = h_2 - h_1 = \int_1^2 c_p(T) dT$$

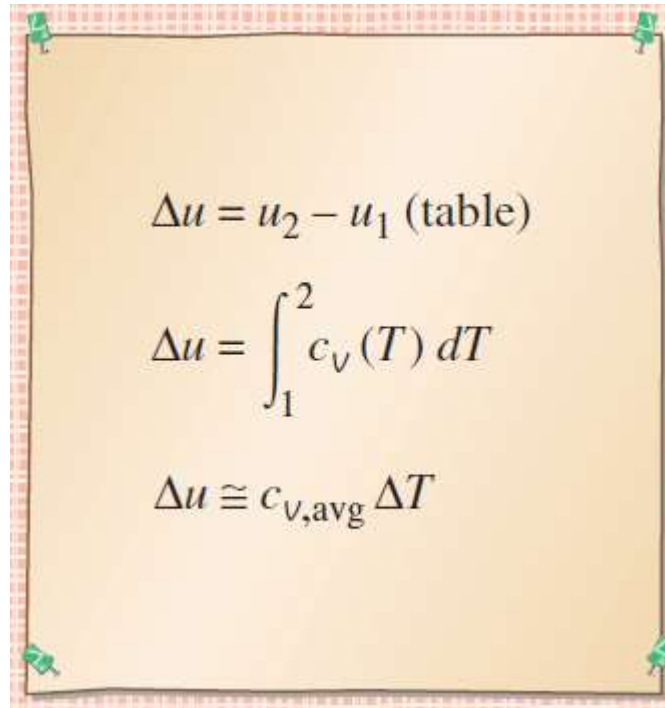
Specific heat of real gases

- Statistical thermodynamics provides rationale for variation
- Data from charts & tables are used for getting U & H
- Choose an arbitrary reference state and integrate from the reference state

Air		
T, K	$u, kJ/kg$	$h, kJ/kg$
0	0	0
•	•	•
•	•	•
300	214.07	300.19
310	221.25	310.24



Three approaches to ΔU & ΔH


$$\Delta u = u_2 - u_1 \text{ (table)}$$
$$\Delta u = \int_1^2 c_v(T) dT$$
$$\Delta u \cong c_{v,\text{avg}} \Delta T$$

Specific heat relationships of ideal gases

$$\begin{aligned}h &= u + RT, \\dh &= du + R dT \\dh &= c_p dT \text{ and } du = c_v dT\end{aligned}$$



The relationship between c_p , c_v and R

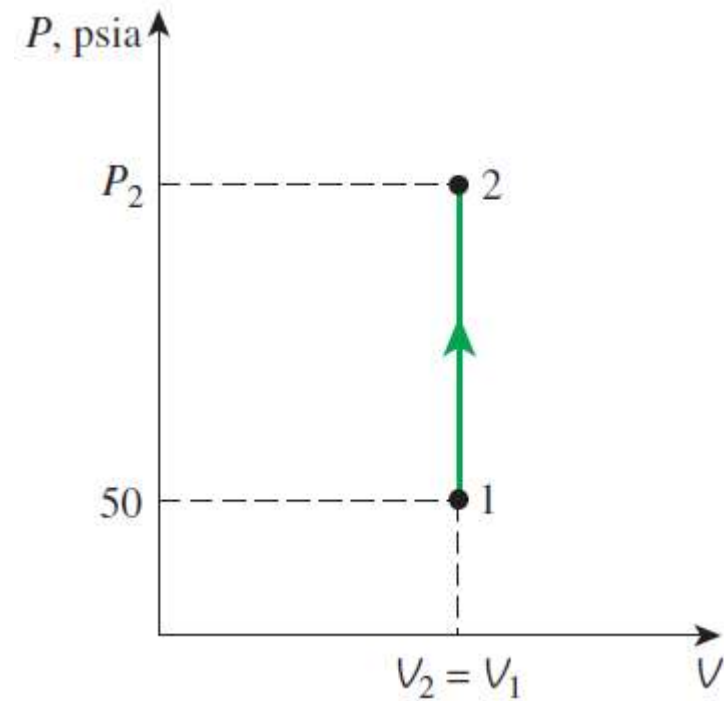
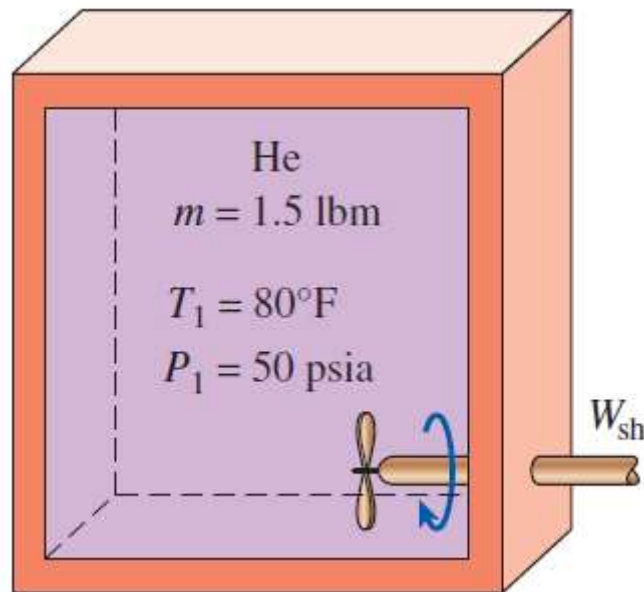
$$c_p = c_v + R \quad (\text{kJ/kg} \cdot \text{K})$$

On a molar basis

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

$$k = \frac{c_p}{c_v} \quad \text{Specific heat ratio}$$

Stirring & increase in T & P



$$\underbrace{E_{\text{in}} - E_{\text{out}}}_{\text{Net energy transfer by heat, work, and mass}} = \underbrace{\Delta E_{\text{system}}}_{\text{Change in internal, kinetic, potential, etc., energies}}$$

$$W_{\text{sh,in}} = \Delta U = m(u_2 - u_1) = mc_{v,\text{avg}}(T_2 - T_1)$$