

6. The Energy Equation

6.1 Derivation

- From the second law of thermodynamics:

$$Tds = dQ = dU + pdV$$

Heat change
Internal energy change

- Write this equation for a small element of mass δm with

$$dQ = \delta m dq$$

$$dU = \delta m de$$

$$dV = \delta m d\left(\frac{1}{\rho}\right) \leftarrow \text{Volume of unit mass}$$

- Where

$$e = \frac{p}{(\gamma - 1)\rho} \quad \text{Internal energy per unit mass} \quad (6.1)$$

$$\gamma = \frac{c_p}{c_v} \quad \text{Ratio of specific heats}$$

- And s is the entropy per unit mass
- If L is the sum of sources and sinks of energy, we can divide by dt to get

$$\rho T \frac{ds}{dt} = \rho \left[\frac{de}{dt} + p \frac{d}{dt} \left(\frac{1}{\rho} \right) \right] = -L$$

- We can write this more simply using mass conservation:

$$\rho \frac{de}{dt} + p\rho \left(\frac{-1}{\rho^2} \right) \frac{d\rho}{dt} = -L$$

$$\Rightarrow \rho \frac{de}{dt} - \frac{p}{\rho} \frac{d\rho}{dt} = -L \quad (6.2)$$

$$\Rightarrow \rho \frac{de}{dt} + p \underline{\nabla} \cdot \underline{u} = -L \quad (6.3)$$

- Now take \underline{u} . [equation of motion (5.1)] - the “mechanical energy equation” to get

$$\rho \frac{d}{dt} \left(\frac{1}{2} u^2 \right) = -\underline{u} \cdot \underline{\nabla} p + \underline{u} \cdot \rho \underline{g} \quad (6.4)$$

- Add this to (6.3) to get

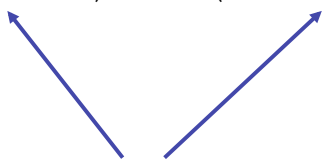
$$\begin{aligned} \rho \frac{d}{dt} \left(\frac{1}{2} u^2 + e \right) &= -L - p \underline{\nabla} \cdot \underline{u} - \underline{u} \cdot \underline{\nabla} p + \underline{u} \cdot \rho \underline{g} \\ &= -L - \underline{\nabla} \cdot p \underline{u} + \underline{u} \cdot \rho \underline{g} \end{aligned} \quad (6.5)$$

- Aside: for a scalar A:

$$\begin{aligned}
 \rho \frac{dA}{dt} &= \frac{d(\rho A)}{dt} - A \frac{d\rho}{dt} \\
 &= \frac{d(\rho A)}{dt} + \rho A \underline{\nabla} \cdot \underline{u} \\
 &= \frac{\partial(\rho A)}{\partial t} + \underline{u} \cdot \underline{\nabla}(\rho A) + \rho A \underline{\nabla} \cdot \underline{u} \\
 &= \frac{\partial(\rho A)}{\partial t} + \underline{\nabla} \cdot (\rho A) \underline{u}
 \end{aligned}$$

- And so writing g in terms of the gravitational potential ψ the energy equation (6.5) becomes:

$$\frac{\partial}{\partial t} \left(\frac{1}{2} \rho u^2 + \rho e \right) + \underline{\nabla} \cdot \left(\frac{1}{2} \rho u^2 + \rho e \right) \underline{u} = -L - \underline{\nabla} \cdot p \underline{u} - \underline{u} \cdot \rho \underline{\nabla} \psi \tag{6.6}$$



 “ρA”

- But we can write

$$\begin{aligned}
 \underline{\nabla} \cdot (\rho \psi) \underline{u} &= (\rho \psi) \underline{\nabla} \cdot \underline{u} + \underline{u} \cdot \underline{\nabla}(\rho \psi) \\
 &= (\rho \psi) \underline{\nabla} \cdot \underline{u} + \psi \underline{u} \cdot \underline{\nabla} \rho + \rho \underline{u} \cdot \underline{\nabla} \psi \\
 &= \psi (\rho \underline{\nabla} \cdot \underline{u} + \underline{u} \cdot \underline{\nabla} \rho) + \rho \underline{u} \cdot \underline{\nabla} \psi \\
 &= \psi \left(-\frac{\partial \rho}{\partial t} \right) + \rho \underline{u} \cdot \underline{\nabla} \psi
 \end{aligned}$$

- And so in a steady state, the energy equation becomes:

$$\underline{\nabla} \cdot \left(\frac{1}{2} \rho u^2 + \rho e + p + \rho \psi \right) \underline{u} = -L \quad (6.7)$$

- Where $\rho e + p = \frac{\gamma}{\gamma - 1} p$ is the *enthalpy*

Within some volume, the net effect (L) of the sources and sink of energy is equal (in a steady state) to the flux of energy through the surface of the volume.

6.2 Aside on equations of state

- In general, $p = p(\rho, T)$ and for an **ideal gas**

$$p = nk_B T$$

or

$$p = \frac{k_B}{m} \rho T$$

n = total number of particles per unit volume

m = mean particle mass

- where $\rho = mn$
- For a fully-ionised H plasma

$$n \equiv n_p + n_e = 2n_e$$

$$\rho \equiv n_p m_p + n_e m_e \approx n_e m_p$$

Barotropic Equations of State: $p(\rho)$

- Means that p can be written as a function of ρ only (e.g. for an ideal gas implies that ρ and T have some additional relation). E.g.:
- Isothermal $\Rightarrow T = \text{constant}$
so $p \propto \rho$
- For this to be a good approximation, require
 - Temperature for thermal equilibrium isn't very sensitive to the heating/cooling rate
 - In time-dependent problems, there is time for the system to reach this constant T thermal equilibrium

- Adiabatic $p = K\rho^\gamma$

- This is derived from the ideal gas laws on the assumption that there is no heat exchange with surroundings (i.e. no external heating/cooling) - changes in the internal energy result purely from pdV work. (Q4: use (6.2) to prove this)
- A fluid element behaves adiabatically if K is constant as the element's properties change. An isentropic fluid is one in which all the elements have the same value of K.

Answer to Q4

Start with the energy equation:

$$\rho \frac{de}{dt} - \frac{p}{\rho} \frac{d\rho}{dt} = -L$$

$$\frac{\rho}{\gamma - 1} \frac{d}{dt} \left(\frac{p}{\rho} \right) - \frac{p}{\rho} \frac{d\rho}{dt} = -L$$

$$\frac{1}{\gamma - 1} \left(\frac{dp}{dt} - \frac{p}{\rho} \frac{d\rho}{dt} - \frac{(\gamma - 1)p}{\rho} \frac{d\rho}{dt} \right) = -L$$

$$\frac{1}{\gamma - 1} \left(\frac{dp}{dt} - \frac{\gamma p}{\rho} \frac{d\rho}{dt} \right) = -L$$

$$\frac{\rho^\gamma}{\gamma - 1} \frac{d}{dt} \left(\frac{p}{\rho^\gamma} \right) = -L$$

Hence if L=0, then following the motion, $p=K\rho^\gamma$