

Lecture 11 141116

Il pdf delle lezioni puo' essere scaricato da

http://www.fisgeo.unipg.it/~fiandrin/didattica_fisica/cosmic_rays1617/

Connection with thermodynamics:

Ideal Gas Law

Isotropic gas of point particles in
Thermodynamic Equilibrium:

$$\frac{1}{2}m\overline{\sigma_x^2} = \frac{1}{2}m\overline{\sigma_y^2} = \frac{1}{2}m\overline{\sigma_z^2} = \frac{1}{6}m\overline{\sigma^2} = \frac{1}{2}k_bT$$

Temperature is defined in terms of kinetic energy of the thermal motions!

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Ideal Gas Law: in
terms of temperature T
and number-density n :
($\rho = nm = \mu n m_H$, $R = k_b / m_H$)

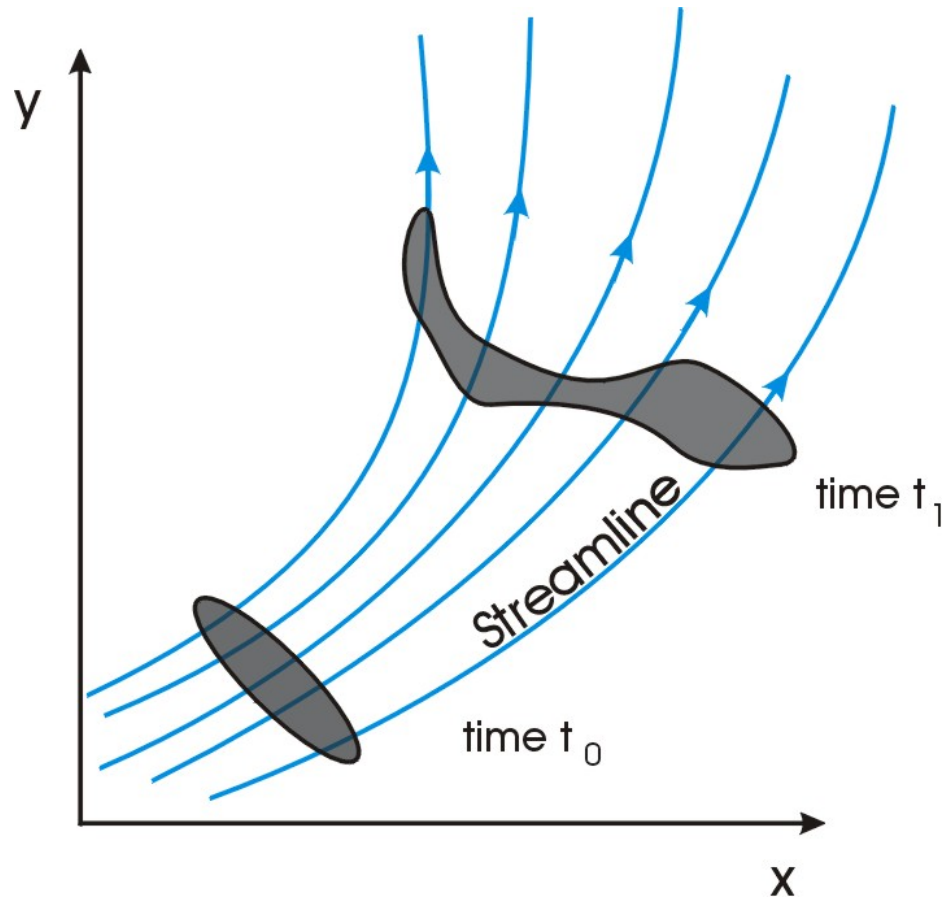
$$p = \frac{1}{3}\rho\overline{\sigma^2}$$



$$P(\rho, T) = nk_bT = \frac{\rho \mathcal{R} T}{\mu}$$

Digressione: fluidi viscosi

Density Changes and Mass Conservation

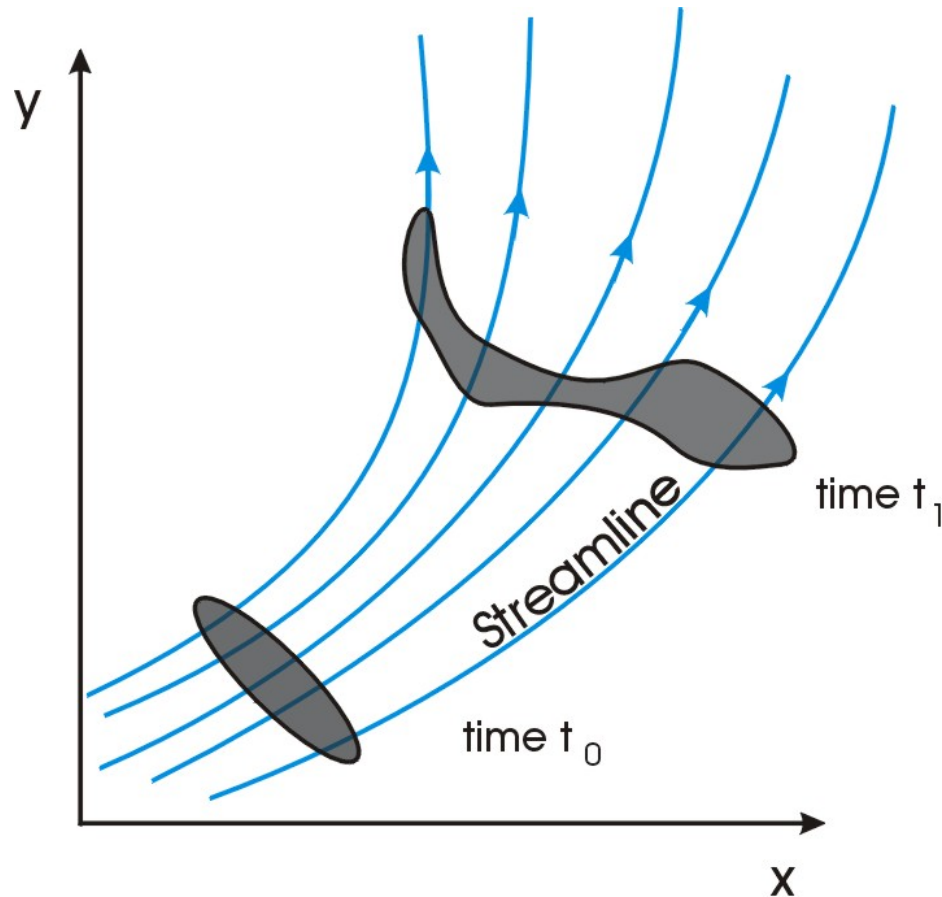


Two-dimensional example:

A fluid filament is deformed and stretched by the flow;

Its area changes, but the mass contained in the filament can NOT change

Density Changes and Mass Conservation



Two-dimensional example:

A fluid filament is deformed and stretched by the flow;

Its area changes, but the mass contained in the filament can NOT change

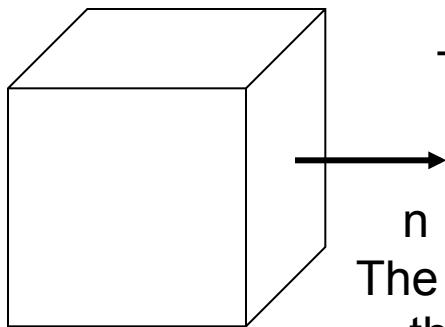
So: the mass density must change in response to the flow!

Mass conservation law

The mass cannot be created nor destroyed (in non-relativistic classical dynamics)

Therefore in a volume V the mass can change only because some of it leaves or enters the volume

The amount of mass through $d\mathbf{A}$ per time unit is $dF = \rho \vec{v} \cdot d\vec{A}$ Convention:
dF>0 if
outgoing



The flux

$$\left(\frac{dM}{dt}\right)_{out} = \int_S \rho \vec{v} \cdot d\vec{A}$$

The outgoing flux must be balanced by the change of mass in the volume

$$\frac{\partial M_V}{\partial t} = -\left(\frac{dM}{dt}\right)_{out}$$

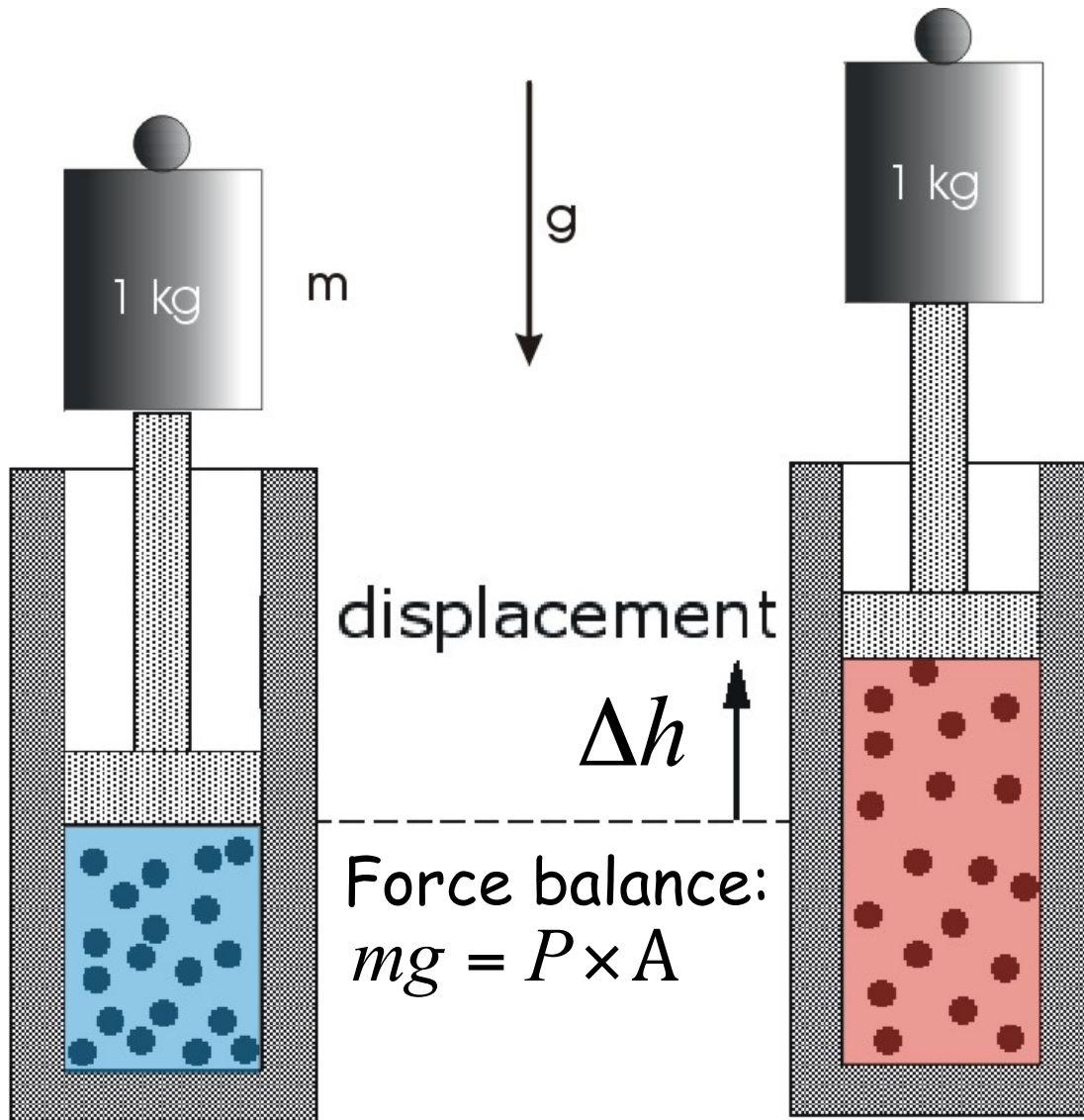
$$\frac{\partial M_V}{\partial t} = \frac{\partial}{\partial t} \left(\int_V \rho dV \right)$$

$$\frac{\partial}{\partial t} \left(\int_V \rho dV \right) = - \int_S \rho \vec{v} \cdot d\vec{A} = - \int_V \nabla \cdot (\rho \vec{v}) dV$$

$$\frac{\partial \rho}{\partial t} = -\nabla \cdot (\rho \vec{v})$$

$$\boxed{\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \vec{v}) = 0}$$

Thermodynamics



$$\Delta W = mg\Delta h$$

$$\Delta U_{\text{gas}} = \Delta Q - \Delta W$$

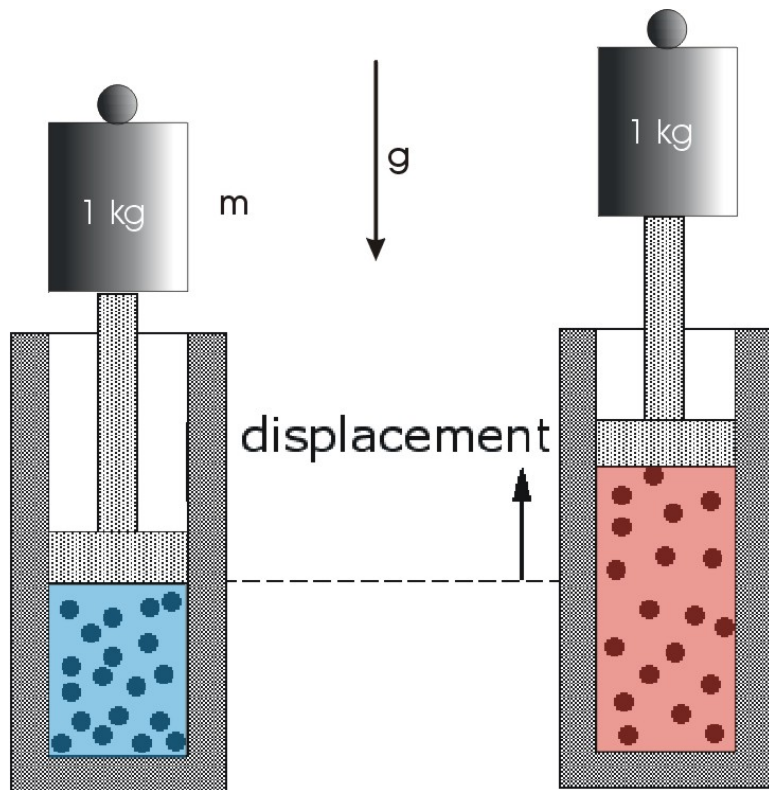
$$= \Delta Q - mg\Delta h$$

$$= \Delta Q - PA \Delta h$$

$$= \Delta Q - P \Delta V$$

First law of thermodynamics:

$$dU_{gas} = dQ - pdV$$



Change in internal energy
=
heat added by external sources
-
work done by gas

Entropy:

A measure of
'disorder'

Second Thermodynamic law:

$$dQ = TdS$$

$$dS \geq 0$$

$$\frac{dS}{dt} \geq 0$$

For an isolated system



The Adiabatic Gas Law: the behaviour of pressure

Thermodynamics:

$$dQ \equiv T dS = dU + P dV$$

Special case: **adiabatic change**

$$dQ = T dS = 0$$

U = internal energy, T = temperature, S = entropy
and V = volume

The Adiabatic Gas Law: the behaviour of pressure

Thermodynamics:

$$dQ \equiv T dS = dU + P dV$$

Special case: **adiabatic change**

$$dQ = T dS = 0$$

Gas of **point particles** of mass m :

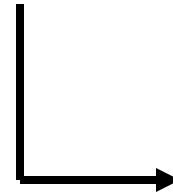
$$\text{Internal energy: } U = n V \times \left[\frac{1}{2} m (\overline{\sigma_x^2} + \overline{\sigma_y^2} + \overline{\sigma_z^2}) \right] = \frac{1}{2} \rho V \overline{\sigma^2}$$

Pressure:

$$P = \frac{1}{3} n m (\overline{\sigma_x^2} + \overline{\sigma_y^2} + \overline{\sigma_z^2}) = \frac{1}{3} \rho \overline{\sigma^2}$$

Thermal equilibrium:

$$\frac{1}{2}m\overline{\sigma_x^2} = \frac{1}{2}m\overline{\sigma_y^2} = \frac{1}{2}m\overline{\sigma_z^2} = \frac{1}{6}m\overline{\sigma^2} = \frac{1}{2}k_bT$$


$$P = \frac{\rho \mathcal{R} T}{\mu}, \quad U = \frac{3}{2} \frac{\rho \mathcal{R} T \mathcal{V}}{\mu}$$

Thermal equilibrium:

$$\frac{1}{2}m\overline{\sigma_x^2} = \frac{1}{2}m\overline{\sigma_y^2} = \frac{1}{2}m\overline{\sigma_z^2} = \frac{1}{6}m\overline{\sigma^2} = \frac{1}{2}k_bT$$

$$\downarrow \quad P = \frac{\rho \mathcal{R} T}{\mu}, \quad U = \frac{3}{2} \frac{\rho \mathcal{R} T \mathcal{V}}{\mu}$$

Adiabatic change:

$$dU + P d\mathcal{V} = 0 \longrightarrow d\left(\frac{3\rho \mathcal{R} T \mathcal{V}}{2\mu}\right) + \left(\frac{\rho \mathcal{R} T}{\mu}\right) d\mathcal{V} = 0$$

Adiabatic Gas Law: a polytropic relation

Thermal equilibrium: $\frac{1}{2}m\overline{\sigma_x^2} = \frac{1}{2}m\overline{\sigma_y^2} = \frac{1}{2}m\overline{\sigma_z^2} = \frac{1}{6}m\overline{\sigma^2} = \frac{1}{2}k_bT$

$$\downarrow \quad P = \frac{\rho \mathcal{R} T}{\mu}, \quad U = \frac{3}{2} \frac{\rho \mathcal{R} T \mathcal{V}}{\mu}$$

Adiabatic change:

$$dU + P d\mathcal{V} = 0 \longrightarrow d\left(\frac{3\rho \mathcal{R} T \mathcal{V}}{2\mu}\right) + \left(\frac{\rho \mathcal{R} T}{\mu}\right) d\mathcal{V} = 0$$

Chain rule for 'd' -operator:

$$d(f g) = (df) g + f (dg) \longrightarrow \frac{5}{3} P d\mathcal{V} + \mathcal{V} dP = 0.$$

(just like differentiation!)

$$\frac{dP}{P} + \frac{5}{3} \frac{d\mathcal{V}}{\mathcal{V}} = d \log (P \mathcal{V}^{5/3}) = 0$$

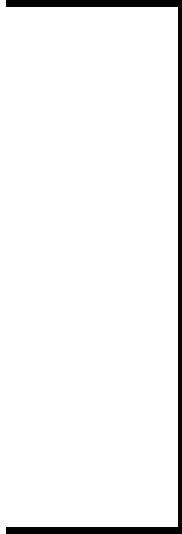
Adiabatic Gas Law: a polytropic relation

Adiabatic pressure change:

$$P \times \mathcal{V}^{5/3} = \text{constant}$$

For small volume:
mass conservation!

$$M = \rho \mathcal{V} = \text{constant}$$


$$P \rho^{-5/3} = \text{constant}$$

Specific Heat and Entropy

Specific Volume:
contains unit mass

$$\bar{v} \equiv \frac{1}{\rho}$$

Thermodynamics
of unit mass:

$$dq = T ds = de + P d\left(\frac{1}{\rho}\right)$$

Specific Heat and Entropy

Specific Volume
contains unit mass

$$\bar{V} \equiv \frac{1}{\rho}$$

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of a unit mass:

$$dq = T ds = de + P d\left(\frac{1}{\rho}\right)$$

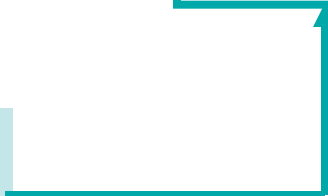
Specific energy e
and pressure P :

$$e \equiv \frac{3}{2} \frac{\mathcal{R}T}{\mu} = \frac{3}{2} \frac{k_b T}{m}, \quad P = \frac{\rho \mathcal{R} T}{\mu}$$

Specific heat coeff.
at constant volume

$$c_v = \frac{\partial e}{\partial T} = \frac{3}{2} \frac{k_b}{m}$$

ρ is kept constant!
 $d(1/\rho) = 0$



Specific Heat and Entropy

Specific Volume
contains unit mass

$$\bar{V} \equiv \frac{1}{\rho}$$

Thermodynamics
of a unit mass:

$$dq = T ds = de + P d\left(\frac{1}{\rho}\right)$$

Specific energy e
and pressure P :

$$e \equiv \frac{3}{2} \frac{\mathcal{R}T}{\mu} = \frac{3}{2} \frac{k_b T}{m}, \quad P = \frac{\rho \mathcal{R} T}{\mu}$$

Specific heat coeff.
at constant volume

$$c_v = \frac{\partial e}{\partial T} = \frac{3}{2} \frac{k_b}{m}$$

$$dq = d\left(e + \frac{P}{\rho}\right) - \frac{dP}{\rho}$$

$$c_p - c_v = \frac{k_b}{m} = \frac{\mathcal{R}}{\mu}$$

Specific heat coeff. at
constant pressure: $dP=0$

$$c_p = \frac{\partial(e + P/\rho)}{\partial T} = \frac{5}{2} \frac{k_b}{m}$$

$$\begin{aligned}
 dq &= c_v dT + \left(\frac{\rho \mathcal{R} T}{\mu} \right) d \left(\frac{1}{\rho} \right) \\
 &= c_v dT - \left(\frac{\mathcal{R} T}{\rho \mu} \right) d\rho \\
 &= c_v T \left[\frac{dT}{T} - \left(\frac{c_p}{c_v} - 1 \right) \frac{d\rho}{\rho} \right]
 \end{aligned}$$

Thermodynamic law for a unit mass,
rewritten in terms of specific heat coefficients

$$\begin{aligned}
 dq &= c_v dT + \left(\frac{\rho \mathcal{R} T}{\mu} \right) d \left(\frac{1}{\rho} \right) \\
 &= c_v dT - \left(\frac{\mathcal{R} T}{\rho \mu} \right) d\rho \\
 &= c_v T \left[\frac{dT}{T} - \left(\frac{c_p}{c_v} - 1 \right) \frac{d\rho}{\rho} \right]
 \end{aligned}$$

Definition specific entropy s

$$T ds = c_v T \left[\frac{dT}{T} - (\gamma - 1) \frac{d\rho}{\rho} \right]$$

$$s = c_v \log \left(\frac{T}{\rho^{\gamma-1}} \right) + \text{constant}$$

$$s = c_v \log (P \rho^{-\gamma}) + \text{constant}$$

γ is the specific heat ratio
 $= 5/3$ for ideal gas of point particles

$$\begin{aligned}
 dq &= c_v dT + \left(\frac{\rho \mathcal{R} T}{\mu} \right) d \left(\frac{1}{\rho} \right) \\
 &= c_v dT - \left(\frac{\mathcal{R} T}{\rho \mu} \right) d\rho \\
 &= c_v T \left[\frac{dT}{T} - \left(\frac{c_p}{c_v} - 1 \right) \frac{d\rho}{\rho} \right]
 \end{aligned}$$

Definition specific entropy s

$$T ds = c_v T \left[\frac{dT}{T} - (\gamma - 1) \frac{d\rho}{\rho} \right]$$

$$s = c_v \log \left(\frac{T}{\rho^{\gamma-1}} \right) + \text{constant}$$

$$\log T - (\gamma - 1) \log \rho = \text{constant}$$

with

$$\gamma \equiv \frac{c_p}{c_v} = \frac{5}{3}.$$

$$s = c_v \log (P \rho^{-\gamma}) + \text{constant}$$

Case of constant entropy
(adiabatic gas) : $ds = 0$

$$T \rho^{-(\gamma-1)} = \text{constant} \quad , \quad P \rho^{-\gamma} = \text{constant}$$

(Self-)gravity

$$\mathbf{f}_{\text{gr}} = \rho \mathbf{g} = -\rho \nabla \Phi$$
$$\mathbf{g}(\mathbf{x}, t) = -\nabla \Phi(\mathbf{x}, t) = -\begin{pmatrix} \frac{\partial \Phi}{\partial x} \\ \frac{\partial \Phi}{\partial y} \\ \frac{\partial \Phi}{\partial z} \end{pmatrix}$$

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

Self-gravity and Poisson's equation

Potential: two contributions!

$$\Phi(\boldsymbol{x}, t) = \Phi_{\text{ext}}(\boldsymbol{x}, t) + \Phi_{\text{self}}(\boldsymbol{x}, t)$$

Poisson equation for the potential associated with self-gravity:

$$\nabla^2 \Phi_{\text{self}}(\boldsymbol{x}, t) = 4\pi G \rho(\boldsymbol{x}, t)$$

Accretion flow around
Massive Black Hole

Self-gravity and Poisson's equation

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Poisson equation for Potential associated with self-gravity:

$$\nabla^2 \Phi_{\text{self}}(\boldsymbol{x}, t) = 4\pi G \rho(\boldsymbol{x}, t)$$

$$\nabla^2 \Phi \equiv \frac{\partial^2 \Phi}{\partial x^2} + \frac{\partial^2 \Phi}{\partial y^2} + \frac{\partial^2 \Phi}{\partial z^2}$$

Laplace operator

Summary: Equations describing ideal (self-)gravitating fluid

Equation of Motion:

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

Continuity Equation:
behavior of mass-density

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}) = 0$$

Ideal gas law
&
Adiabatic law:
Behavior of pressure
and temperature

$$P(\rho, T) = nk_b T = \frac{\rho \mathcal{R} T}{\mu}$$

$$P \rho^{-5/3} = \text{constant}$$

Poisson's equation: self-gravity

$$\nabla^2 \Phi_{\text{self}}(\mathbf{x}, t) = 4\pi G \rho(\mathbf{x}, t)$$

Conservative Form of the Equations

Aim: To cast all equations in the same *generic form*:

$$\frac{\partial}{\partial t} \begin{pmatrix} \text{density of} \\ \text{quantity} \end{pmatrix} + \nabla \cdot \begin{pmatrix} \text{flux of that} \\ \text{quantity} \end{pmatrix} = \begin{pmatrix} \text{external sources} \\ \text{per unit volume} \end{pmatrix}$$

Reasons:

1. Allows quick identification of conserved quantities
2. This form works best in constructing numerical codes for *Computational Fluid Dynamics*
3. Shock waves are best studied from a conservative point of view

Generic Form:

$$\frac{\partial}{\partial t} \begin{pmatrix} \text{density of} \\ \text{quantity} \end{pmatrix} + \nabla \cdot \begin{pmatrix} \text{flux of that} \\ \text{quantity} \end{pmatrix} = \begin{pmatrix} \text{external sources} \\ \text{per unit volume} \end{pmatrix}$$

Transported quantity is a scalar S , so flux \mathbf{F} must be a vector!

$$\frac{\partial S}{\partial t} + \nabla \cdot \mathbf{F} = q(\mathbf{x}, t)$$

Component form:

$$\frac{\partial S}{\partial t} + \left(\frac{\partial F_x}{\partial x} + \frac{\partial F_y}{\partial y} + \frac{\partial F_z}{\partial z} \right) = q$$

Generic Form:

$$\frac{\partial}{\partial t} \begin{pmatrix} \text{density of} \\ \text{quantity} \end{pmatrix} + \nabla \cdot \begin{pmatrix} \text{flux of that} \\ \text{quantity} \end{pmatrix} = \begin{pmatrix} \text{external sources} \\ \text{per unit volume} \end{pmatrix}$$

Transported quantity is a vector \mathbf{M} , so the flux must be a tensor \mathbf{T} .

$$\frac{\partial \mathbf{M}}{\partial t} + \nabla \cdot \mathbf{T} = \mathbf{Q}(\mathbf{x}, t)$$

Component form:

$$\frac{\partial}{\partial t} \begin{pmatrix} M_x \\ M_y \\ M_z \end{pmatrix} + \begin{pmatrix} \frac{\partial T_{xx}}{\partial x} + \frac{\partial T_{yx}}{\partial y} + \frac{\partial T_{zx}}{\partial z} \\ \frac{\partial T_{xy}}{\partial x} + \frac{\partial T_{yy}}{\partial y} + \frac{\partial T_{zy}}{\partial z} \\ \frac{\partial T_{xz}}{\partial x} + \frac{\partial T_{yz}}{\partial y} + \frac{\partial T_{zz}}{\partial z} \end{pmatrix} = \begin{pmatrix} Q_x \\ Q_y \\ Q_z \end{pmatrix}$$

The fact that the flux of a vector field is a rank 2 tensor can be understood as follows: the transported quantity is a vector with 3 arbitrary components, each of them can be transported in 3 independent directions \rightarrow so there are 3×3 independent quantities...exactly the number of components of a rank 2 tensor

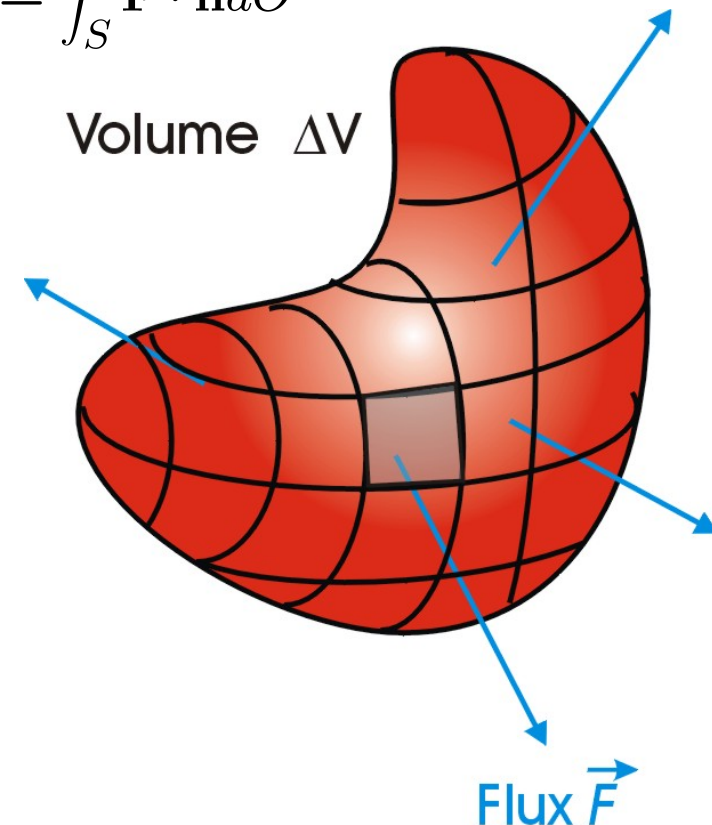
Integral properties: Stokes Theorem

$$\frac{\partial}{\partial t} \left(\begin{array}{c} \text{density of} \\ \text{quantity} \end{array} \right) + \nabla \cdot \left(\begin{array}{c} \text{flux of that} \\ \text{quantity} \end{array} \right) = \left(\begin{array}{c} \text{external sources} \\ \text{per unit volume} \end{array} \right)$$

Let integrate the equation over a volume V and use the Stokes theorem $\int_V \nabla \cdot \mathbf{F} dV = \int_S \mathbf{F} \cdot \mathbf{n} dO$

$$\frac{\partial}{\partial t} \left(\int_V dV S \right) = \int_V dV q(\mathbf{x}, t) - \oint_{\partial V} d\mathbf{O} \cdot \mathbf{F}$$

The integral relation states the amount of quantity S in a volume can change only due to sources in V or by a flux of S into or out from V



Examples: mass and momentum conservation

Mass conservation: already in conservation form!

Continuity Equation:
transport of the scalar ρ

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}) = 0$$

Excludes 'external mass sources' due to processes like two-photon pair production etc.

Examples: mass- and momentum conservation

Mass conservation: already in conservation form!

Continuity Equation:
transport of the scalar ρ

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}) = 0$$

Momentum conservation: transport of a vector!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

Algebraic Manipulation

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + \nabla \cdot (\rho \mathbf{V} \otimes \mathbf{V} + P \mathbf{I}) = -\rho \nabla \Phi$$

As advertised: *Algebraic Manipulation!*

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

Starting point: Equation of Motion

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}) = 0$$

$$\begin{aligned} \rho \frac{\partial \mathbf{V}}{\partial t} &= \frac{\partial(\rho \mathbf{V})}{\partial t} - \mathbf{V} \frac{\partial \rho}{\partial t} \\ &= \frac{\partial(\rho \mathbf{V})}{\partial t} + \mathbf{V} (\nabla \cdot (\rho \mathbf{V})) \end{aligned}$$

Use:

1. chain rule for differentiation
2. continuity equation for density

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}) = 0$$

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$$\frac{\partial(\rho \mathbf{V})}{\partial t} + (\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = -\nabla P - \rho \nabla \Phi$$

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + \boxed{(\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V}} = -\nabla P - \rho \nabla \Phi$$

$$\boxed{(\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = \nabla \cdot (\rho \mathbf{V} \otimes \mathbf{V})}$$

Use divergence chain rule for dyadic tensors

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + \boxed{(\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V}} = -\boxed{\nabla P} - \rho \nabla \Phi$$

$(\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = \nabla \cdot (\rho \mathbf{V} \otimes \mathbf{V})$

$\nabla P = \nabla \cdot (P \mathbf{I})$

Rewrite pressure gradient as a divergence

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + (\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = -\nabla P - \rho \nabla \Phi$$

$$(\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = \nabla \cdot (\rho \mathbf{V} \otimes \mathbf{V})$$

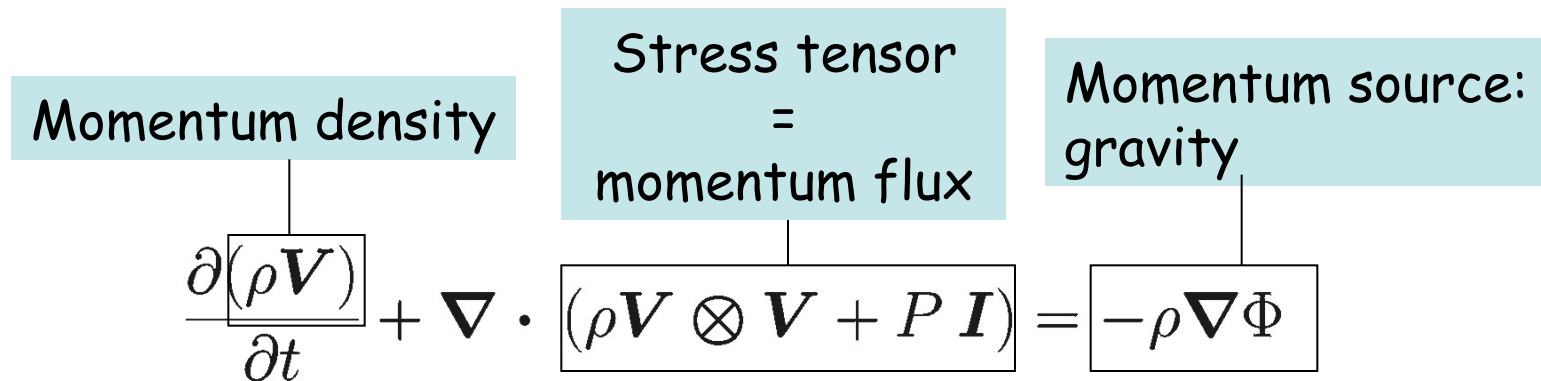
$$\nabla P = \nabla \cdot (P \mathbf{I})$$

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + \nabla \cdot (\rho \mathbf{V} \otimes \mathbf{V} + P \mathbf{I}) = -\rho \nabla \Phi$$

As advertised: Algebraic Manipulation!

$$\rho \left[\frac{\partial \mathbf{V}}{\partial t} + (\mathbf{V} \cdot \nabla) \mathbf{V} \right] = -\nabla P - \rho \nabla \Phi$$

$$\frac{\partial(\rho \mathbf{V})}{\partial t} + (\nabla \cdot (\rho \mathbf{V})) \mathbf{V} + \rho (\mathbf{V} \cdot \nabla) \mathbf{V} = -\nabla P - \rho \nabla \Phi$$



The tensor $\mathbf{R}_{ik} = \rho \mathbf{V}_i \mathbf{V}_k + p \delta_{ik}$ is the Reynolds stress tensor for an ideal fluid and represents the momentum flux