

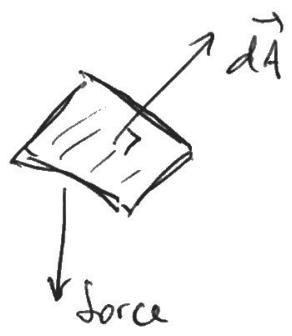
# Dissipative effects in fluid flow

When flow is not homogeneous (the gradients  $\nabla v$ ,  $\nabla T$ ,  $\nabla p$  are not identically zeros  $\Rightarrow v, T, p$  are not const)

$\Rightarrow$  the system is not exactly in thermodynamic equilibrium, and there are dissipative (irreversible) processes. (Obs: exception - rotation as a "solid" body).

We consider "sufficiently slow" motion, such that dissipative processes are small and can be studied in the first-order approximation w.r.t.  $\nabla v, \nabla T, \nabla p$ .

## Viscous flow:



Force in a general case:

(the Cauchy stress theorem based  
on conservation of moments & linear & angular,

$$\left[ (\text{force})_i = \sigma_{ij} dA_j \right], \quad i = 1, 2, 3$$

Here we use coordinates  $r = (x_1, x_2, x_3)$  and assume summation w.r.t. repeated indices:  $\sigma_{ij} dA_j := \sum_{j=1}^3 \sigma_{ij} dA_j$ .

By the Cauchy theorem,  $\sigma_{ij}$  is a symmetric tensor called the stress tensor. We define

$$\sigma_{ij} = -p \delta_{ij} + \sigma'_{ij} \quad \text{where } p = p(T, \sigma) \text{ is the}$$

pressure given by the equilibrium thermodynamics, and  $\sigma'_{ij}$  is the remaining symmetric part, called viscous tensor.

In equilibrium,  $\sigma'_{ij} = 0$ . Also, fluid is isotropic. (59)

Hence,  $\sigma'_{ij}$  is proportional to  $\partial v_i / \partial x_j$  as a tensor (maintaining the form of relation under rotations).

Obs: There are no linear tensor terms proportional to  $\nabla T$  or  $\nabla p$ .

Velocity gradient:  $\frac{\partial v_i}{\partial x_j} = \underbrace{\frac{1}{2} \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right)}_{\text{symmetric}} + \underbrace{\frac{1}{2} \left( \frac{\partial v_i}{\partial x_j} - \frac{\partial v_j}{\partial x_i} \right)}_{\text{antisymmetric}} \rightarrow \text{vorticity (rotation)}$

$\sigma'_{ij} = 0$  for a rotating fluid (exact thermodyn. equilibrium)

$\Rightarrow \sigma'_{ij}$  depends only on the symmetric part, which yields two independent tensor terms:

$$\left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right) \quad \text{and} \quad \underbrace{\left( \frac{\partial v_k}{\partial x_k} \right)}_{\text{div } v} \delta_{ij} \quad (\text{see tensor analysis})$$

We choose  ~~$\sigma'_{ij} = \frac{1}{2} \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right) + \frac{2}{3} \zeta \left( \frac{\partial v_k}{\partial x_k} \right) \delta_{ij}$~~

$$\sigma'_{ij} = \eta \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right) + \left( \zeta - \frac{2}{3} \eta \right) \left( \frac{\partial v_k}{\partial x_k} \right) \delta_{ij}$$

where  $\eta$  and  $\zeta$  are viscous coefficients.

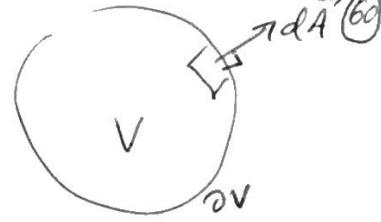
Obs:  $\eta$  and  $\zeta$  depend on thermodynamic variables,

e.g.  $\eta = \eta(T, g)$  or  $\eta(T, P)$ .

Obs:  $\zeta$  is called the second viscosity (defines dissipation due to volume changes)

2<sup>nd</sup> Newton's law:

$$\int_V \rho \frac{dv_i}{dt} dV = \oint_{\partial V} \sigma_{ij} dA_j, \quad i=1,2,3$$



Stokes formula:  $\oint_{\partial V} \sigma_{ij} dA_j = \int_V \frac{\partial \sigma_{ij}}{\partial x_j} dV$

$$\int_V \left( \rho \frac{dv_i}{dt} - \frac{\partial \sigma_{ij}}{\partial x_j} \right) dV = 0 \quad \text{for any volume } V.$$

$$\Rightarrow \underbrace{\rho \frac{d v_i}{d t}}_{\text{material derivative}} - \underbrace{\frac{\partial \sigma_{ij}}{\partial x_j}}_{\text{ }} = 0$$

$$\sigma_{ij} = -p \delta_{ij} + \sigma'_{ij} = \dots$$

$\Rightarrow$   
after  
long derivation  
(exercise)

$$\boxed{\rho \left( \frac{\partial v}{\partial t} + v \cdot \nabla v \right) = -\nabla p + \eta \Delta v + \left( \frac{2}{3} + \frac{\eta}{3} \right) \text{grad } dV v}$$

Navier - Stokes equations.

Obs Can add body forces, e.g.  
gravity:  $+ \rho \vec{g}$ .

Obs Here  $p = p(T, \rho)$  is defined by equilibrium thermodynamic relations.

Boundary conditions: no-slip condition on a rigid wall:  $v=0$  for  $r \in$  rigid boundary.



## Particular cases :

Eg (1) Neglecting dependence on temperature,

$$P = P(\rho) \text{ and taking } \eta = \text{const}, \gamma = \text{const},$$

we obtain the closed system of equations

for  $\rho(r, t)$  and  $v(r, t)$ : continuity eq. + N.S. eq.

(2) Neglecting variations of density,  $\rho \approx \text{const.}$

Continuity eq.  $\Rightarrow \boxed{\frac{d \nabla v}{dt} = 0}$  incompressibility condition.

$$\text{N.S. eq.} \Rightarrow \boxed{\frac{\partial v}{\partial t} + v \cdot \nabla v = - \frac{\nabla P}{\rho} + \gamma \Delta v}$$

incompressible Navier - Stokes

where  $\gamma := \frac{\eta}{\rho}$  is called the kinematic viscosity ( $\eta$  is the dynamic viscosity)

Neglecting dependence of  $\gamma$  on temperature,  
 ( N.S. eq + incomp. condition ) is the closed system  
 of equations. Here pressure  $P$  is uniquely  
 defined for the given  $v$  and boundary conditions  
 ( no need for thermodynamics ! ).

In general, we need one more equation that  
determines the evolution of thermodynamic (second)  
 variable, e.g., temperature  $T(r, t)$ .

Energy is conserved, so we can model dissipative processes by extra terms in the energy conservation law

$$\frac{\partial}{\partial t} \left( \frac{\rho v^2}{2} + g e \right) + \nabla \cdot \Gamma = 0,$$

Here  $\overset{\text{def}}{\Gamma} = (\Gamma_1, \Gamma_2, \Gamma_3)$  is the energy flux given by

$$\Gamma_i = \underbrace{\rho v_i \left( \frac{v^2}{2} + h \right)}_{\substack{\text{energy flux} \\ \text{in ideal flow}}} + \underbrace{q_i}_{\substack{\text{dissipative} \\ \text{contribution}}},$$

In the first-order approximation,  $q = (q_1, q_2, q_3)$  is proportional to gradients of velocity  $v$  and gradients of thermodynamical variables (we will choose  $T$  and  $P$ ).

(1) terms proportional to  $\partial v_i / \partial x_j$ :

Consider  $T \equiv \text{const} \Rightarrow$  there is no heat transfer  $\Rightarrow$  energy flux is due to mechanical work only.

This means that  $q$  is due to work of viscous forces.

change of energy = work

$\underset{\text{"}}{\underbrace{\text{energy flux} \cdot dt}}$

$\underset{\text{"}}{\underbrace{\text{force} \cdot \text{displacement}}}$

$$q_j dA_j \cdot dt = -\sigma'_{ij} dA_j \cdot v_i dt.$$

(63)

After dropping  $\frac{dt}{dt} \frac{d\mathbf{v}}{dt}$ , we obtain

$$q_i = -\sigma'_{ij} v_j \quad \text{or} \quad \boxed{\mathbf{q} = \sigma' \mathbf{v}} \quad (\text{matrix form})$$

(viscous stress tensor)

This is the energy flux due to viscosity.

(2) terms proportional to  $\nabla T$ . Due to isotropy of fluid  $\Rightarrow q_i = -\lambda \nabla T$  with some coefficient, called the thermal conductivity.

Obs  $\lambda$  is in general a function of local thermodynamic variables, e.g.,  $\lambda = \lambda(\rho, T)$ .

(3) there ~~are~~ cannot be terms proportional to  $\nabla P$ ,  $\theta$  as we will prove later below.

We obtained the energy equation in the form:

$$\frac{\partial}{\partial t} \left( \frac{\rho v^2}{2} + \rho e \right) = -\nabla \cdot \left[ \rho v \left( \frac{v^2}{2} + h \right) - \sigma' v - \lambda \nabla T \right],$$

where energy density  $e$ , enthalpy density  $h$  and thermal conductivity  $\lambda$  are functions of local thermodynamic variables  $(T, \rho)$  - ~~at~~ local equilibrium condition.

Expression for the viscous stress tensor  $\sigma'$  was given above (see derivation of N.S. eq.).

## Entropy equation

thermodynamics  
We will use :  $h = e + \frac{P}{\rho}$

$$\left. \begin{aligned} de = T ds + \frac{P}{\rho^2} df \Rightarrow \frac{\partial e}{\partial t} = T \frac{\partial s}{\partial t} + \frac{P}{\rho^2} \frac{\partial f}{\partial t} \end{aligned} \right\}$$

$$\left. \begin{aligned} dh = T ds + \frac{1}{\rho} dp \Rightarrow \nabla h = T \nabla s + \frac{1}{\rho} \nabla p \Rightarrow \nabla p = \rho \nabla h - P \nabla s. \end{aligned} \right\}$$

fluid dynamics  
 $\frac{\partial \rho}{\partial t} + \text{div}(\rho v) = 0$  continuity eq.

$$g \left( \frac{\partial v}{\partial t} + v \cdot \nabla v \right) = - \nabla p + \nabla \sigma' , \quad \sigma'_{ij} = \eta \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right) + \left( g - \frac{2\eta}{3} \right) \frac{\partial v_k}{\partial x_k} \delta_{ij}$$

$$\left. \begin{aligned} \frac{\partial}{\partial t} \left( \frac{\rho v^2}{2} + \rho e \right) &= - \text{div} \left[ \rho v \left( \frac{v^2}{2} + h \right) - \sigma' v - 2 \nabla T \right] \end{aligned} \right\}$$

$$\frac{\partial}{\partial t} \left( \frac{\rho v^2}{2} + \rho e \right) = \left( \frac{v^2}{2} + e \right) \frac{\partial \rho}{\partial t} + \rho v \cdot \frac{\partial v}{\partial t} + g T \frac{\partial s}{\partial t} + \frac{P}{\rho} \frac{\partial p}{\partial t} =$$

long derivation  
using all above eqs.  
(exercise)

$$\frac{\partial}{\partial t} \left( \frac{\rho v^2}{2} + \rho e \right) = \underbrace{g T \frac{\partial s}{\partial t}}_{\text{heat received}} = \underbrace{\sigma'_{ij} \frac{\partial v_i}{\partial x_j}}_{\text{viscosity}} + \underbrace{\text{div}(\lambda \nabla T)}_{\text{thermal conduction}}$$

$$\frac{\partial}{\partial t} (gs) = \cancel{g \frac{\partial \rho}{\partial t} + g \frac{\partial s}{\partial t}} = \text{div}(\dots) + \lambda \frac{(\nabla T)^2}{T^2} +$$

$$+ \frac{\eta}{2T} \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} - \frac{2}{3} \delta_{ij} \frac{\partial v_k}{\partial x_k} \right)^2 + \frac{\zeta}{T} (\text{div} v)^2$$

$$\frac{\partial}{\partial t} \int g s dV = \cancel{\int \text{div}(\dots) dV} + \lambda \int \frac{(\nabla T)^2}{T^2} dV + \frac{\eta}{2T} \int (\dots)^2 dV + \frac{\zeta}{T} \int (\dots)^2 dV$$

(vanishes at  $\infty$ )

total entropy  
(in whole space)

(65)

2<sup>d</sup> law of thermodynamics:  $\frac{dS}{dt} > 0$ .

$\Rightarrow \boxed{\lambda \geq 0, \gamma \geq 0, \beta \geq 0}$  all dissipative coefficients are non-negative.

Obs If one adds  $+\delta \nabla p$  to the energy flux, this yields the term proportional to  $+\gamma \int \nabla p \cdot \nabla T dV$  in the total entropy change. Since, it is not definite, from the 2<sup>d</sup> law of thermod.  $\Rightarrow \underline{\delta = 0}$ .

Full system of equations:

5 dependent variables: velocity field  $v(r,t)$  and thermodynamic fields, e.g.,  $\rho(r,t)$  and  $T(r,t)$ .

5 equations: continuity eq, N.S. eqs, energy conservation eq.

+ thermodynamic relations  $e(\rho, T)$  and  $h = e + \frac{P}{\rho}$ .

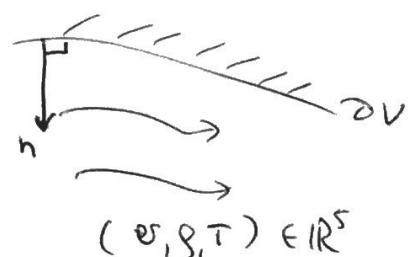
+ dissipative coefficients  $\lambda(\rho, T)$ ,  $\gamma(\rho, T)$  and  $\beta(\rho, T)$ .

Boundary conditions (example)

For a rigid isolating wall:

$r \in \partial V$ : (1)  $v = 0$  (no-slip)

~~slip~~  
~~slip~~



(2)  $n \cdot \nabla T = 0$  (no heat flux through)  
the Boundary

Obs. Recall that heat flux is:  $-\lambda \nabla T$ .

## Particular case (incompressible flow)

Main assumptions:

(1) Flow is slow with  $\omega \sim \frac{L}{t_*} \ll c$ , where

$L$  and  $t_*$  are characteristic length and time of the flow,  
 $c = \sqrt{\left(\frac{\partial P}{\partial \rho}\right)_S}$  (adiabatic derivative) is the sound speed.

$$\underline{\text{Obs}}: \left(\frac{\partial P}{\partial \rho}\right)_S = c^2 \Rightarrow (\text{"integrating" w.r.t. } g) \quad P \sim \rho c^2$$

From the N.S. equation:  $\underbrace{\rho \mathbf{v} \cdot \nabla \mathbf{v}}_{\text{convective term}} \sim \nabla P$

$$\Rightarrow (\text{by order of magnitude}) \quad \frac{\rho v^2}{L} \sim \frac{\delta P}{L} \Rightarrow \delta P \sim \rho v^2 \ll P \sim \rho c^2$$

(second-order small!)

Hence, hence, relative variations of pressure are small:

$$P = P_0 + \delta P, \quad P_0 = \text{const}, \quad \delta P \ll P_0.$$

For the temperature  $T = T_0 + \delta T$  with  $T_0 = \text{const}$  and  $\delta T$  small, we have

$$\rho(T, P) \approx \rho_0 + \left(\frac{\partial \rho}{\partial T}\right)_{T_0, P_0} \delta T + \left(\frac{\partial \rho}{\partial P}\right)_{T_0, P_0} \delta P$$

$\approx \rho(T_0, P_0)$

Neglecting the second-order term  $\delta P \sim \rho v^2$ , we have

$$\rho \approx \rho_0 + \left(\frac{\partial \rho}{\partial T}\right)_{T_0, P_0} \delta T = \rho_0 (1 - \alpha \delta T).$$

where  $\alpha = -\frac{1}{\rho_0} \left(\frac{\partial \rho}{\partial T}\right)_{T_0, P_0}$  is the coefficient of thermal expansion.

(2) Thermal expansion is weak if  $\delta T \ll 1/\alpha$ .

In this case  $\delta p = \cancel{\rho_0} - \rho_0 \propto \delta T \ll \rho_0$ , i.e.,

the density is approximately constant.

### Resulting equations of motion:

(A) Continuity eq.:  $\frac{\partial \rho}{\partial t} + \mathbf{v} \cdot \nabla \rho + \rho dN \mathbf{v} = 0$

$$\underbrace{\frac{\delta \rho}{\rho_0}}_{\sim \frac{\delta p}{\rho_0}} + \underbrace{\mathbf{v} \cdot \nabla \rho}_{\sim \frac{\mathbf{v} \delta \rho}{L}} + \underbrace{dN \mathbf{v}}_{\sim \frac{\rho_0 \mathbf{v}}{L}} = 0$$

$\Rightarrow$  in the first-order approximation:  $\boxed{dN \mathbf{v} = 0}$

Incompressibility cond.

### (B) Navier-Stokes eqs. (with gravity)

$$\rho \left( \frac{\partial \mathbf{v}}{\partial t} + (\mathbf{v} \cdot \nabla) \mathbf{v} \right) = -\nabla p + \gamma \Delta \mathbf{v} + \rho \vec{g}$$

The buoyancy term:  $\rho \vec{g} = \rho_0 \vec{g} + \delta \rho \vec{g} = \underbrace{\rho_0 \vec{g}}_{\text{this const. term can be included into } P.} - \rho_0 \alpha \delta T \vec{g}$

Hence, in the first-order approximation (dividing by  $\rho \approx \rho_0$ ):

$$\boxed{\frac{\partial \mathbf{v}}{\partial t} + \mathbf{v} \cdot \nabla \mathbf{v} = -\frac{\nabla p}{\rho_0} + \gamma \Delta \mathbf{v} - \alpha \Delta T \vec{g}} \quad \text{with } \Delta T = T - T_0$$

This is the Boussinesq approximation for buoyancy.

(c) Heat equation:

$$\rho T \frac{ds}{dt} = \sigma_i' \frac{\partial v_i}{\partial x_j} + \text{div}(\lambda \nabla T)$$

where  $\frac{d}{dt} = \frac{\partial}{\partial t} + \mathbf{v} \cdot \nabla$  and  $\sigma_i' = \eta \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right)$ . Recall:  $\frac{\partial v_k}{\partial x_k} = 0$

In the first-order approximation:

$$P = P_0 + \underbrace{\delta P}_{\sim v^2 \text{ (second-order)}} \rightarrow \text{neglect}$$

$$T = T_0 + \underbrace{\delta T}_{\text{first-order}} \rightarrow \text{keep } p.$$

$$\text{Term 1: } \rho T \frac{ds}{dt} \approx s_0 T_0 \left( \frac{\partial s}{\partial T} \right)_{P_0, P_0} \frac{dT}{dt} = s_0 c_p \frac{dT}{dt}$$

(Recall that  $c_p := T \left( \frac{\partial s}{\partial T} \right)_p$  is the heat capacity)

$$\text{Term 2: } \sigma_{ij}' \frac{\partial v_i}{\partial x_j} \xrightarrow[\text{symmetry}]{\leftrightarrow} \frac{1}{2} \sigma_{ij}' \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right) = \frac{\eta}{2} \left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right)^2$$

Term 3:  $\text{div}(\lambda \nabla T) \approx \lambda \Delta T$  with  $\lambda$  taken at  $T_0, P_0$ .

Resulting heat equation (dividing by  ~~$\rho s_0 c_p$~~ )

$$\boxed{\frac{\partial T}{\partial t} + \mathbf{v} \cdot \nabla T = \chi \Delta T + \frac{\gamma}{2c_p} \underbrace{\left( \frac{\partial v_i}{\partial x_j} + \frac{\partial v_j}{\partial x_i} \right)^2}_{\text{summed w.r.t. i and j}}}$$

where  $\gamma = \eta / s_0$  and

(summed w.r.t. i and j)

$\chi = \frac{\lambda}{\rho c_p}$  is called thermometric conductivity.

The full system:  $\boxed{\text{div } \mathbf{v} = 0} + \boxed{\text{Boussinesq eq}} + \boxed{\text{heat eq}}$

determine the fields  $\mathbf{v}(r,t)$ ,  $P(r,t)$  and  $T(r,t)$ , where all coefficients ( ~~$\rho, \alpha, \chi, c_p$~~ ) are constants. Obs:  $v \equiv 0 \Rightarrow \boxed{\frac{\partial T}{\partial t} = \chi \Delta T}$  stand. heat. eq.