

Lecture 2 -

Oil

Is popular for its ease of transport

Oil worsens the green-house effect but it's ^{has} high energy density, they are getting easier to extract.

They have high sulfur concentration.

Natural Gas → best fossil fuels.

Very efficient energy production, difficult to transport but needs infrastructure.

It produces the least CO₂ and is relatively clean.

Easier to use in small quantities.

To ship it needs to be transported by special ships.

Until before Ukraine, the use of gas a proportional to the regional production

Coal → Worst in every way

- ↳ most CO₂ producing
- ↳ high sulfur quantity
- ↳ not very efficient.

Things to worry about fossil fuels: direct and indirect pollution

As engineers to help stop global warming we need to reduce

the CO₂ we produce and prepare for the changed climate as it will take time to reequilibrate.

Solutions for mitigation:

↳ Nuclear

↳ Clean

↳ New ideas coming (SMR and fusion)

↳ Available for long periods

→ Flaws:

- High cost

- Requires safety

- Nuclear residue

↳ Hydroelectricity

↳ Ments

↳ Clean and renewable

↳ New small-scale machines

→ Flaws

↳ Limited by geography

↳ Some safety issues.

→ Newer Renewables

↳ Improvements in old renewables in different places.

↳ Wind and Solar are not programmable.

↳ CO₂ capture and sequestration

↳ Hydrogen

↳ Can be very dangerous, so it has to be limited to certain uses.

↳ Hydrogen is not readily available, it needs to be produced.

- ↳ We have different types of hydrogen depending on the energy source we use to generate it.
 - ↳ Possible fuel cells to produce electrical energy.
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Thermodynamics describes the energy transfer between a large number of elements. This is not possible, so there are two approaches to deal with that.

One is through observations and deriving phenomenological aspects. This is how the first and second principles were derived, these are statements not derived.

While we cannot study at a small scale but we can use probabilistic derivations, making assumptions we can derive macroscopic phenomena. This finds the statistical mechanics.

We use the phenomenological approach since we only care about macroscopic levels. But it is wrong since we are only looking at the macro scale, we need to place postulates, but it's top down.

We use an intermediate approach, using the postulate on statistical mechanics. From statistics we get temperature and we use the postulates on the temperature.

$\underline{L} = \Delta E_k + \Delta E_p \rightarrow$ This is not necessarily true (proved by Joule) as it does not consider heat which is another way

Work (only valid for the mechanical)

$$E_k = \frac{1}{2} m v^2$$

$$E_p = Mgz \quad E_p \text{ is irrelevant for us.}$$

system.

to transfer energy.

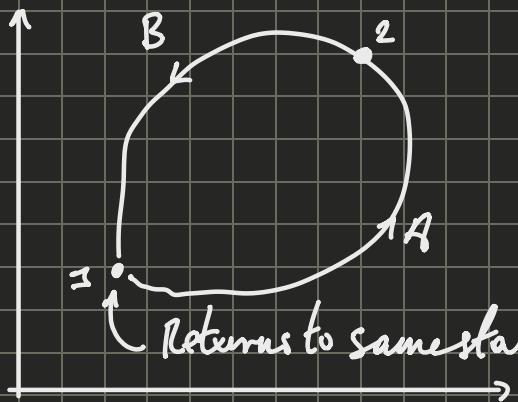
We also need a measure of the stored energy.

Microscopic changes en masse leave a macroscopic trace

Simple Thermodynamic System

State \rightarrow the set of conditions which describe the system

\hookrightarrow can be defined at the microscopic level with variables.



$$\oint (\delta L + \delta Q) = 0 = \int_{1A}^2 (\delta L + \delta Q) - \int_{1B}^1 (\delta L + \delta Q)$$

$$= f(1, 2) = U_2 - U_1 = \Delta U$$

Returns to same state since it's cyclic.

What enters the body is positive, what leaves is negative.

State function \rightarrow function that only depends on the state

Difference in internal energy is 0, proving the validity of the first principle of thermodynamics.

$$\left\{ \begin{array}{l} \Delta U = L + Q \\ dU = \delta L + \delta Q \end{array} \right. \rightarrow \text{First Principle of Thermodynamics.}$$

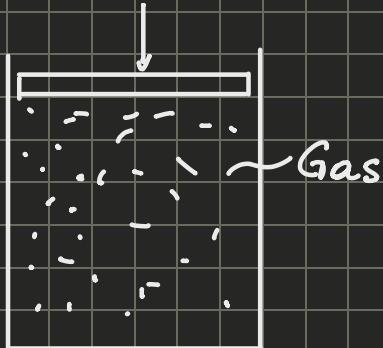
$\Rightarrow U$ is a state function

δ \Rightarrow not state function since they depend on the path.

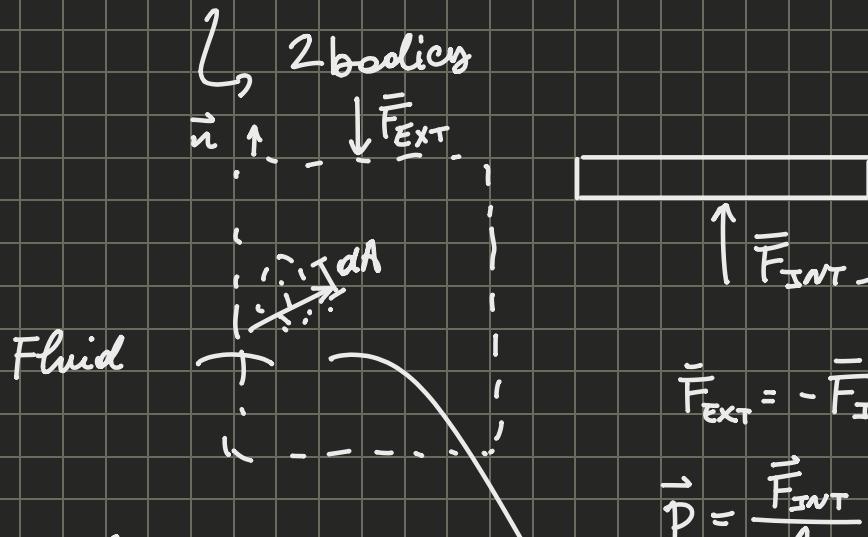
We have named E_k , work in our system they are needed, also E_{pot} for dams.
What are L and Q ?

Characterisation of Work / \rightarrow Heat Tomorrow

Work Modelling



The particles are in motion, they can heat the walls through momentum transfer



If we consider out fluid as a continuous medium it allows to maintain the macroscopic properties while maintaining the ability for our fluid to have a microscopic gradient, which at the same time allows us to not care about the microscopic randomness.

$$\bar{F}_{ext} = -\bar{F}_{int}$$

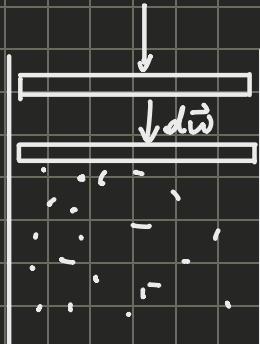
$$\vec{P} = \frac{\bar{F}_{int}}{A} = \lim_{dA \rightarrow 0} \frac{\delta \bar{F}_{int}}{dA} = P \vec{n}$$

↳ Pressure

always normal to area as the only relevant ones are normal

$$\Rightarrow \bar{F}_{int} = \int \delta \bar{F}_{int} = \int_A P \vec{n} dA \Rightarrow \bar{F}_{ext} = - \int_A P \vec{n} dA$$

What happens if we are able to move the piston



If $\text{d}w$ is fast it cannot be a simple system since it will generate kinetic energy.

So we want it to be slow so we can have infinitesimal states of equilibria.

If we do it quick we also need more force to generate the kinetics

$$\delta L = \vec{F}_{ext} \cdot d\vec{\omega} = - \int_A P \vec{n} dA \cdot d\vec{\omega} = - P \int_A \vec{n} d\vec{\omega} dA$$

↓

$$= -PdV \rightarrow \text{Revers. ble}$$

Work exchanged by a simple thermodynamic system that keeps the system the same and is reversible.

A process like this is not real since it takes a lot of time. If the velocity is high enough the force needs to be higher since we need to generate all the energy which we need for all "waste" processes that come along

$$\text{If } \vec{F}_{\text{ext}} \neq \vec{F}_{\text{int}} \rightarrow \vec{F}_{\text{ext}} = -\vec{F}_{\text{int}} + \vec{F}_w \rightarrow SL = -PdV + \vec{F}_w d\vec{w}$$

Wanted

$$= -PdV + SL_w$$

Energy is positive if it enters the fluid.

Always positive

\Rightarrow It is irreversible

So we need to put more energy to put work in less total work out

