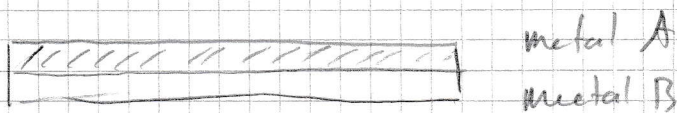
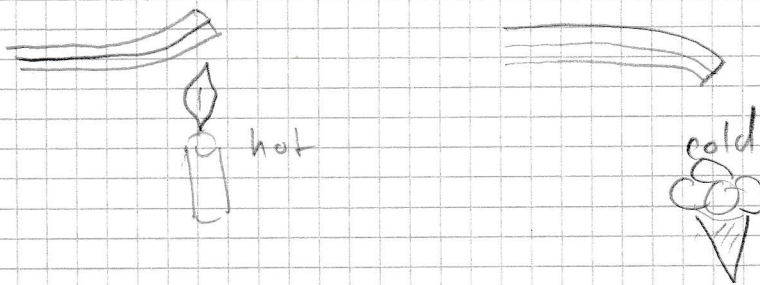
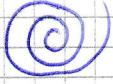


Bimetal thermometer

strip of 2 metals A, B with different coefficients of linear thermal expansion

If $T \nearrow$ (or \searrow), metals A & B will expand (contract) differently:



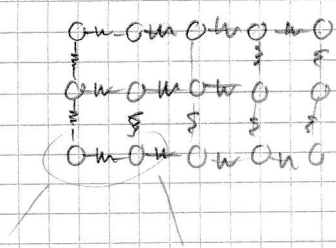
- Exp**
- 1) bi-metal strip
 - 2) snapping metal: thermostat
 - 3) spring thermometer:  - bimetal, or retro projector, works up

Exp IR camera to "see" (measure) temperature

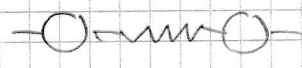
- ball (metal) on wood: impact piston glow
- rub shoe on floor
- metal plate,

Thermal expansion

microscopic picture (of matter) & expansion

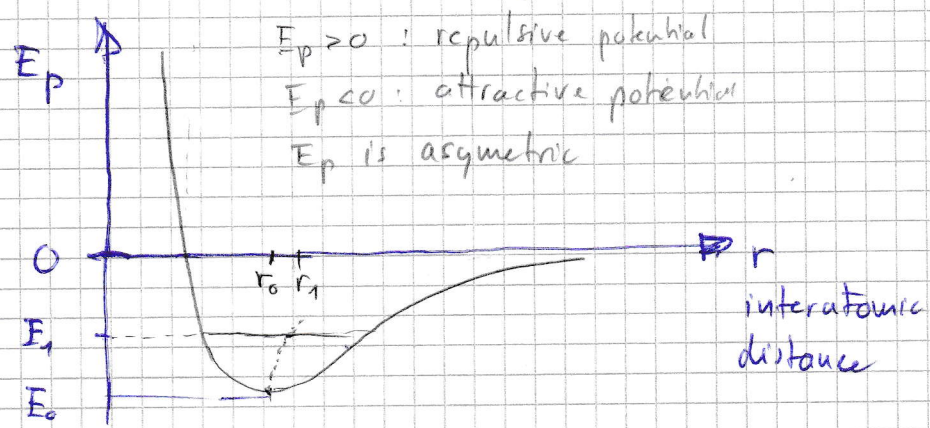


lattice of atoms with interatomic interaction (crystal structure)



binding energy between atoms: E_p potential energy
(remember coupled oscillators)

interatomic potential (potential energy)



$E_p > 0$: repulsive potential
 $E_p < 0$: attractive potential
 E_p is asymmetric

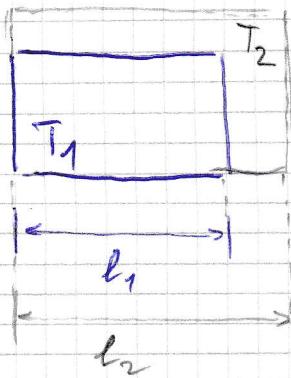
at $T=0$, $E_{kin} = E_k = 0$, no kinetic energy (atoms at rest)
and $E_0 = E_k + E_p = E_p$, r_0 : avg. distance between 2 atoms

at $T_1 > 0$, $E_1 = E_k + E_p > E_0$, as $E_k \neq 0$
 $r_1 > r_0$

As $T \nearrow$, $r \nearrow$ \forall atoms in a crystal / material

Macroscopically: if all atoms grow further apart from each other with increasing temperature the solid (macroscopic object composed of many atoms) will expand with temperature (increase its volume)

• macroscopic picture (of expansion)



$$l_2 - l_1 = \Delta l$$

$$T_2 - T_1 = \Delta T$$

$$\parallel \frac{\Delta l}{l} = \alpha_L \cdot \Delta T$$

↑ linear expansion coefficient $[\alpha_L] = \frac{1}{K}$

(slide) linear exp coefficients

$$\text{typ.} \sim 10^{-6} \text{ K}^{-1}$$

(Exp) Eisen Draht

or → do volume expansion first

Thermal expansion (1D), reminder

distance between atoms
increases (on avg)
=> expansion

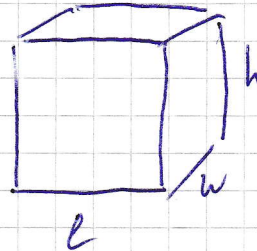
$$\left\| \frac{\Delta l}{l} = \alpha_l \cdot \Delta T \right.$$

↑
linear expansion coefficient

$$[\alpha_l] = \frac{1}{K}$$

(slide): Linear exp. coeff. values, (table)

for a volume:



$$V_1 = l \cdot w \cdot h$$

at ambient T

$$V_2 \text{ at } T_2 > T_1$$

$$\frac{\Delta V}{V} = \frac{V_2 - V_1}{V_1}$$

$$V_1 = l \cdot w \cdot h$$

$$V_2 = (l + \Delta l) \cdot (w + \Delta w) \cdot (h + \Delta h) = l \left(1 + \frac{\Delta l}{l}\right) \cdot (w + \Delta w) \cdot (h + \Delta h)$$

$$= \underbrace{l \cdot w \cdot h}_{V_1} \cdot \left(1 + \frac{\Delta l}{l}\right) \cdot \left(1 + \frac{\Delta w}{w}\right) \cdot \left(1 + \frac{\Delta h}{h}\right)$$

$$= V_1 \cdot \underbrace{\left(1 + \alpha_l \cdot \Delta T\right)^3}_{\alpha_l \cdot \Delta T}$$

$$\cong V_1 \cdot \left(1 + 3\alpha_l \cdot \Delta T\right)$$

$$\alpha_l \sim 10^{-6}$$

=> $\alpha_l \cdot \Delta T \ll 1$ at "reasonable" temperatures

and $\frac{\Delta V}{V} = \frac{V_2 - V_1}{V_1}$

$$\boxed{\frac{\Delta V}{V} \cong 3\alpha_l \cdot \Delta T = \alpha_v \cdot \Delta T}$$

$$[\alpha_v] = \frac{1}{K}$$

~~slide~~

exp: Iron wire expansion (slide)

slides: 1) α_l, α_v ; 2) volume expansion/road

o/e

exp \bar{F}_e min

$$75 \text{ V}, 28 \text{ A} = 21 \text{ kW}$$

$$l = 2.80 \text{ m}$$

• upon cooling, check the "kicks" (down and up again)
corresponds to phase transition

$$(1 + \epsilon)^n \approx 1 + n\epsilon$$

$\epsilon \ll 1$

Case of water:

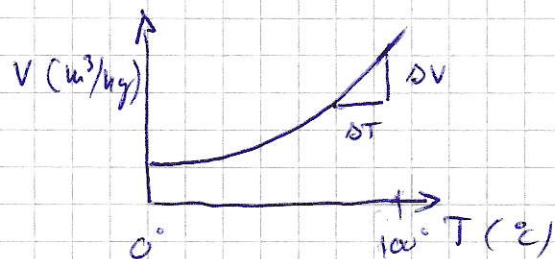
def: specific volume v : ratio of a substance volume to its mass

$$v = \frac{V}{m} = \frac{1}{\rho} \quad \rho: \text{density}$$

$$[v] = \text{m}^3/\text{kg}$$

$$[\rho] = \text{kg}/\text{m}^3$$

slide v vs T for water



Volume expansion α_v T:

$$\frac{\Delta v}{v} = \alpha_v \Delta T$$

$$\alpha_v = \left(\frac{\Delta v}{\Delta T} \right) \cdot \frac{1}{v}, \quad \text{slope of graph } v \text{ vs } T \propto \alpha_v$$

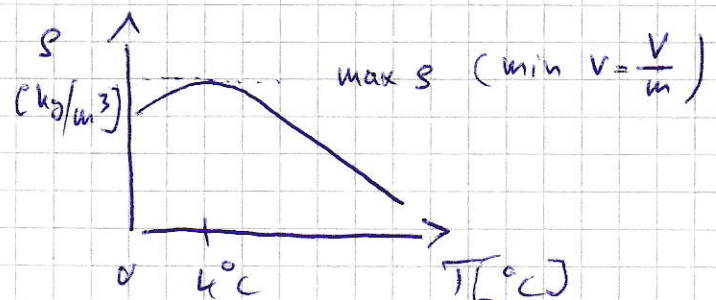
↑
prop. to

Note: $\alpha_v \sim$ const at high T (const. slope)

$\alpha_v \rightarrow 0$ as T decreases

$\alpha_v < 0$ below 4°C

show slide for ρ vs T

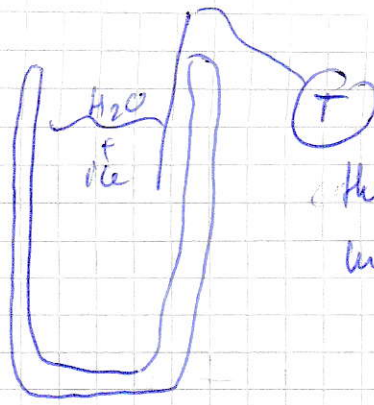


water has a point of max. density at 4°C

\Rightarrow water at 4°C will settle at the bottom of any water body (lake, pond, ...)

slide + Exp: 4°C water %

Exp: 4° water



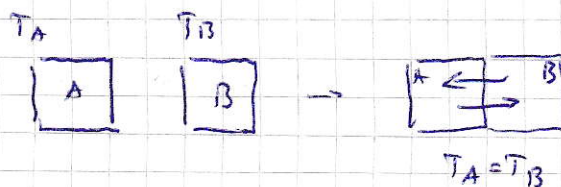
thermometer

meas. T vs depth

Heat transport and exchange

4

remember: thermal equilibrium -
reached upon
energy exchange



heat = energy that spontaneously passes from a hotter body
 Q , $[Q] = J$ to a colder body through thermal contact
 (via ~~heat~~ conduction, radiation or convection)

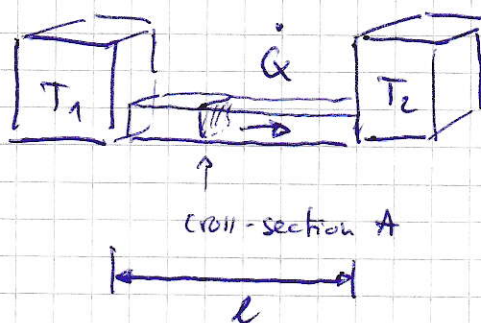
also mentioned: Temperature is a measure of the internal energy of a body,
and reflects the kinetic energy of the constituents of
this body (atoms, molecules)

It is possible to increase the temp. of an object (body)
by doing mechanical work on it as well (not only
by transferring heat to it via conduction, radiation or
convection)

Exp:

- ball thrown on wall (?)
- shoe sliding on floor

(observe with IR camera.)

heat conductionheat Q heat current: $\dot{Q} = \frac{dQ}{dt}$ 

$T_1 > T_2$

$$\dot{Q} = \lambda \cdot A \cdot \frac{\Delta T}{l}$$

λ : thermal conductivity of
material
 $[\lambda] = \frac{W}{m \cdot K}$, $W = \frac{J}{s}$ %

(slide) thermal conductivity table

NB: graphite, $\sim 500 \frac{W}{m \cdot K}$ on substrate

$\sim 2000 \frac{W}{m \cdot K}$ free standing

Expt: bar with nails in holes



on Cu side nails fall fast

stainless side, nails fall only later / or not at all
Steel

Heat capacity : ratio of heat added to (or removed from) a body to the resulting temperature change

5

$$K = \frac{\Delta Q}{\Delta T}$$

heat added (or removed) ↓
↑ heat capacity resulting change of temperature

$$[K] = \frac{J}{K}$$

specific heat capacity : (heat capacity per unit mass)
mass of the object

$$\Delta Q = c \cdot m \cdot \Delta T$$

↓ mass of the object
↑ specific heat capacity

$$[c] = \frac{J}{kg \cdot K}$$

also molar heat capacity:

$$\Delta Q = C \cdot N \cdot \Delta T$$

↑ nb of molar

remember
1 mole of a material
= N_A elementary entities
of this material (atoms,
molecules, ...)

$$N_A = 6.02 \cdot 10^{23}$$

Avogadro number

slide label for heat capacity
↑ specific

if c large, it means that one needs a large amount of heat (ΔQ) to increase the temperature (ΔT) of the ~~object~~ considered material
i.e. material "stores" a lot of energy per degree

Heat and work

(slide) Joule's experiment

increase the temp. of a system by doing work on it (or by adding heat: e.g. radiation)
 i.e., its internal energy (Joules)

⇒ equivalence mechanical energy and heat

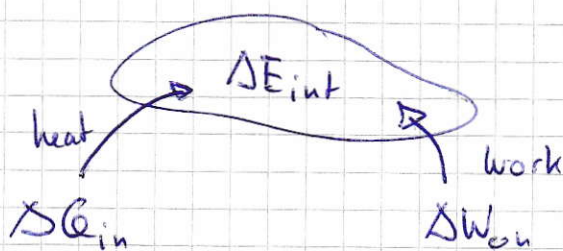
4.184 J of mechanical energy needed to increase the temp. of 1g of water by 1°C

$$(mgh = m'c \cdot \Delta T)$$

\uparrow \uparrow
 mass falling mass of water

0/0

⇒ | relationship between internal energy (or measured by the temperature), work done on a system and the heat transferred to a system? |



ΔE_{int} : change of internal energy of the system

ΔQ_{in} : heat transferred to the system

ΔW_{on} : work done on the system

|| 1st law of thermodynamics || $\Delta E_{int} = \Delta Q_{in} + \Delta W_{on}$ conservation of energy

• if the system is a machine and does (produces) a work,

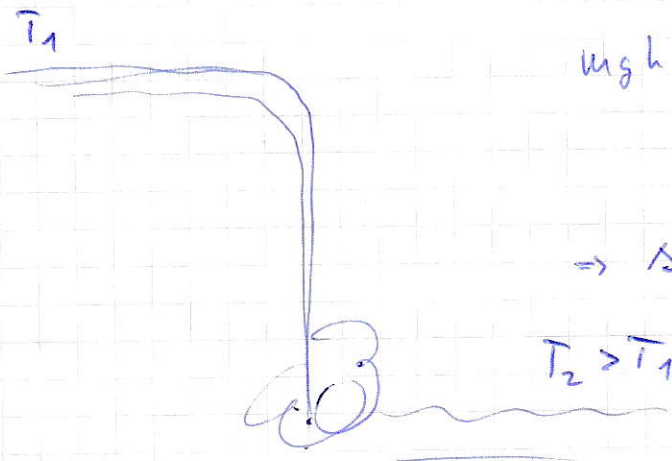
$$\Delta E_{int} = \Delta Q_{in} - \Delta W_{by}$$

ΔW_{by} : work done by machine

• note: E_{int} is a function of the state of the system and depends on pressure, volume, temperature.

Q, W are independent of the state of the system; they ^{are a} measure of transferred energy

example: waterfall $\rightarrow T_2 > T_1$, $\Delta T = ?$



$$mgh = m \cdot c \cdot \Delta T$$

\uparrow
specific heat capacity

$$\Rightarrow \Delta T = \frac{gh}{c}$$

$$T_2 > T_1$$

the same
if we consider a given
mass of water in before
and after fall

for a 100m (!) waterfall

$$\Delta T = \frac{9.81 \frac{\text{m}}{\text{s}^2} \cdot 100 \text{m}}{4.184 \frac{\text{J}}{\text{kg} \cdot \text{K}}} \approx 0.2 \text{ K}$$

$\checkmark = \frac{\text{N}}{\text{kg}}$

(4.184 $\frac{\text{J}}{\text{g} \cdot \text{K}}$)