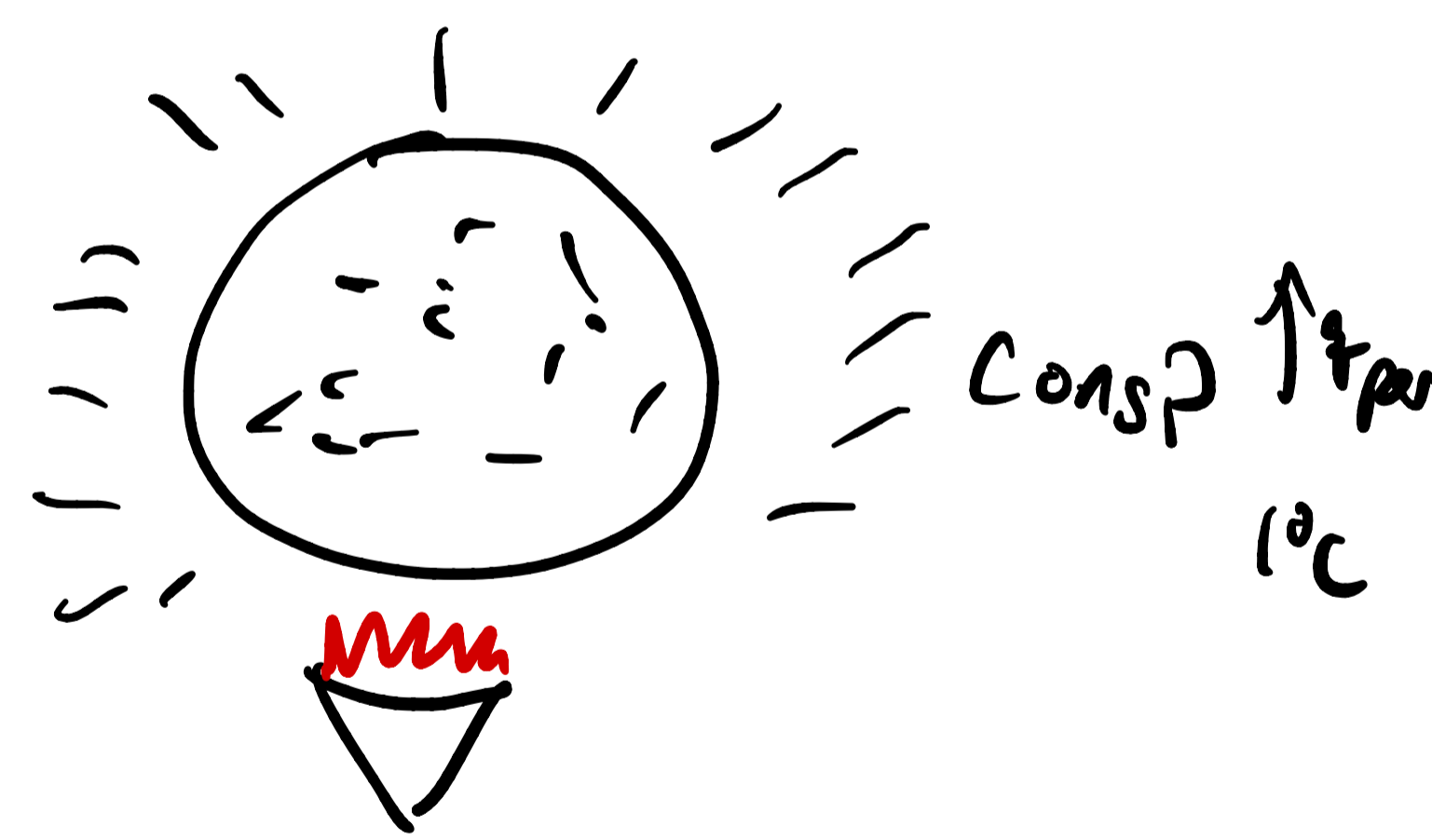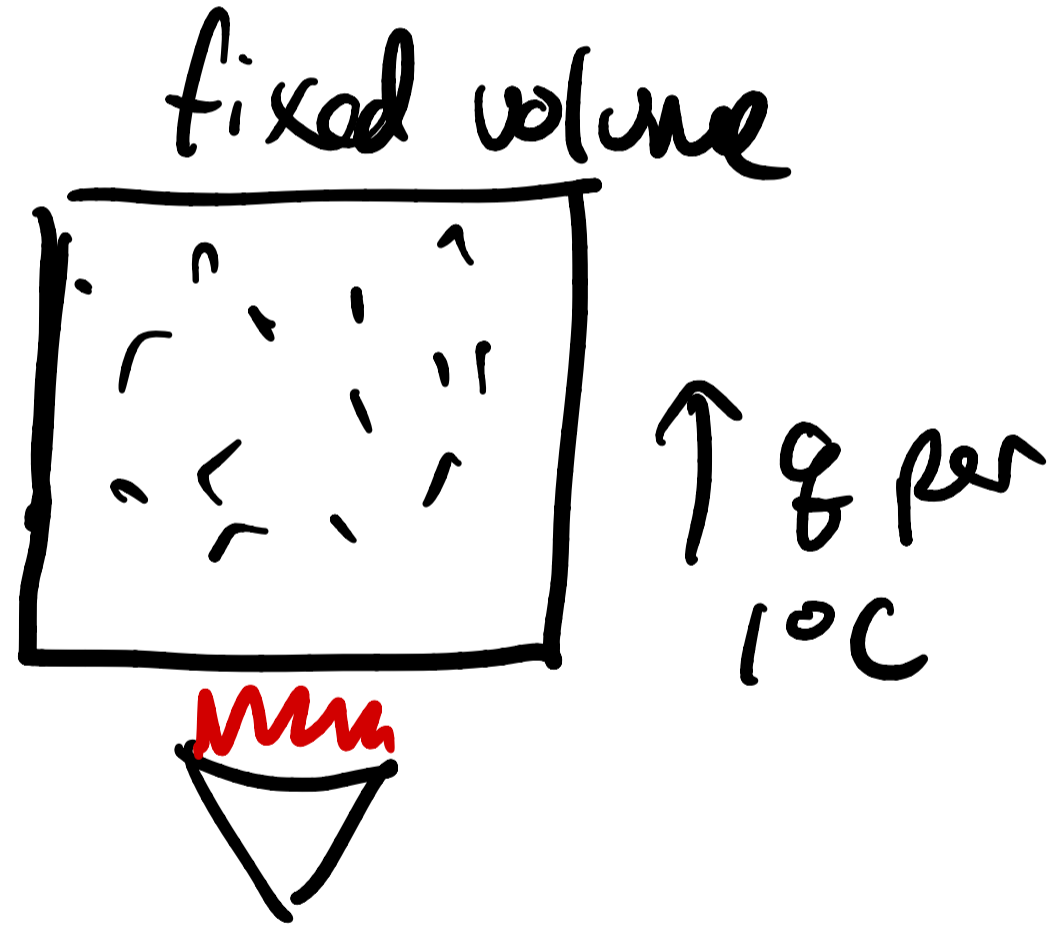# Lecture 6

heat capacity (extensive quantity)

$$dq = CdT$$

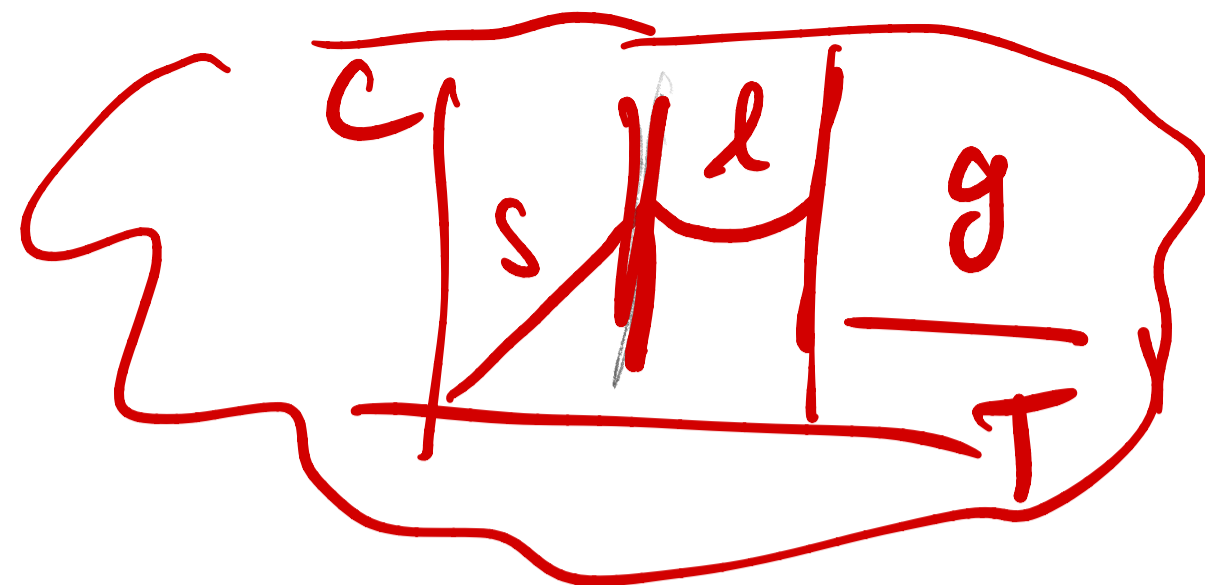$$C_v = \left(\frac{\partial q}{\partial T}\right)_V \qquad C_p = \left(\frac{\partial q}{\partial T}\right)_P$$

fixed volume

$\uparrow q$ per $1°C$

consp $\uparrow q$ per $1°C$

$$C_p > C_v$$

How much heat flows:

$$dq = C(T) \, dT$$

$$q = \int_{T_1}^{T_2} C(T) \, dT \approx C \int_{T_1}^{T_2} dT = C(T_2 - T_1)$$
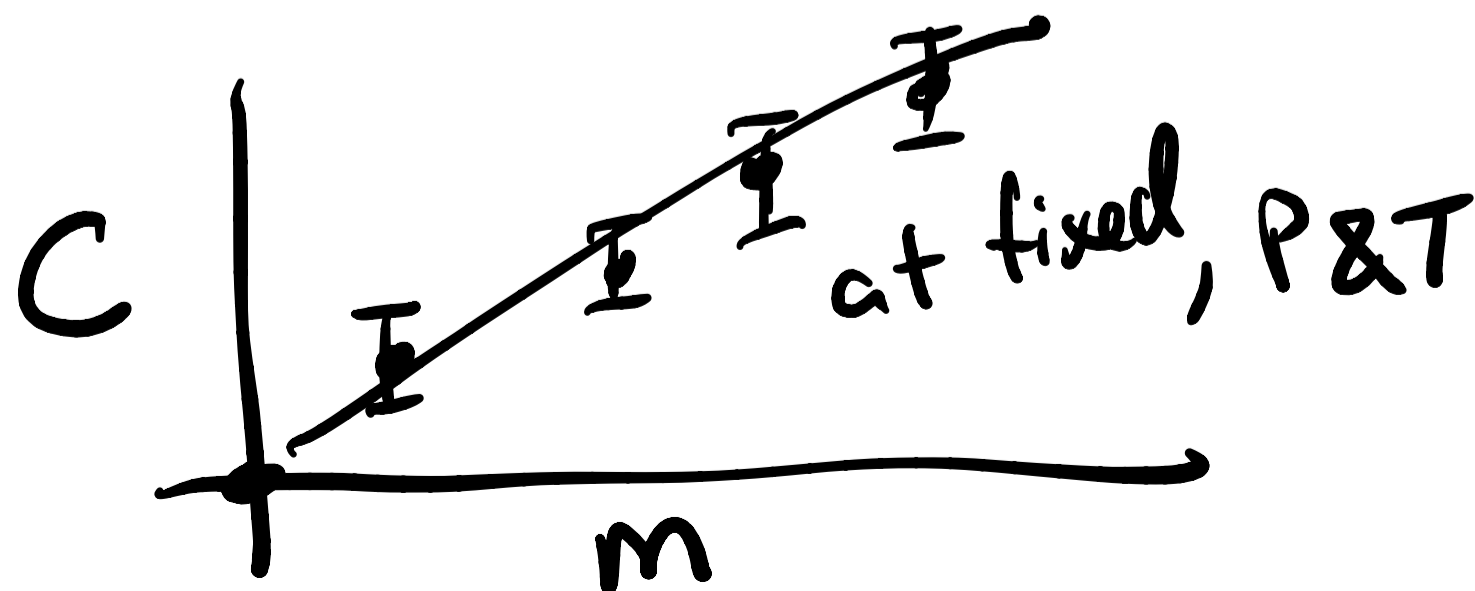
if T_1 & T_2 aren't that diff
& phase transition, $C(T) \approx C$

"$q = n C \Delta T$"
$\uparrow$ molar

"$q = m c \Delta T$"
$\uparrow$ per mass

C vs m, at fixed, P&T

How much does $\mathcal{E}$ change

$$d\mathcal{E} = dq + d\omega \qquad \text{(1st law)}$$

$$= CdT - PdV$$

$$\Delta\mathcal{E} = \int_a^b (CdT - PdV)$$

a) Const volume, $dV = 0$ $\qquad C = C_V$

$$\Delta\mathcal{E} = C_V \Delta T$$

b) const P, $C = C_p$

$$\Delta\mathcal{E} = C_p \Delta T - P\Delta V$$

$$C_v = \left(\frac{\partial \mathcal{E}}{\partial T}\right)_V = C_v \frac{\partial T}{\partial T} = C_v$$

$$\underline{\qquad\qquad\qquad}$$

$$d\mathcal{E} = C_v dT$$

$$dq = d\mathcal{E}$$

(a) C_v is how much energy changes for ~~for~~ some change in temp
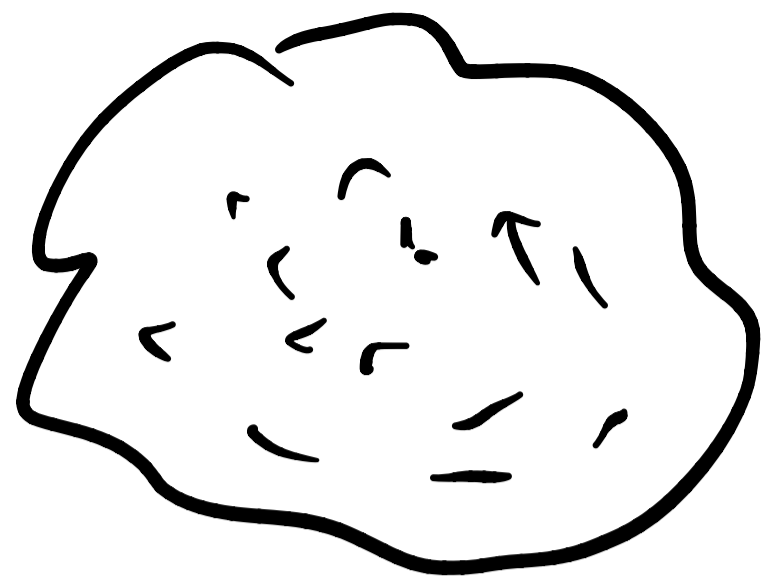
b) $d\mathcal{E} = dq - PdV \implies dq = d\mathcal{E} + PdV$

$$C_p = \left(\frac{d(\mathcal{E} + PV)}{dT}\right)_P \qquad d\mathcal{E} + PdV$$

Enthalpy

$$H \equiv E + PV$$

$$d(H) = d(E) + d(PV)$$

$$= dE + VdP + PdV$$

Consider $dP = 0$

@ const P

$$dH = dE + PdV = dq$$

$$c_p = \left(\frac{\partial H}{\partial T}\right)_P$$

Ideal gas: (monatomic ideal gas)

2 assumptions: - take up no space
- don't interact

① $PV = nRT$

② $\mathcal{E} = \frac{3}{2}nRT$

$\mathcal{E}_{total} = \overset{\propto T}{KE} + \cancel{PE} \; 0$

$$C_V^{ig} = \left(\frac{\partial \mathcal{E}}{\partial T}\right)_V = \frac{3}{2}nR \quad , \quad c_V = \frac{C_V}{n} = \frac{3}{2}R$$
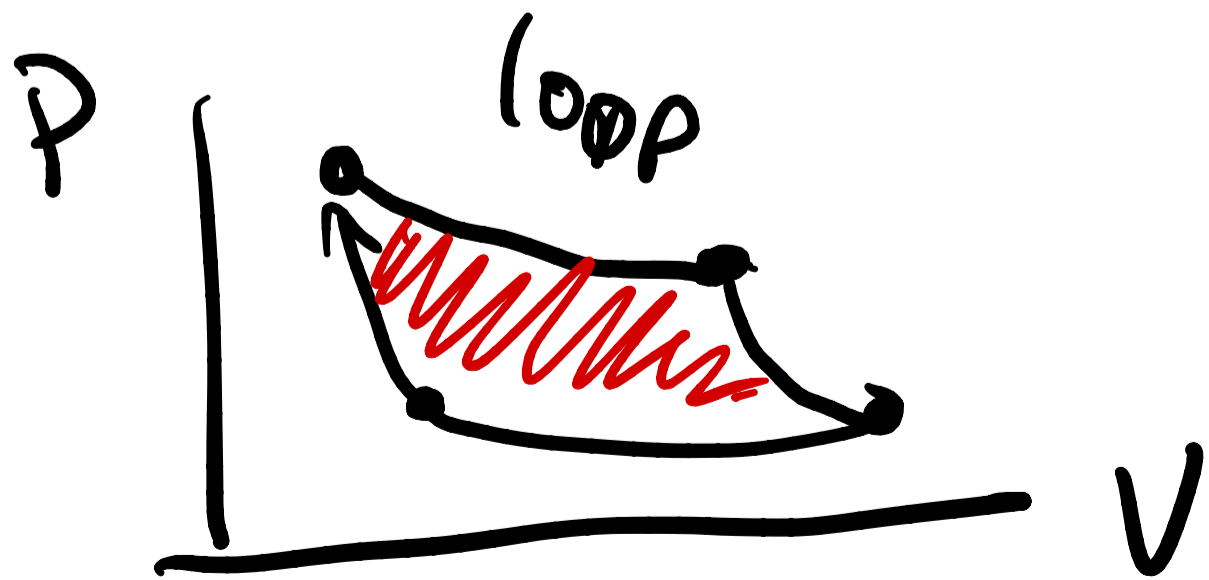
$$C_P^{ig} = \left(\frac{\partial(\mathcal{E}+PV)}{\partial T}\right)_P = \frac{\partial}{\partial T}\left(\frac{3}{2}nRT + nRT\right) = \frac{5}{2}nR$$

$C_p > C_V$, ideal gas

$$C_p - C_V = nR$$

$\rightarrow$ true for any # atoms

Preview: how to do work
through a cyclic process

$$\omega = \int_{cycle} -P \, dV$$
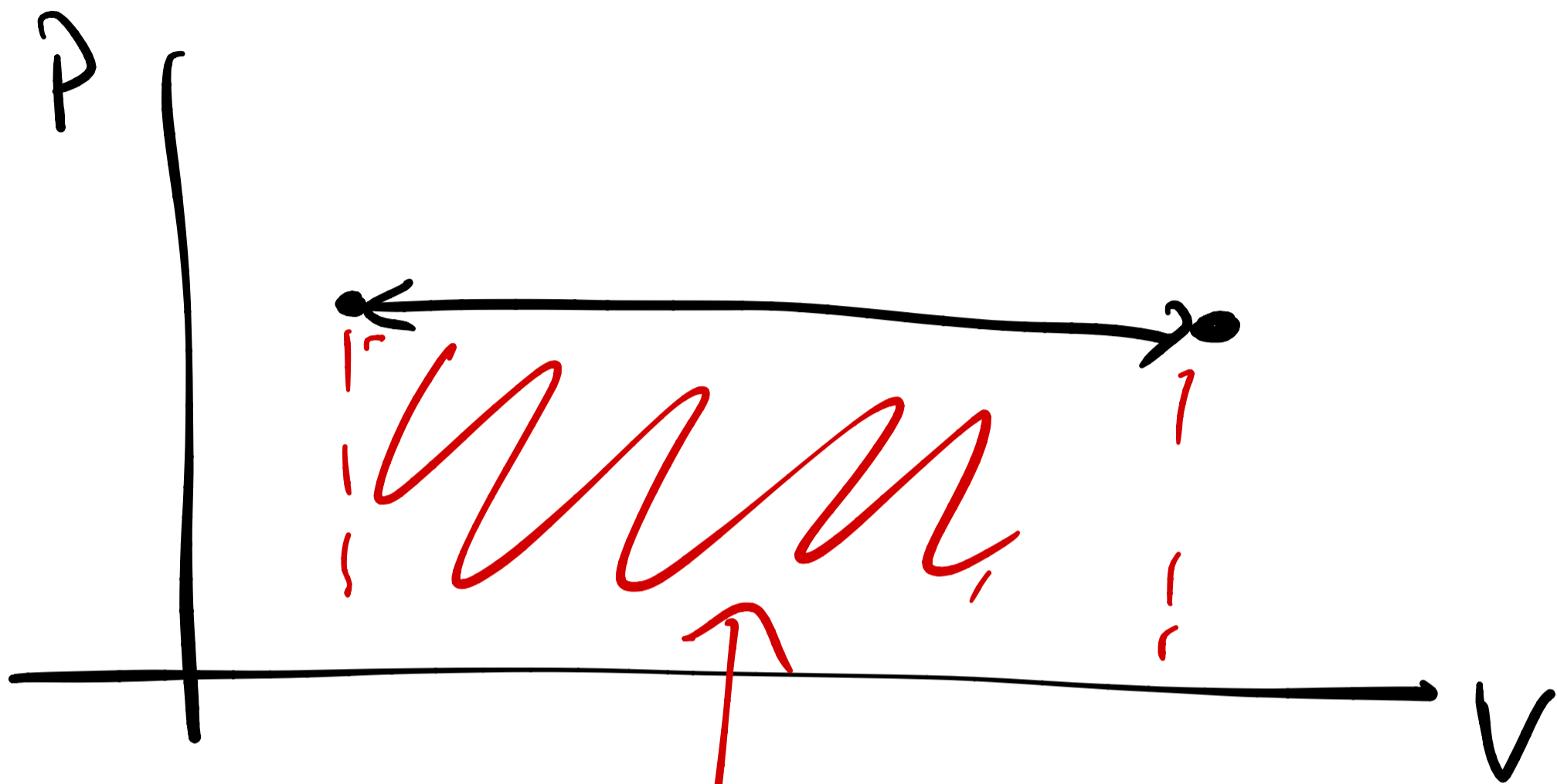
Change state:

- Const P, $dP = 0$

- Cont V, $dV = 0$

- Const T, $dT = 0$

- Adiabatic, $dq = 0$

need w, q, ΔE

① dP $\qquad$ $(PV = nRT)$

true in general

$$\underline{w} = -P\Delta V$$

$$q = C_P \Delta T \longleftarrow \text{ for an ideal gas,} \quad \left\{ \Delta T = \frac{P_2 V_2}{nR} - \frac{P_1 V_1}{nR} \right.$$

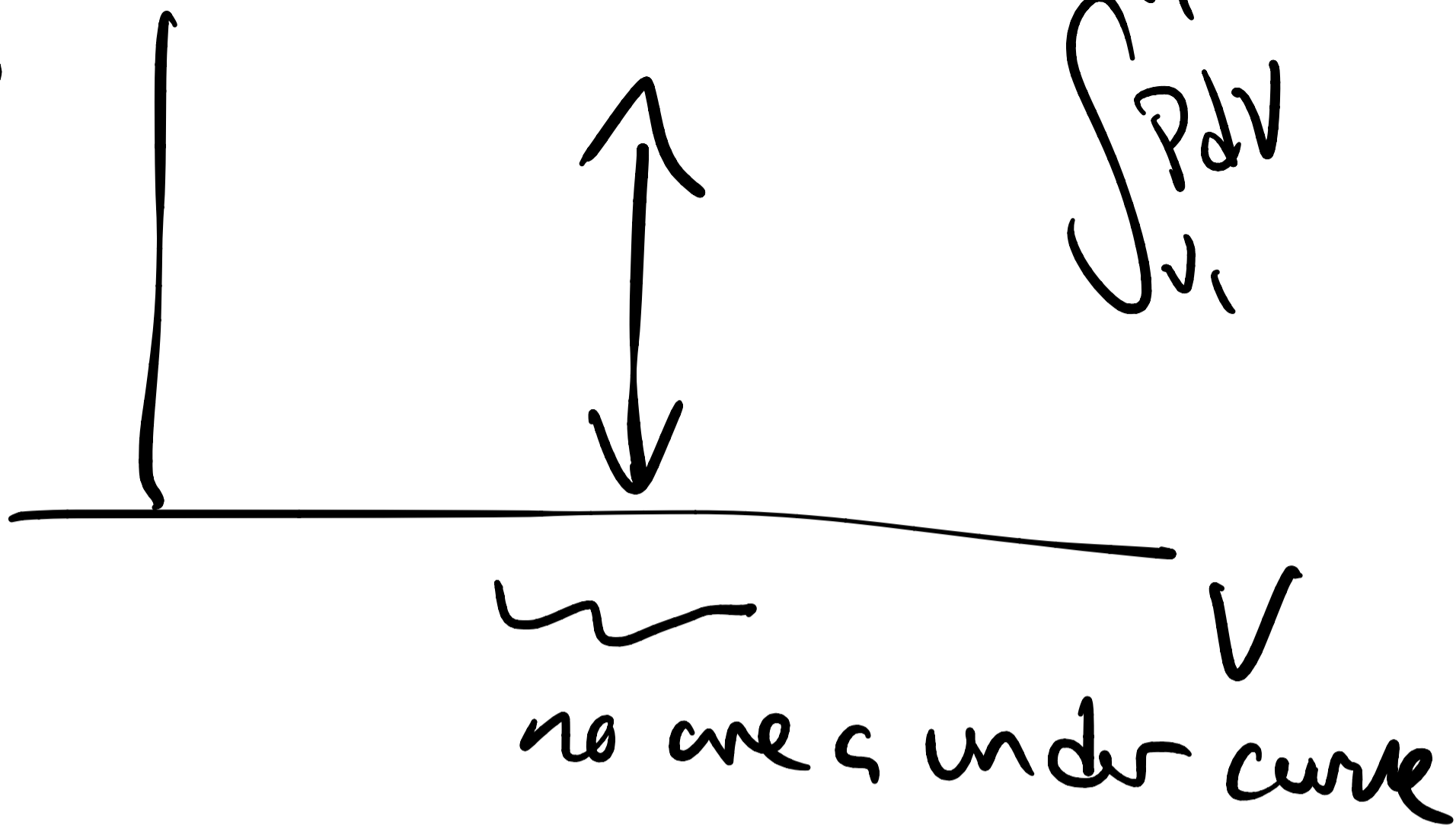$$\Delta E = q + w = C_P \Delta T - P\Delta V \left. \right\} = \frac{P}{nR} \Delta V$$

② $dV = 0$?

($dw = 0$)

$w = 0$

$q = C_V \Delta T$

$\Delta E = C_V \Delta T$

$$\int_{V_i}^{V_i} PdV$$
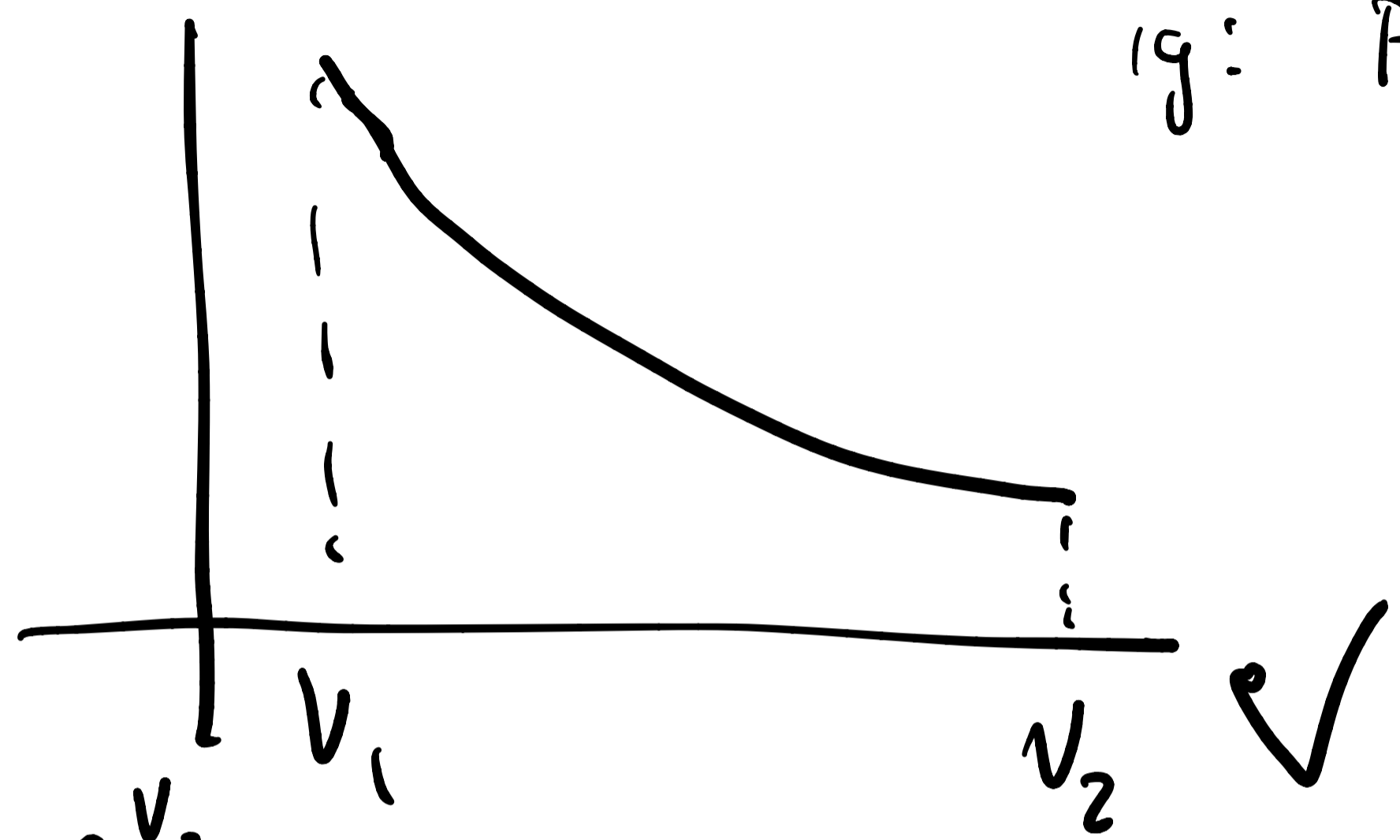
no area under curve

V

(for an ideal gas,
follows from energy
formula)

③ $dT = 0$

What does this look like

ig: $P = nRT \cdot \dfrac{1}{V}$

Isothermal expansion or compression

$$w = -\int_{V_1}^{V_2} P\,dV = -\int_{V_1}^{V_2} \frac{nRT}{V}\,dV$$

$$= -nRT\int_{V_1}^{V_2} \frac{1}{V}\,dV$$

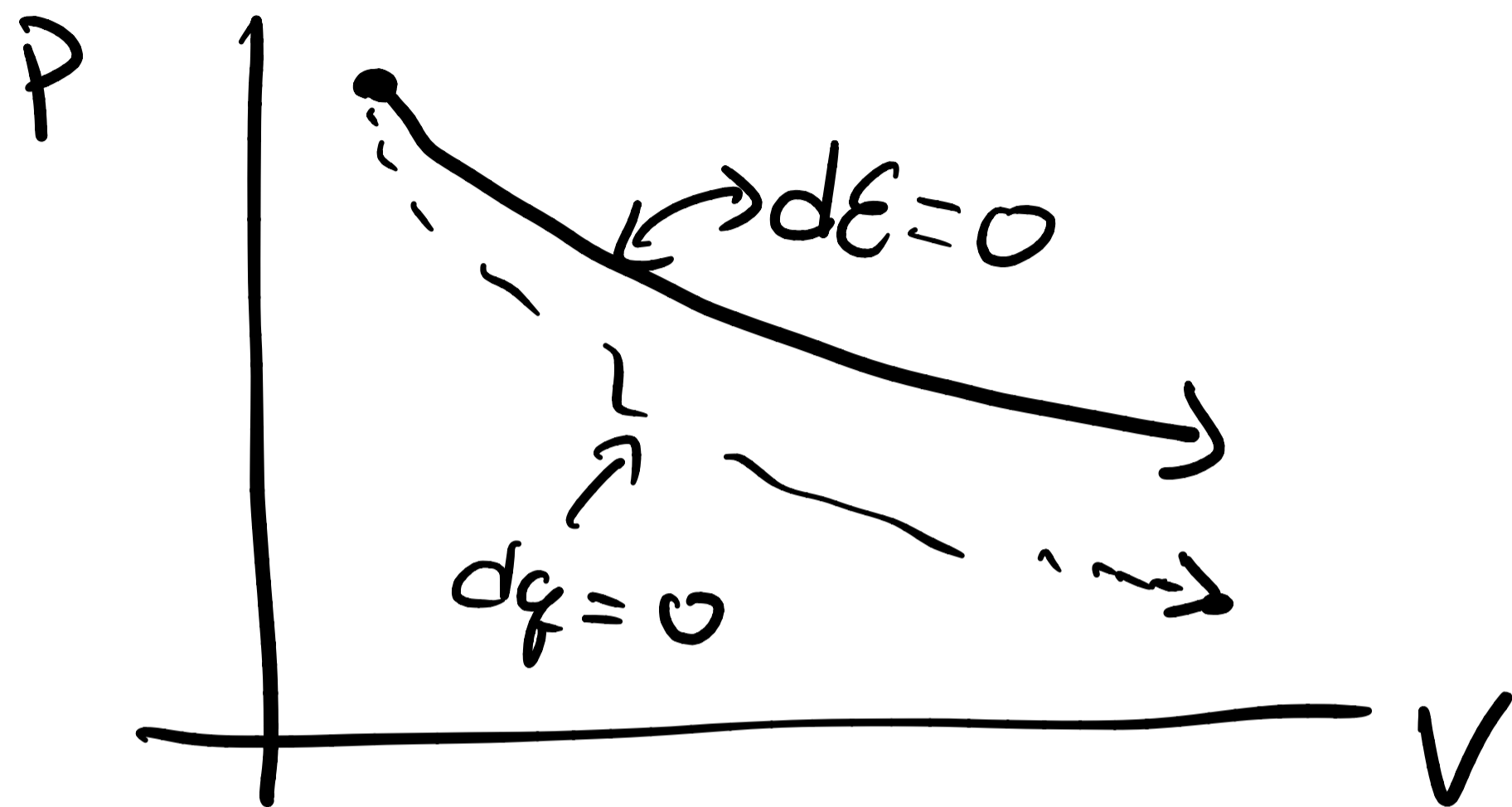$$= -nRT \ln(V_2/V_1)$$

question

$q > 0,$

$q < 0,$

$q = 0$

i.g. $\quad d\mathcal{E} = d\left(\frac{3}{2}nRT\right) = \frac{3}{2}nRdT = 0$

$0 = dq + dw = q + w \implies q = -w$

$q = + nRT\ln\left(V_2/V_1\right)$

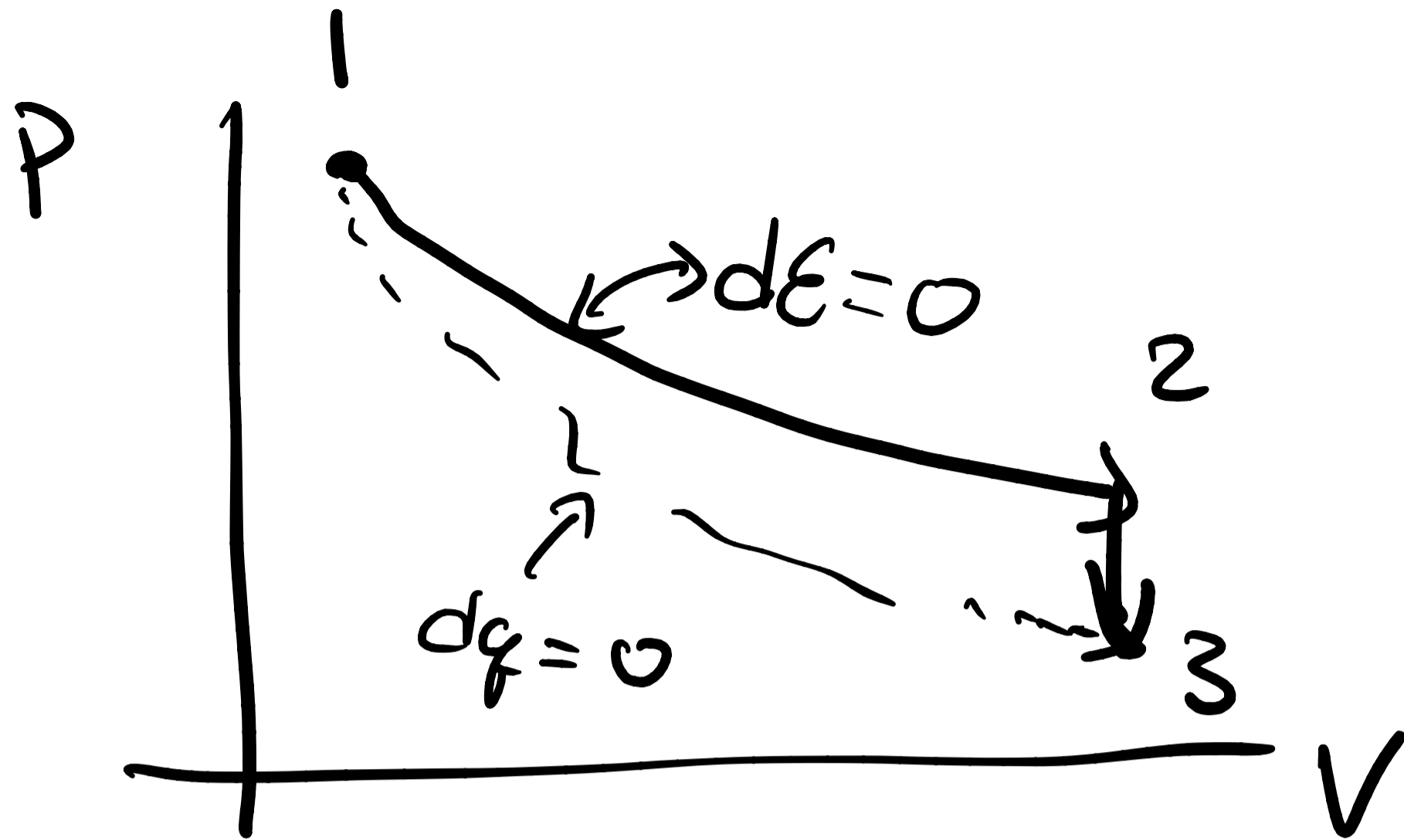$q > 0$ for expansion

④ Adiabatic expansion, $dq = 0$

pressure drops faster due to cooling

$$dE = \cancel{dq} + d\omega = -P(v)\,dV$$

$$= -\frac{nR}{V}T(v)\,dV$$

how much does it cool? ← next time

P (y-axis), V (x-axis)

1 (point)

$\rightarrow dE = 0$

$dq = 0$

2

3

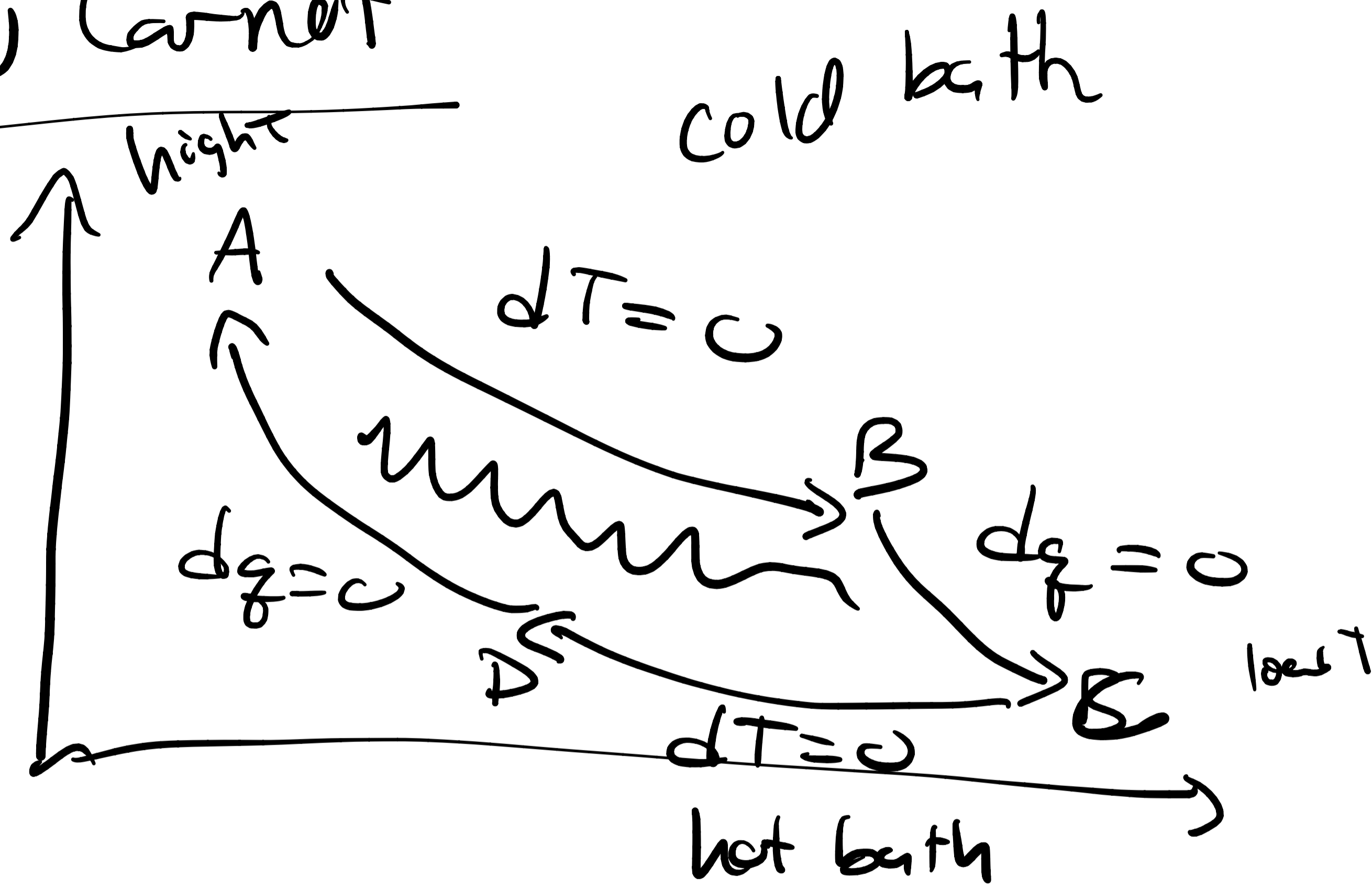$1 \rightarrow 2 \quad dT = 0$

$2 \rightarrow 3 \quad dV = 0$

Change in E from Pt $1 \rightarrow$ Pt 3

$$= Pt\, 1 \rightarrow 2 + Pt_2 \rightarrow Pt_3$$

next time . . .

Preview Carnot

high

cold bath

A

$dT = 0$

B

$dq = 0$

$dq = 0$

D

$dT = 0$

E   lowest T

hot bath

4 steps

temps T_h & T_c

efficiency $= w/q_{in} = 1 - T_c/T_h$