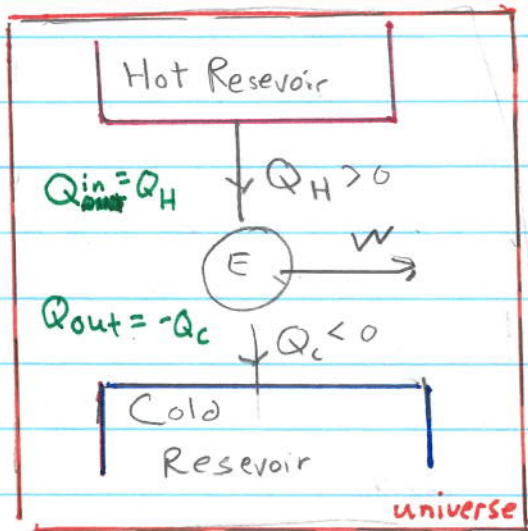


# Carnot Efficiency Revisited

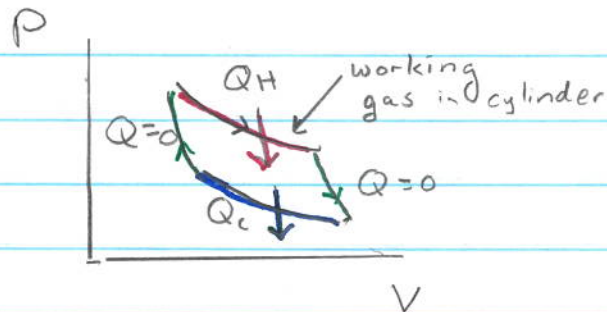


$$\textcircled{1} \Delta S_{\text{Hot}}^{\text{reservoir}} = -\frac{Q_H}{T_H} \equiv -\frac{Q_{\text{in}}}{T_H}$$

$$\textcircled{2} \Delta S_{\text{Cold}}^{\text{res}} = -\frac{Q_c}{T_c} \equiv \frac{Q_{\text{out}}}{T_c} \quad \begin{matrix} Q_c < 0 \\ Q_{\text{out}} > 0 \end{matrix}$$

$$\textcircled{3} \Delta S_{\text{working}} = 0$$

Substance of engine



This is a closed loop  
The gas starts and stops at same place. So  $\Delta S = 0$ .

$$\Delta S_{\text{universe}} = -\frac{Q_H}{T_H} + -\frac{Q_c}{T_c} \geq 0$$

rearranging:  $-\frac{Q_c}{T_c} \geq \frac{Q_H}{T_H}$  i.e.  $\boxed{\frac{|Q_c|}{T_c} \geq \frac{Q_H}{T_H}}$

Then this implies that the efficiency is less than carnot efficiency:

$$\frac{|Q_c|}{Q_H} \geq \frac{T_c}{T_H} \quad \text{or} \quad \eta \equiv 1 - \frac{|Q_c|}{Q_H} \leq 1 - \frac{T_c}{T_H}$$

↑ efficiency

The condition:

$$\frac{|Q_c|}{T_c} \geq \frac{Q_H}{T_H} \quad \text{is saying equivalently}$$

$$\boxed{S_{\text{out}} \equiv \frac{Q_{\text{out}}}{T_c} \geq S_{\text{in}} \equiv \frac{Q_{\text{in}}}{T_H}}$$

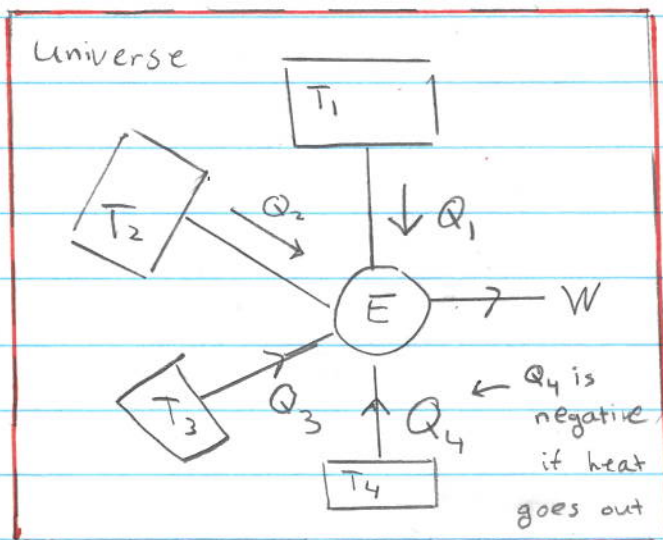
or entropy is produced!

by the engine cycle

## Generalizing

(working substance)

- We can have an engine with many reservoirs at different temperatures. At each reservoir the working substance exchanges heat.



This is reversible if each reservoir is in perfect equilibrium with the working substance

- The generalization is of Carnot to multicarnot is simple

$$\Delta S_{\text{universe}} = - \sum_i \frac{Q_i}{T_i} \geq 0 \text{ or ultimately } - \oint \frac{dQ}{T} \geq 0$$

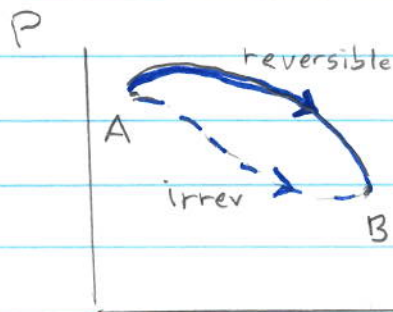
Or dividing into in ( $dQ > 0$ ) & out ( $dQ < 0$ )

$$- \int_{\text{in}} \frac{dQ}{T} + \int_{\text{out}} \frac{|dQ|}{T} \geq 0$$

and thus, **engine produces entropy**

$$S_{\text{out}} = \int_{\text{out}} \frac{|dQ|}{T} > \int_{\text{in}} \frac{dQ}{T} \equiv S_{\text{in}}$$

- For any process which is irreversible taking the system from  $A$  to  $B$  then we have



$$- \int_A^B \frac{dQ_{\text{irrev}}}{T} + \int_A^B \frac{dQ_{\text{rev}}}{T} \geq 0$$

non equilibrium transition from  $A$  to  $B$  of working substance

equilibrium transition  $B$  to  $A$

$$\int_B^A \frac{dQ_{\text{rev}}}{T} = - \int_A^B \frac{dQ_{\text{rev}}}{T}$$

First we go from  $A$  to  $B$  via a non-equilibrium process (irrev), and then back via an equilibrium process  $B$  to  $A$  (reversible)

Leading to

$$\int_A^B \frac{dQ_{\text{rev}}}{T} \geq \int_A^B \frac{dQ_{\text{irrev}}}{T}$$

in equilibrium we have  $dS = dQ_{\text{rev}}/T$

or

$$\Delta S_{A \rightarrow B} \geq \int_A^B \frac{dQ_{\text{irrev}}}{T}$$

Thus we see that the change in entropy  $dS$ , is in general greater than  $dQ/T$ .

- If equilibrium between the reservoirs and working substance is maintained we have

$$\Delta S_{A \rightarrow B} = S(B) - S(A) = \int_A^B \frac{dQ}{T}$$