

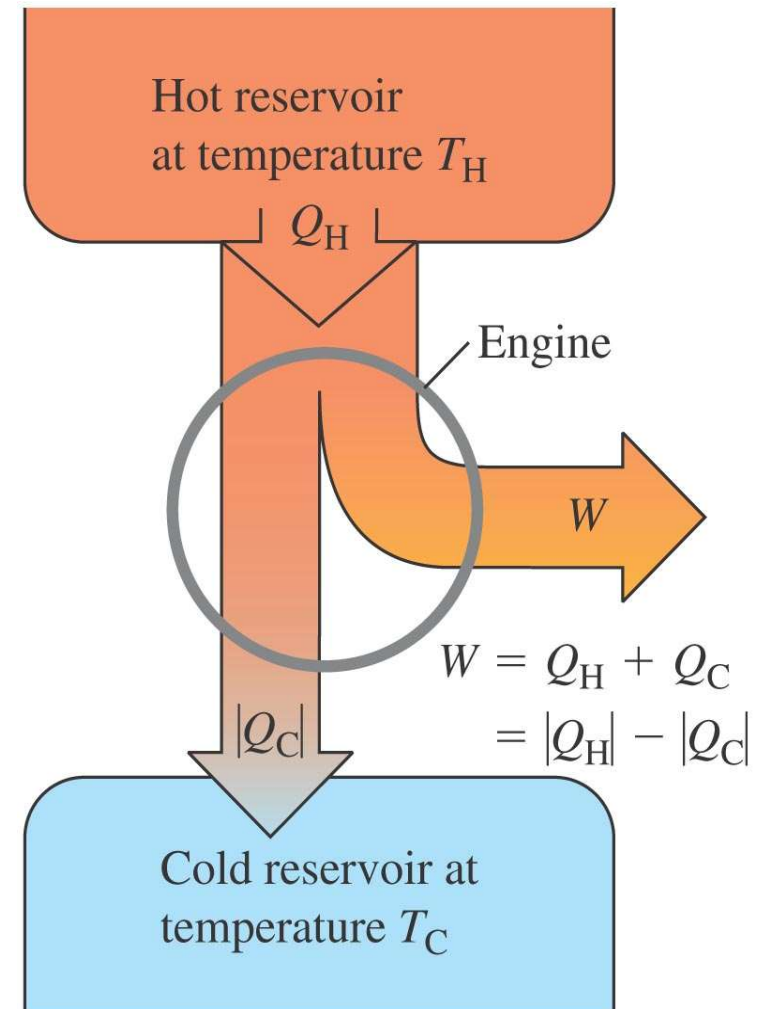
Heat Engines

- Definition: A device that converts a given amount of *heat* into *mechanical energy*.
- All heat engines carry some *working substance* thru a *cyclic process*:

Engine absorbs heat from hot reservoir at T_H

Mechanical work is done by engine

Engine releases residual heat to cold reservoir at T_C

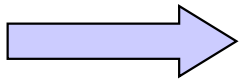


Work Done by a Heat Engine

The heat engine works in a *cyclic* process,

$$\Delta U = 0$$

1st Law gives,
$$\Delta U = Q_{net} - W = 0$$



$$Q_{net} = W$$

where,
$$Q_{net} = Q_H + Q_C = |Q_H| - |Q_C|$$

explicit signs for heats

Efficiency for a Heat Engine

- Thermal Efficiency e is defined as the *ratio* of the mechanical energy output to the heat energy input,

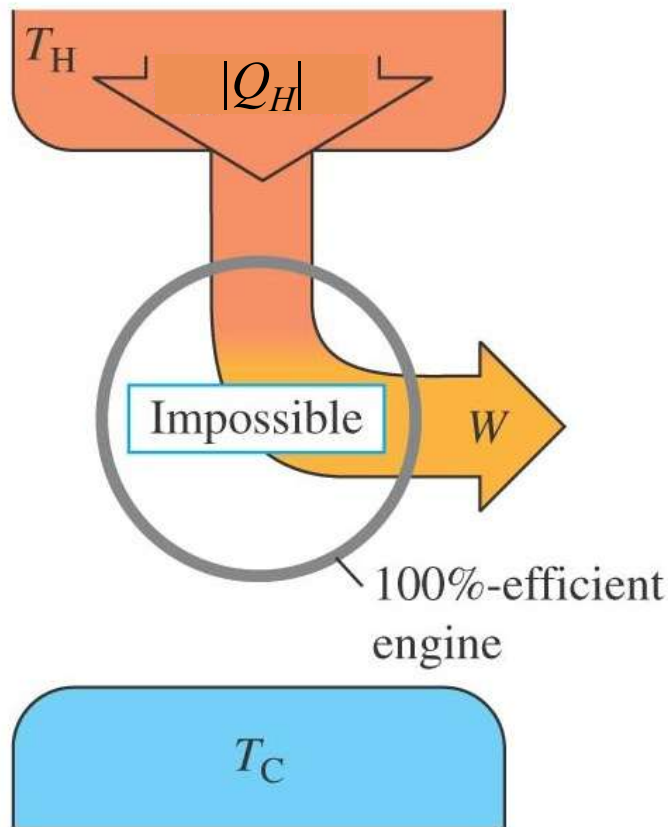
$$e = \frac{W}{Q_H} = \frac{\text{what you get out}}{\text{what you put in}}$$

Substituting $W = Q_H + Q_C$, we have

$$e = \frac{W}{Q_H} = \frac{Q_H + Q_C}{Q_H} = 1 + \frac{Q_C}{Q_H} = 1 - \left| \frac{Q_C}{Q_H} \right|$$

using explicit signs here

A “perfect” (100% efficient) heat engine



A “perfect” heat engine means 100% efficiency ($e=1$). This means that

$$e = 1 - \frac{|Q_C|}{|Q_H|} \rightarrow 1 \text{ means } Q_C = 0$$

All heat absorbed from reservoir T_H is converted into mechanical work W . No residual heat is released back.

The Kelvin-Planck’s statement of the 2nd Law does *not* allow this !



~~$$e_{\text{perfect}} = 1$$~~

and

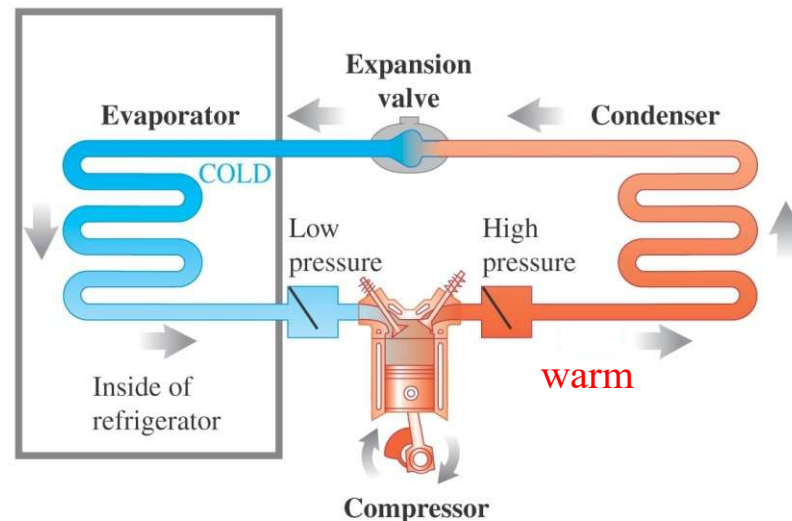
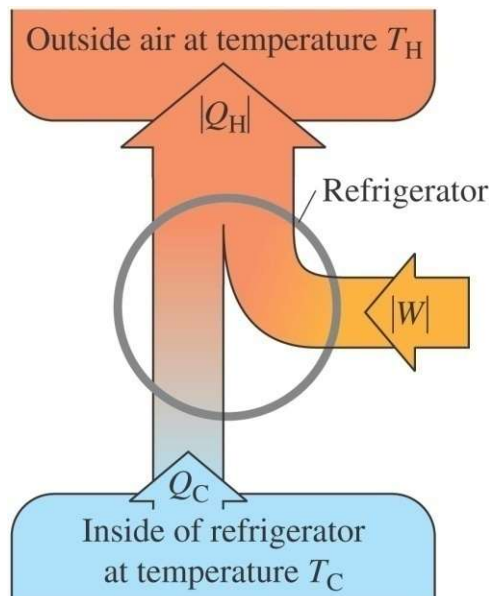
$$e_{\text{realistic}} < 1$$

D_{drinking bird}

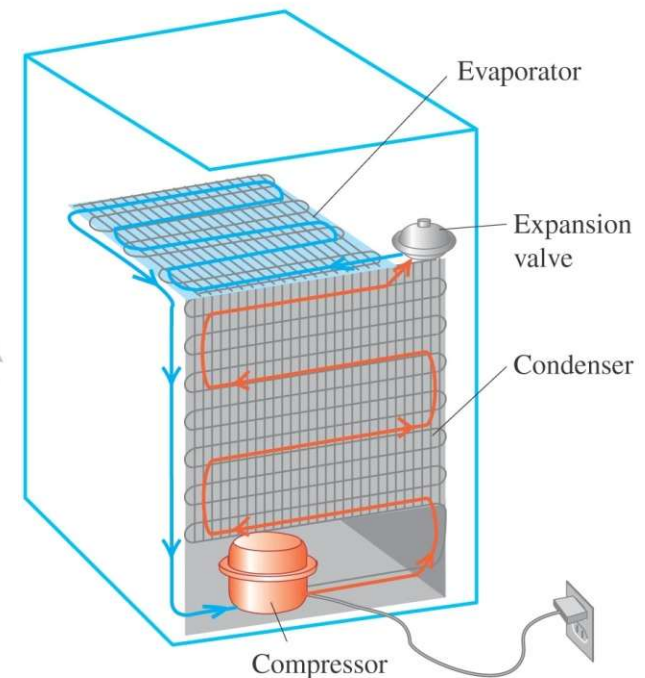
Refrigerators

Refrigerators are basically heat engine running in *reverse*.

- Heat from inside the refrigerator (cold T reservoir) is absorbed and released into the room (high T reservoir) with the *input* of mechanical work.




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Refrigerators

From 1st Law, $\Delta U_{cycle} = 0 = (|Q_C| - |Q_H|) - (-|W|)$
 $|Q_H| = |Q_C| + |W|$  explicit signs

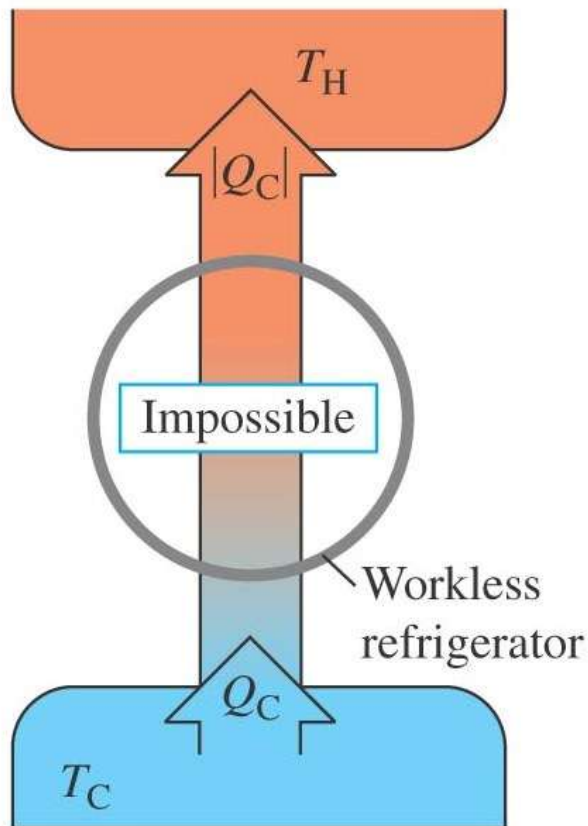
(Note: we have put in the explicit signs according to our sign convention.)

- ✓ $|Q_C|$ (absorbed) → positive
- ✓ $|Q_H|$ (released) → negative
- ✓ $|W|$ (work is done *on* working substance by motor) → negative

Coefficient of Performance for a Refrigerator

$$K = \frac{\text{what you get}}{\text{what you put in}} = \frac{|Q_C|}{|W|} = \frac{|Q_C|}{|Q_H| - |Q_C|}$$

A “perfect” Refrigerator



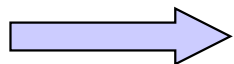
$$K = \frac{|Q_C|}{|W|} = \frac{|Q_C|}{|Q_H| - |Q_C|}$$

A “perfect” refrigerator means ($K = \infty$).
This means that

$$|Q_H| = |Q_C| \quad \text{or} \quad |W| = 0$$

No mechanical work W is needed to transfer heat from the cold reservoir to the hot reservoir.

The Clausius’s statement of the 2nd Law does *not* allow this !



$$K_{realistic} < \infty$$



2nd Law, Disorder, & Available Energy

Two Forms of Energy in any Thermal Process:

Internal Energy

In the *Kinetic-Molecular Model*, this consists of the KE and PE associated with all the *randomly* moving microscopic molecules.

(One typically cannot control the individual *random* motions of all these molecules.)

Macroscopic Mechanical Energy

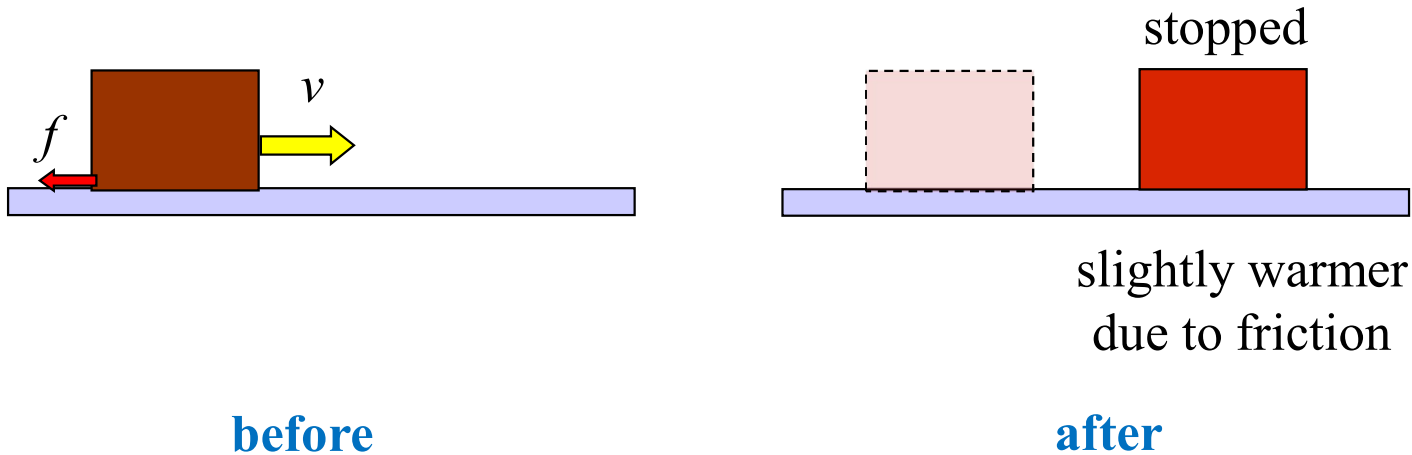
The piston's motion in an automobile engine results from the overall *coordinated* macroscopic motion of the molecules.

(Energy associated with this coordinated [*ordered*] motion can be used for useful work.)

2nd Law, Disorder, & Available Energy

In a natural process (a block sliding to a stop),

e.g.



- The coordinated motion of the block is converted into the slightly *more* agitated random motions of the molecules in the block and table.
- **Macroscopic Mechanical Energy** (KE of the block) is converted into **Internal Energy** through *heat* as a result of friction.

2nd Law, Disorder, & Available Energy

- Now consider the possibility of *reverse* situation... do we typically see a group of randomly moving molecules **all** push a block in a coordinated fashion?
- NO. In other words, one typically cannot *completely* convert the internal energy of a system back into macroscopic mechanical energy.
- However, this does not mean that internal energy is not accessible. An *Heat Engine* is exactly the machine that can perform this conversion but only *partially*.

The 2nd Law of Thermodynamics is basically a statement limiting the *availability* of internal energy for useful mechanical work.

The Carnot Cycle (Most Efficient Heat Engine)

- A *reversible* cycle described by Sadi Carnot in 1824.
- The Carnot Theorem gives the *theoretical limit* to the thermal efficiency of *any* heat engine.
- The Carnot cycle consists of:
 - A cycle operating between two temperatures: T_H and T_C
 - 2 reversible isothermal processes in which $Q \pm$
 - 2 reversible adiabatic processes in which $T_H \leftrightarrow T_C$
 - An Ideal Gas as its working substance

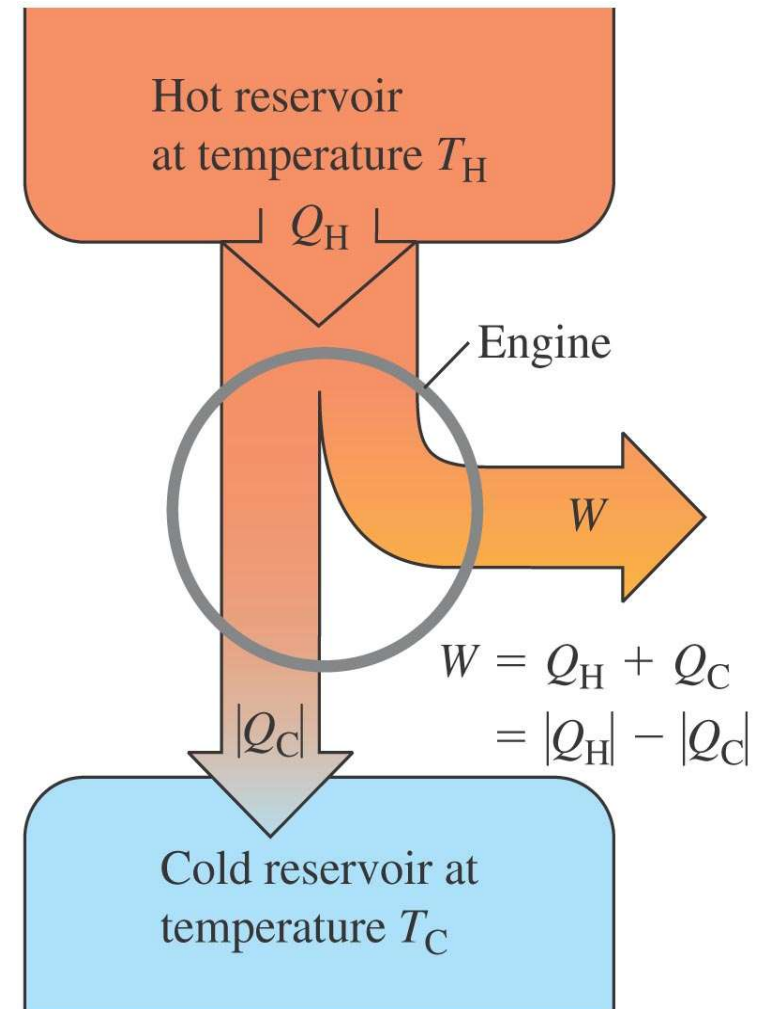
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Mechanical work is done by engine

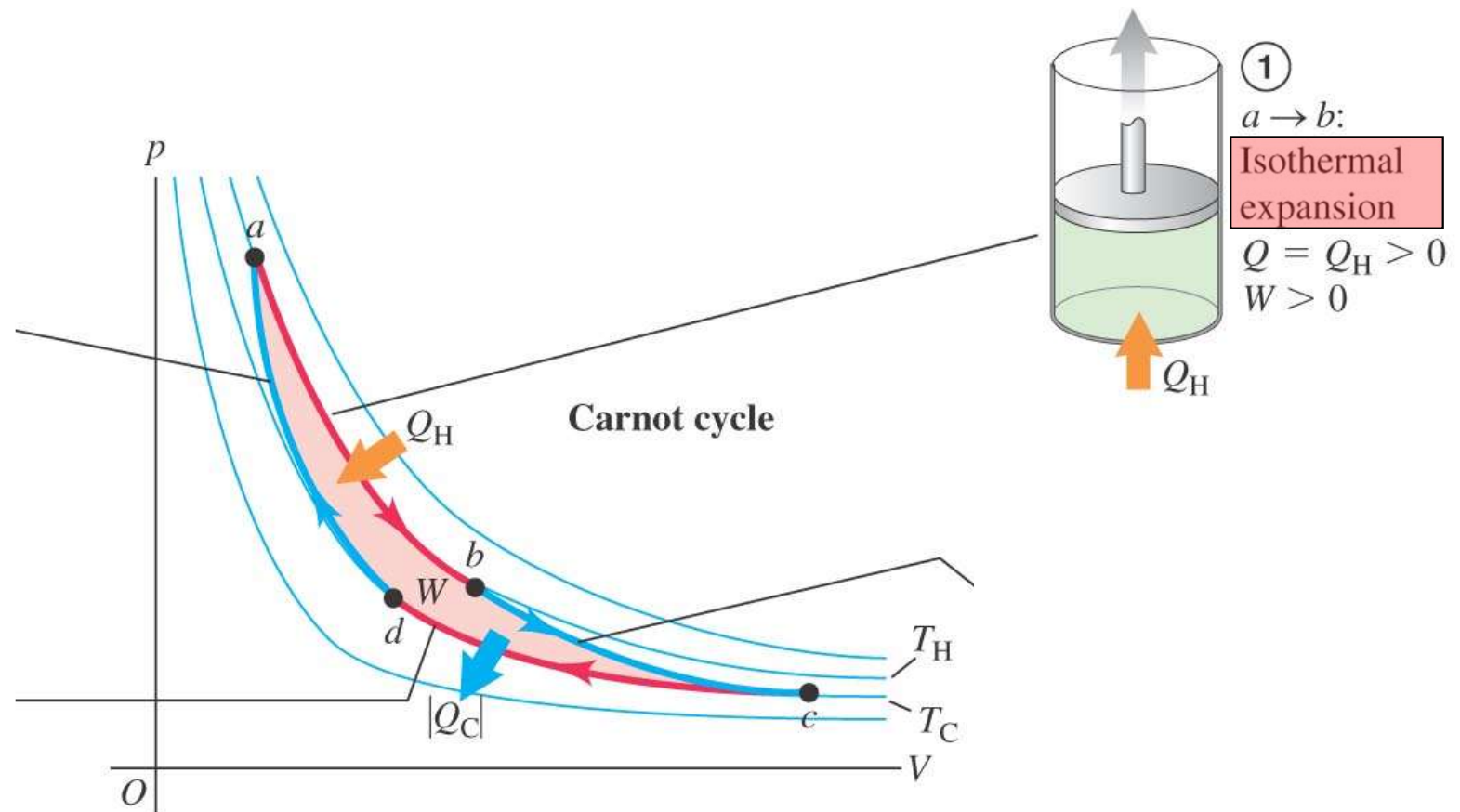
Engine releases residual heat to cold reservoir at T_C



Steps of the Carnot Cycle

[animation](#)

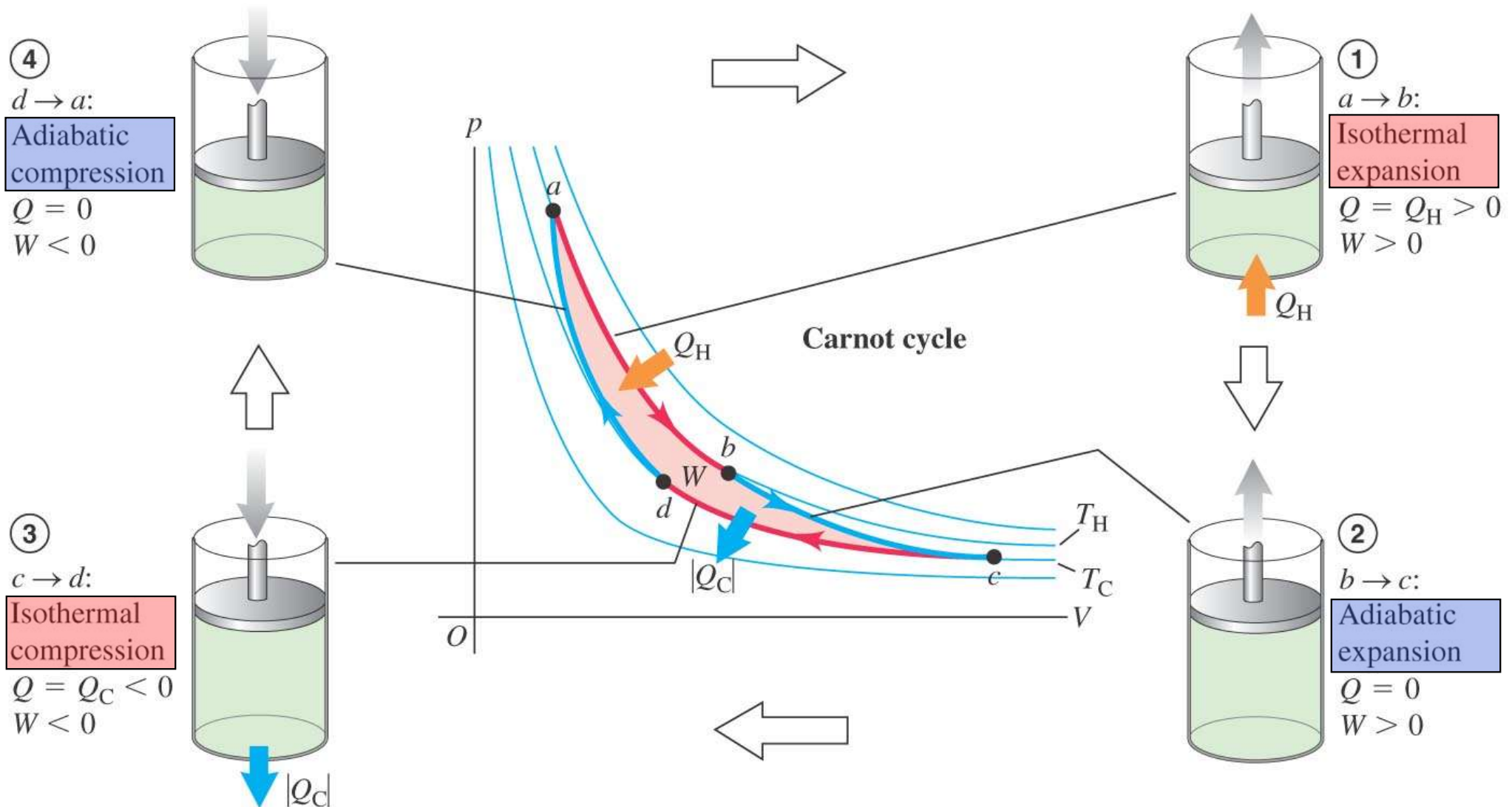
http://galileoandstein.physics.virginia.edu/more_stuff/flashlets/carnot.htm



Steps of the Carnot Cycle

[animation](#)

http://galileoandstein.physics.virginia.edu/more_stuff/flashlets/carnot.htm



Details on the Carnot Cycle

The isothermal expansion ($a \rightarrow b$) and compression ($c \rightarrow d$):

$$\Delta U_{\text{isothermal}} = 0$$

(T is constant and $U(T)$ is a function of T only for an Ideal Gas.)

$$|Q_H| = W_{ab} = nRT_H \ln\left(\frac{V_b}{V_a}\right)$$

($a \rightarrow b$: isothermal expansion)

$$|Q_C| = |W_{cd}| = \left| nRT_C \ln\left(\frac{V_d}{V_c}\right) \right| = nRT_C \ln\left(\frac{V_c}{V_d}\right) \quad (V_c > V_d)$$

($c \rightarrow d$: isothermal compression)

Details on the Carnot Cycle

The adiabatic expansion ($b \rightarrow c$) and compression ($d \rightarrow a$):

$$Q_{adiabatic} = 0 \quad (\text{by definition})$$

From Section 19.8, we learned that $TV^{\gamma-1} = \text{const}$ for adiabatic processes.

$$T_H V_b^{\gamma-1} = T_C V_c^{\gamma-1} \quad (b \rightarrow c: \text{adiabatic expansion})$$

$$T_H V_a^{\gamma-1} = T_C V_d^{\gamma-1} \quad (d \rightarrow a: \text{adiabatic compression})$$

Dividing these two equations gives,

$$\frac{V_b}{V_a} = \frac{V_c}{V_d}$$

Efficiency of the Carnot Cycle

From definition, we have $e = 1 - \frac{|Q_C|}{|Q_H|}$

Using our results for Q_C and Q_H from the *isothermal* processes,

$$e = 1 - \frac{nRT_C \ln(V_c/V_d)}{nRT_H \ln(V_b/V_a)} = 1 - \frac{T_C \ln(V_c/V_d)}{T_H \ln(V_b/V_a)}$$

Then, from the *adiabatic* processes, we have $V_b/V_a = V_c/V_d$

$$\longrightarrow \frac{\ln(V_c/V_d)}{\ln(V_b/V_a)} = 1 \longrightarrow e \equiv 1 - \frac{|Q_C|}{|Q_H|} = 1 - \frac{T_C}{T_H} \quad (T \text{ must be in K})$$

(Carnot Cycle only)

Efficiency of the Carnot Cycle

$$e_{carnot} = 1 - \frac{T_C}{T_H} \quad (\text{Carnot Cycle})$$

General Comments:

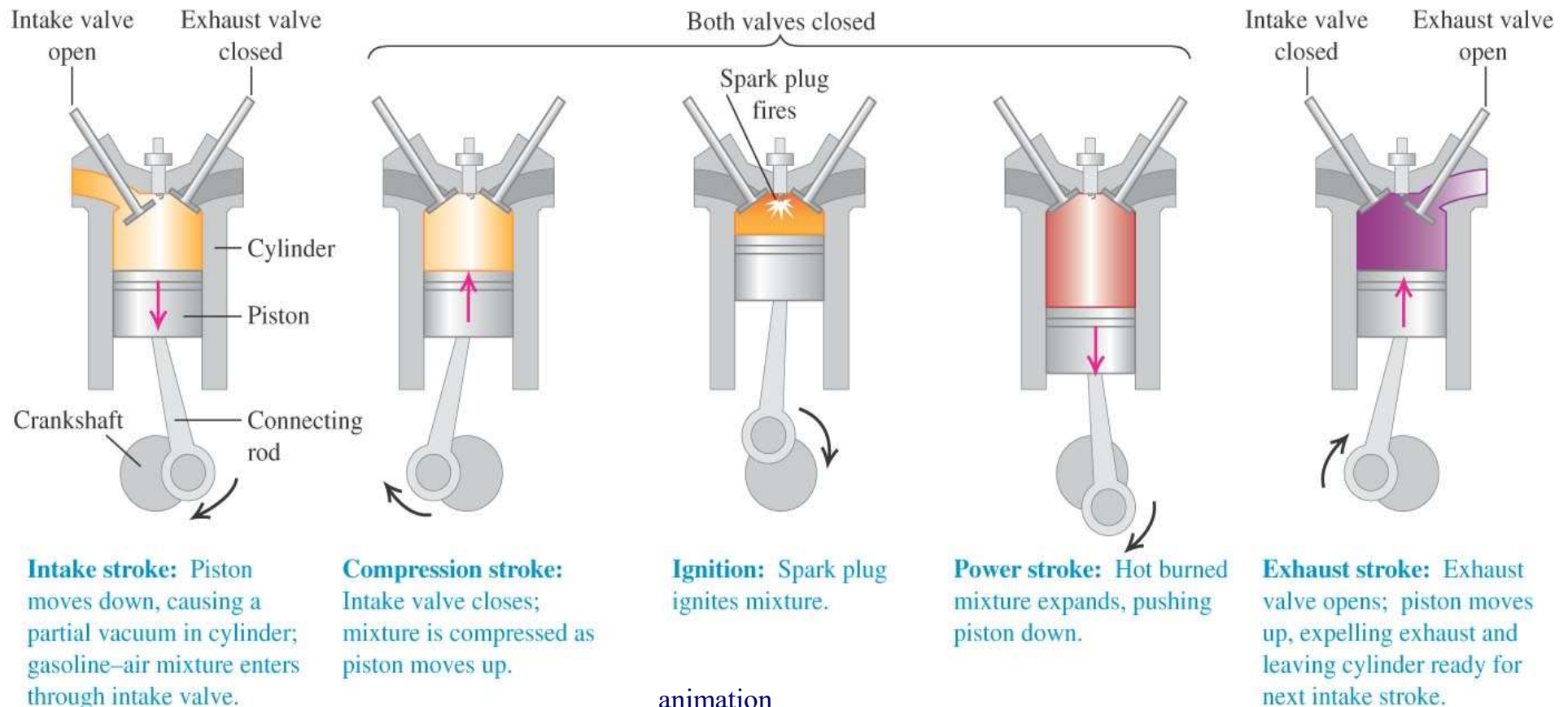
- Higher efficiency if either T_C is lower and/or T_H is higher.
- For any *realistic* thermal process, the cold reservoir is far above absolute zero, i.e., $T_C > 0$.
- Thus, a *realistic* e is *strictly less* than 1! (No 100% efficient heat engine)
- *Realistic* heat engines must take in energy from the high T reservoir for the work that it produces **AND** some heat energy must be *released* back to the lower T reservoir.

(Kelvin-Planck's Statement)

Internal Combustion Engine

The Otto Cycle

A fuel vapor can be compressed, then detonated to rebound the cylinder, doing useful work.

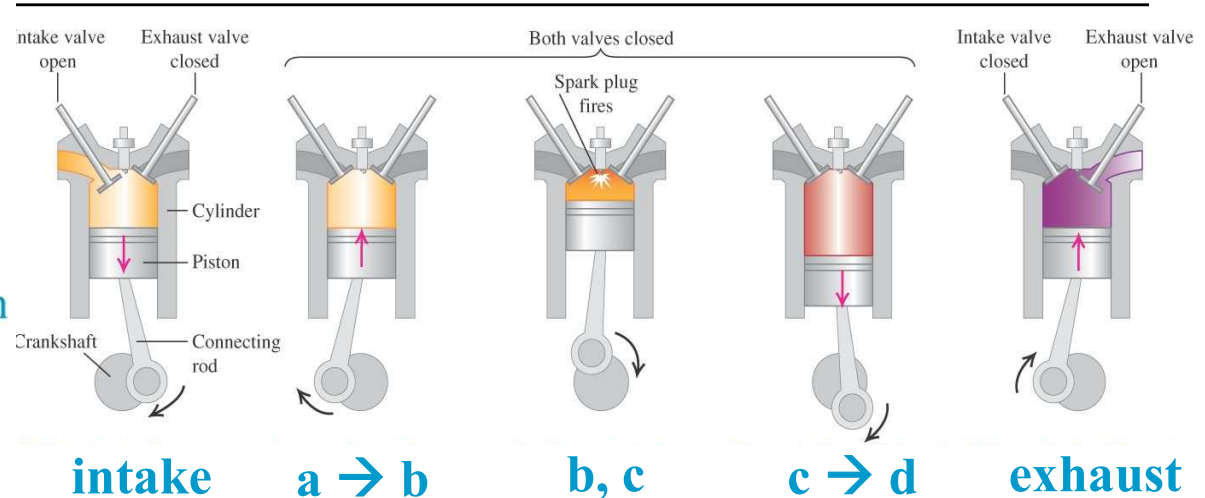
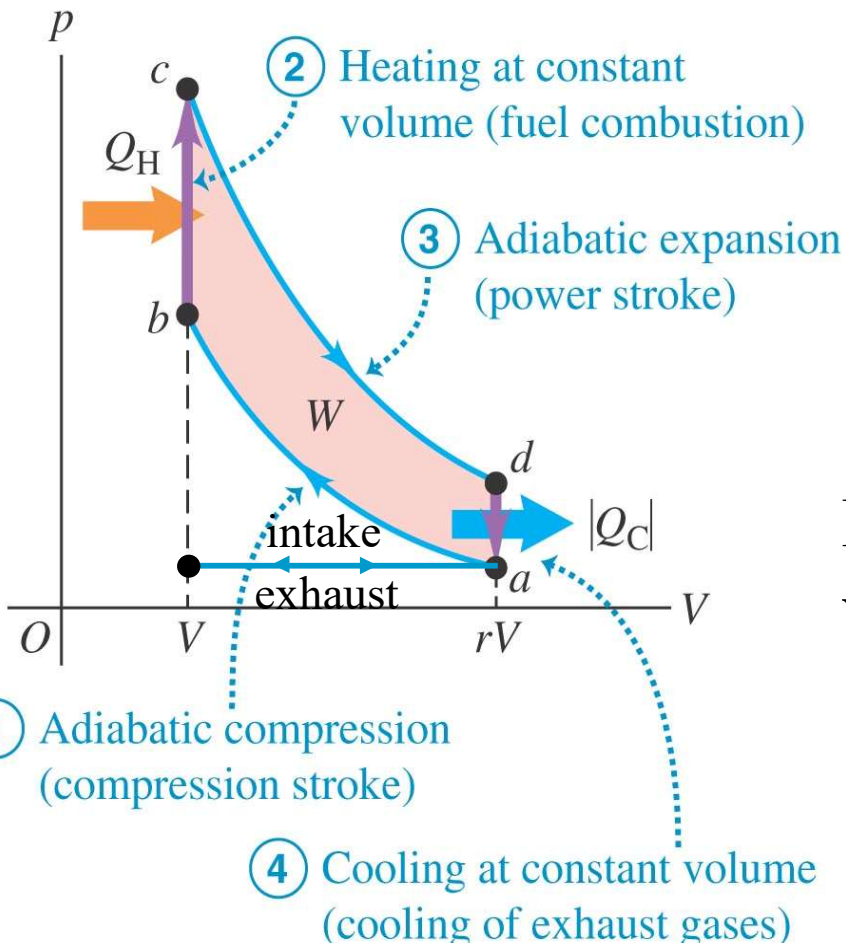


[animation](#)

<http://auto.howstuffworks.com/engine1.htm>

The Otto Cycle

Otto cycle



For the two constant V processes: 2 and 4, we can calculate,

$$Q_H = \Delta U_2 = nC_V (T_c - T_b) > 0$$

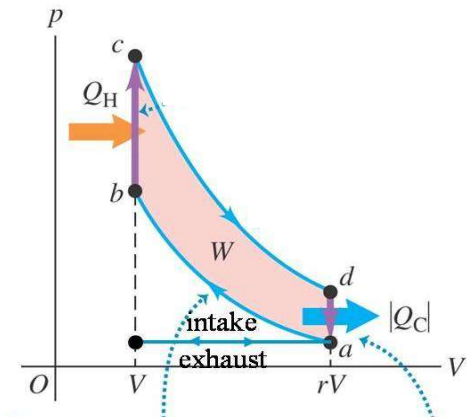
$$Q_C = \Delta U_4 = nC_V (T_a - T_d) < 0$$

r is the compression ratio (8 to 13)

The Otto Cycle

Applying the definition of *efficiency*,

$$e = 1 + \frac{Q_C}{Q_H} = 1 + \frac{nC_V (T_a - T_d)}{nC_V (T_c - T_b)} = 1 + \frac{T_a - T_d}{T_c - T_b}$$



Now, we can utilize the two adiabatic processes: 1 and 3,

$$T_a V_a^{\gamma-1} = T_b V_b^{\gamma-1} \quad \text{and} \quad T_c V_c^{\gamma-1} = T_d V_d^{\gamma-1}$$

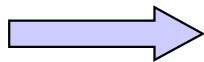
$$T_a (rV)^{\gamma-1} = T_b V^{\gamma-1} \quad T_c V^{\gamma-1} = T_d (rV)^{\gamma-1}$$

$$T_a r^{\gamma-1} = T_b \quad T_c = T_d r^{\gamma-1}$$

The Otto Cycle

Substituting T_b and T_c into the efficiency equation, we have,

$$e = 1 + \frac{T_a - T_d}{T_c - T_b} = 1 + \frac{T_a - T_d}{T_d r^{\gamma-1} - T_a r^{\gamma-1}} = 1 - \frac{T_a - T_d}{(T_a - T_d) r^{\gamma-1}}$$



$$e = 1 - \frac{1}{r^{\gamma-1}}$$

Using a typical value for the compression ratio $r = 8$ and $\gamma = 1.40$ gives,

$$e = 0.56 \quad (\text{or } 56\%)$$

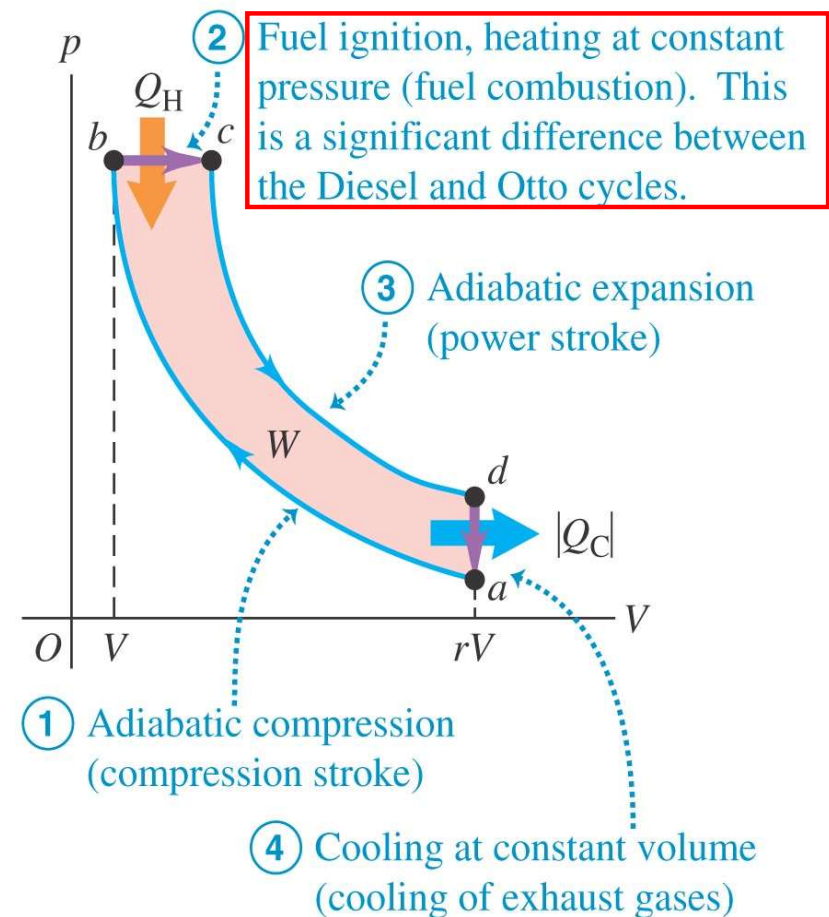
Note: This is a theoretical value.
Realistic gasoline engine typically
has $e \sim 35\%$.

The Diesel Cycle

Key difference:

- No fuel in cylinder at the beginning of the compression stroke (process 1)
- Fuel is injected only moments before ignition in the power stroke
- No fuel until the end of the adiabatic compression can avoid pre-ignition
- Compression ratio r value can be higher (15 to 20)
- Higher temperature can be reached during the adiabatic compression
- Higher e and no need for spark plugs

Diesel cycle



Entropy

Recall from a Carnot Cycle, we have derived the following relationship:

$$\frac{|Q_C|}{|Q_H|} = \frac{T_C}{T_H} \quad \longrightarrow \quad \frac{|Q_H|}{T_H} + \frac{-|Q_C|}{T_C} = 0$$

Formally, we can rewrite this as,

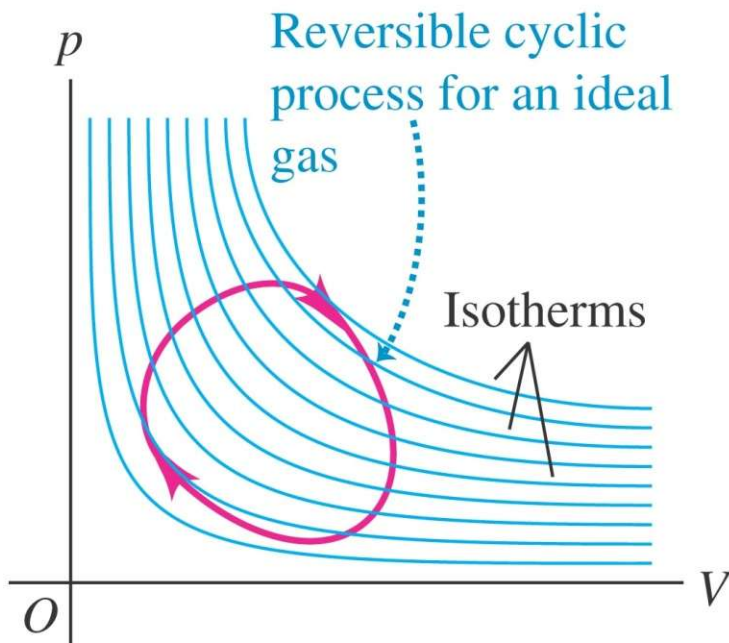
$$\sum_{\text{cycle}} \frac{Q}{T} = 0$$

(We have absorbed the explicit sign back into the variable Q .)

where Q represents the heat absorbed/released along the isotherm at temp T .

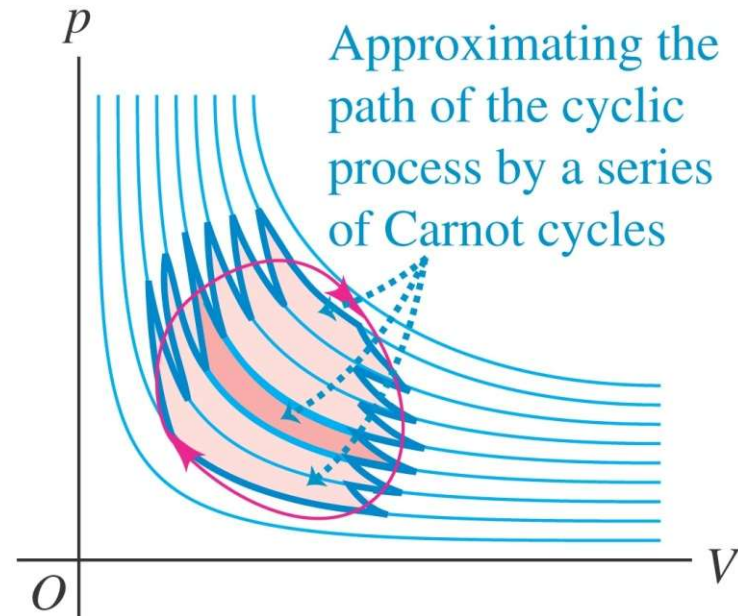
Entropy

Now consider any reversible cycles...



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Any reversible cycles can be approximated as a series of Carnot cycles !



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Entropy

This suggests that the following generalization to be true for any reversible cycles,

$$\oint_{\text{cycle}} \frac{dQ_r}{T} = 0$$

where,

dQ_r is the infinitesimal heat absorbed/released by the system at an infinitesimal *reversible* step at temp T .

\oint_{cycle} denotes the integration evaluated over one complete cycle.

Entropy

We have seen this property previously,

$$\oint_{\text{cycle}} dU = 0 \quad \text{Changes in the internal energy } U \text{ over a closed cycle is zero!}$$

This is a consequence of the fact that U is a **state variable** and dU for any processes depends on the initial and final states only.

Thus the result $\oint_{\text{cycle}} \frac{dQ_r}{T} = 0$ indicates that there is another **state variable** S such that,

$$dS \equiv \frac{dQ}{T}$$

and

$$\oint_{\text{cycle}} dS = 0$$

This new **state variable** S is called the **entropy** of the system.