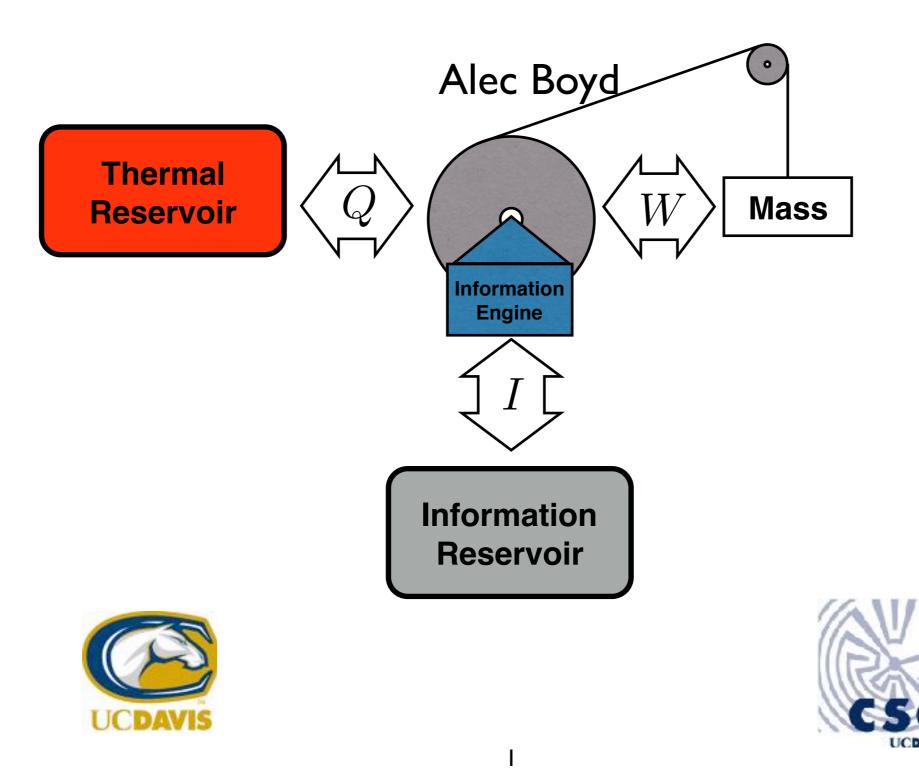
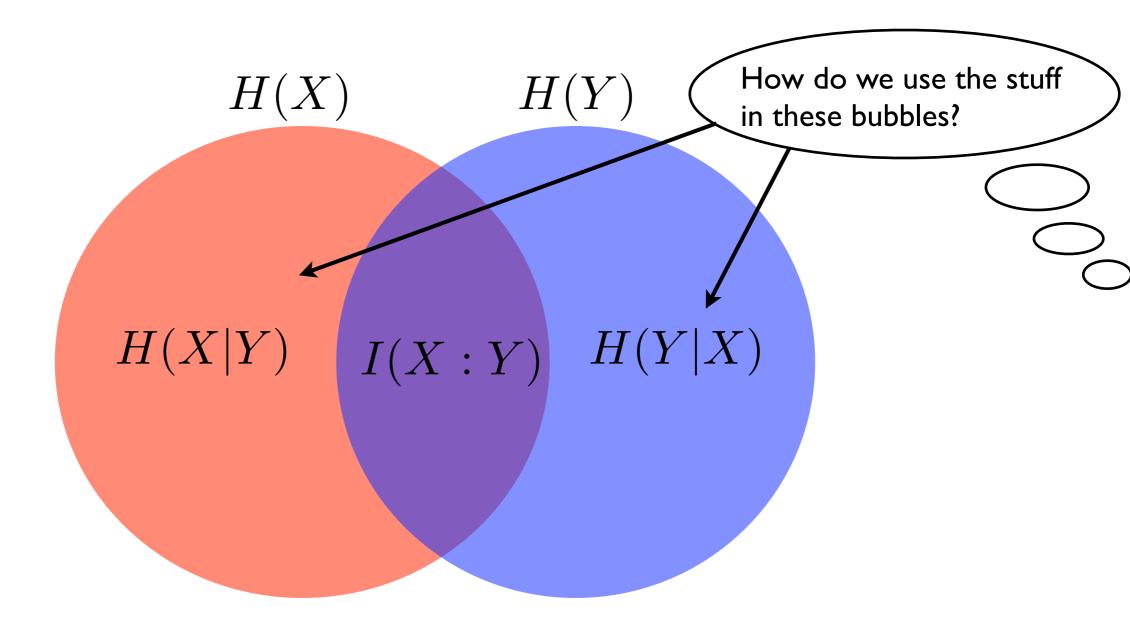
Thermodynamics of Information Part I

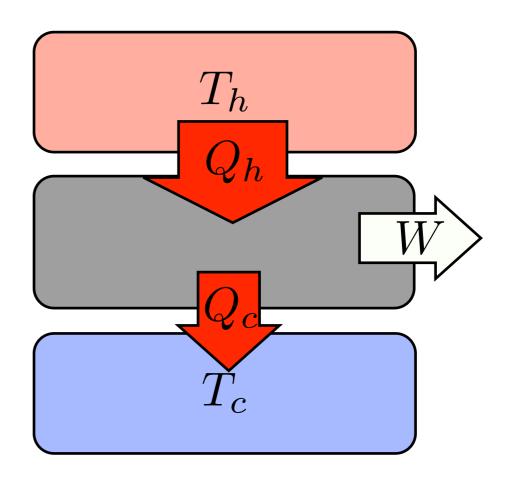


What does information do?



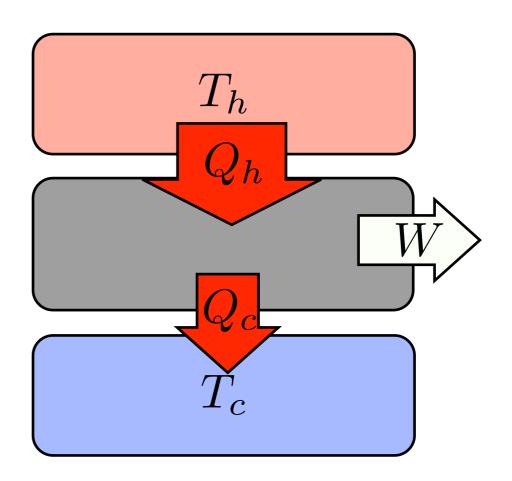
Thermodynamics treats physical systems many elements.

Relevant quantities: I) Energy



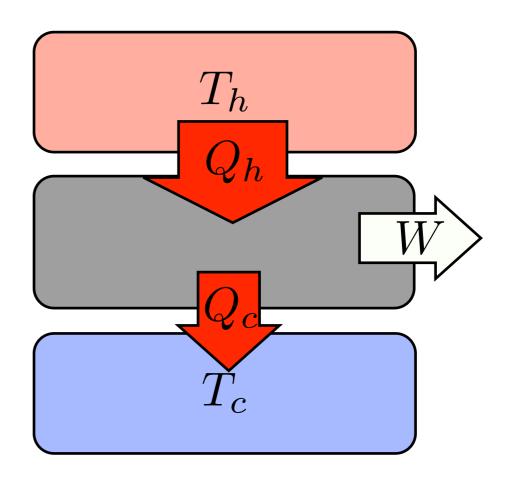
Thermodynamics treats physical systems many elements.

Relevant quantities: 1) Energy 2) Work



Thermodynamics treats physical systems many elements.

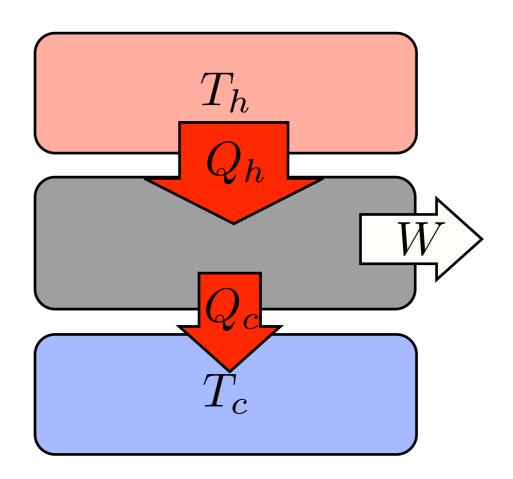
Relevant quantities: I) Energy 2) Work 3) Heat



Thermodynamics treats physical systems many elements.

Relevant quantities:

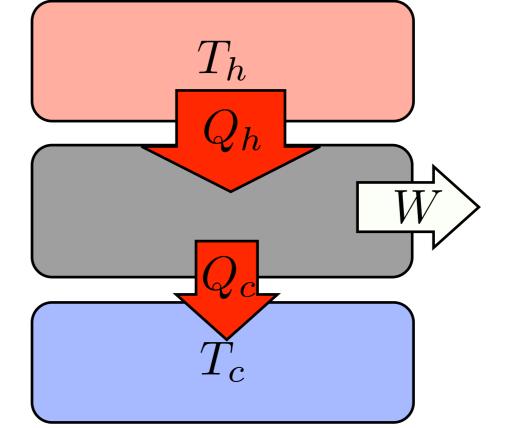
I) Energy
 Work
 Heat
 Temperature



Thermodynamics treats physical systems many elements.

Relevant quantities:

I) Energy
 Work
 Heat
 Temperature
 Entropy



Physical system states:

$$\mathcal{X} = \{x\}$$

Physical system states:

$$\mathcal{X} = \{x\}$$

Probability of physical state at time t:

 $\Pr(X_t = x)$

Physical system states:

$$\mathcal{X} = \{x\}$$

Probability of physical state at time t:

 $\Pr(X_t = x)$

Thermodynamic entropy (Gibbs) of physical system: $S[X_t] = -k_B \sum_{x \in \mathcal{X}} \Pr(X_t = x) \ln \Pr(X_t = x)$

Physical system states:

$$\mathcal{X} = \{x\}$$

Probability of physical state at time t:

 $\Pr(X_t = x)$

Thermodynamic entropy (Gibbs) of physical system:

$$S[X_t] = -k_B \sum_{x \in \mathcal{X}} \Pr(X_t = x) \ln \Pr(X_t = x)$$

Shannon entropy of physical system:

$$H[X_t] = -\sum_{x \in \mathcal{X}} \Pr(X_t = x) \log_2 \Pr(X_t = x)$$

Physical system states:

$$\mathcal{X} = \{x\}$$

Probability of physical state at time t:

 $\Pr(X_t = x)$

Thermodynamic entropy (Gibbs) of physical system:

$$S[X_t] = -k_B \sum_{x \in \mathcal{X}} \Pr(X_t = x) \ln \Pr(X_t = x)$$

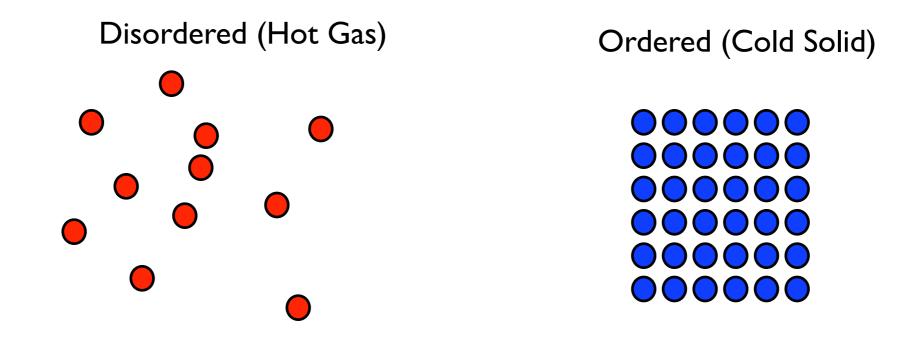
Shannon entropy of physical system:

$$H[X_t] = -\sum_{x \in \mathcal{X}} \Pr(X_t = x) \log_2 \Pr(X_t = x)$$

 $S[X_t] = k_B \ln 2 H[X_t]$

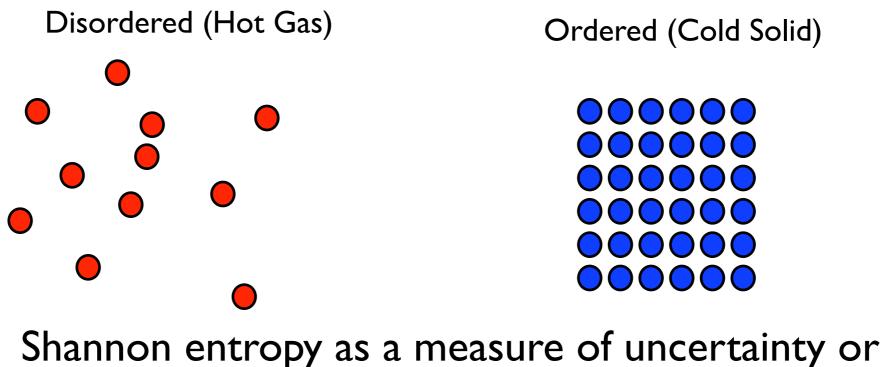
Entropy Comparison

Thermodynamic entropy as a measure of disorder.



Entropy Comparison

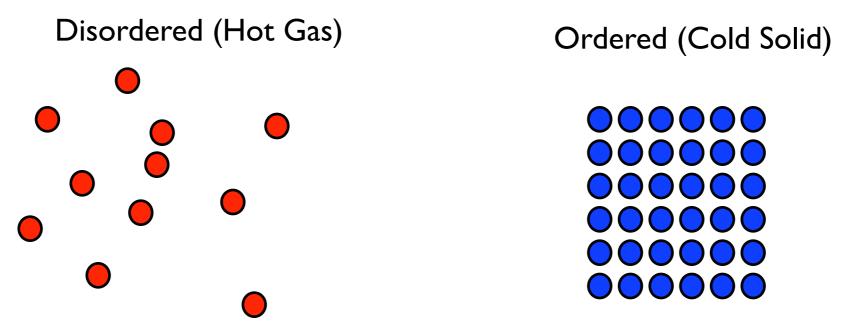
Thermodynamic entropy as a measure of disorder.



unpredictability.

Entropy Comparison

Thermodynamic entropy as a measure of disorder.



Shannon entropy as a measure of uncertainty or unpredictability.

String of measurements of unpredictable variable

...11011101100101111000000001...

String of measurements of predictable variable

...10101010101010101010101...

Classical Physics

Physical states described by positions q_i and their conjugate momenta p_i :

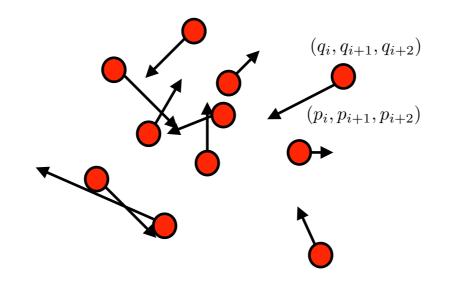
$$\mathcal{X} = \{x\} = \{(q_1, q_2, \dots, q_N, p_1, p_2, \dots, p_N)\}$$

Classical Physics

Physical states described by positions q_i and their conjugate momenta p_i :

$$\mathcal{X} = \{x\} = \{(q_1, q_2, ..., q_N, p_1, p_2, ..., p_N)\}$$

Can be thought of as a collection of particles:

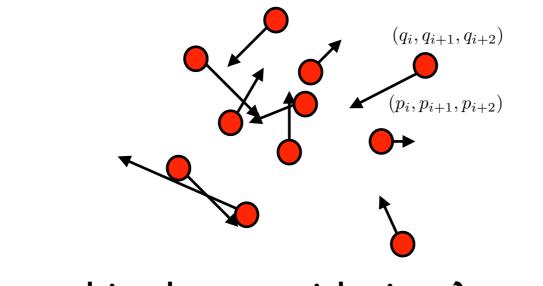


Classical Physics

Physical states described by positions q_i and their conjugate momenta p_i :

$$\mathcal{X} = \{x\} = \{(q_1, q_2, ..., q_N, p_1, p_2, ..., p_N)\}$$

Can be thought of as a collection of particles:



How does this change with time?

First Law of Thermodynamics

Energy of each state specified by Hamiltonian function of state

 $E(x) = \mathcal{H}(\vec{q}, \vec{p})$

First Law of Thermodynamics

Energy of each state specified by Hamiltonian function of state

$$E(x) = \mathcal{H}(\vec{q}, \vec{p})$$

Position changes as velocity and momentum changes as the force:

$$\frac{dq_i}{dt} = \partial_{p_i} \mathcal{H}(\vec{q}, \vec{p}) \qquad \frac{dp_i}{dt} = -\partial_{q_i} \mathcal{H}(\vec{q}, \vec{p})$$

First Law of Thermodynamics

Energy of each state specified by Hamiltonian function of state

$$E(x) = \mathcal{H}(\vec{q}, \vec{p})$$

Position changes as velocity and momentum changes as the force:

$$\frac{dq_i}{dt} = \partial_{p_i} \mathcal{H}(\vec{q}, \vec{p}) \qquad \frac{dp_i}{dt} = -\partial_{q_i} \mathcal{H}(\vec{q}, \vec{p})$$

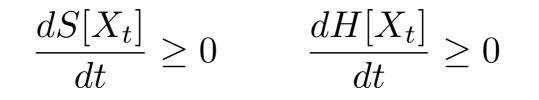
Equations of motion conserve energy

$$\Delta_{t \to t+\tau} E = E(x(t+\tau)) - E(x(t)) = 0$$
$$\frac{dE(x(t))}{dt} = 0$$

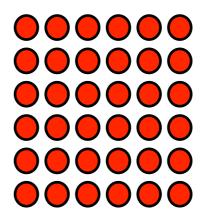
Entropy increases in isolated systems:

$$\frac{dS[X_t]}{dt} \ge 0 \qquad \quad \frac{dH[X_t]}{dt} \ge 0$$

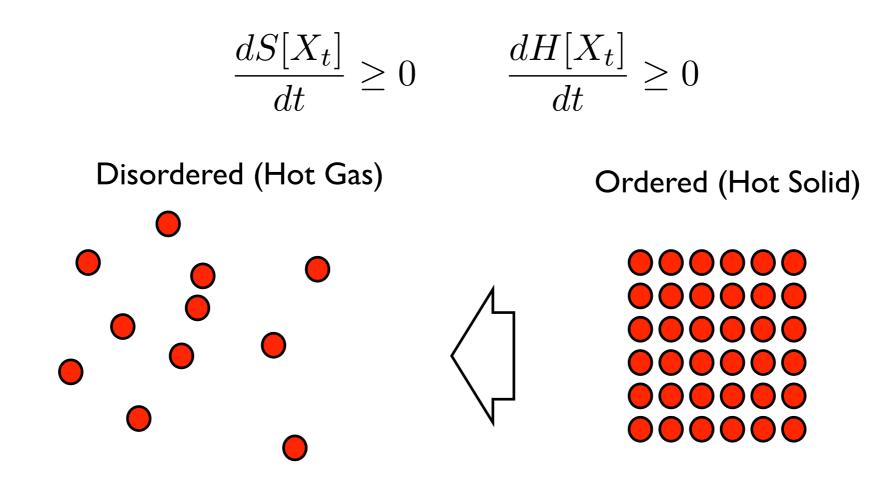
Entropy increases in isolated systems:



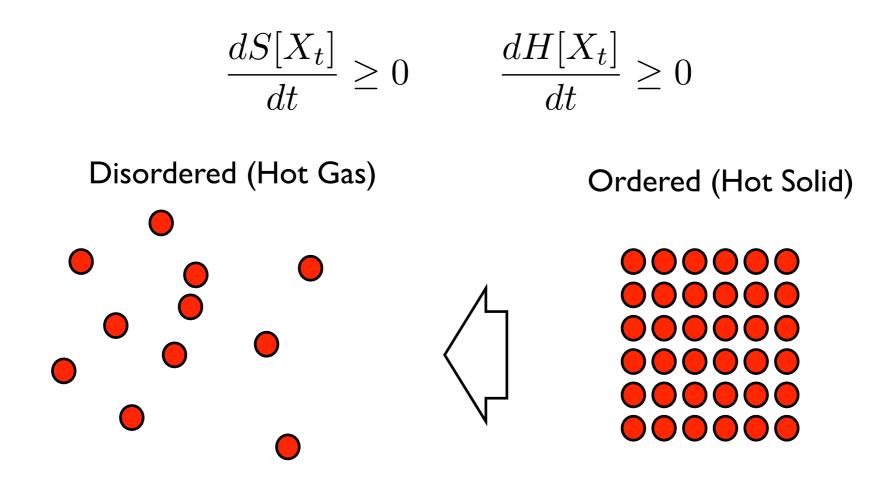
Ordered (Hot Solid)



Entropy increases in isolated systems:



Entropy increases in isolated systems:



Example another and another and another

Energy and Entropy

According to the first law and second law of thermodynamics together, the microstates evolve towards the maximum entropy distribution over a fixed energy E, called the microcanonical ensemble:

Energy and Entropy

According to the first law and second law of thermodynamics together, the microstates evolve towards the maximum entropy distribution over a fixed energy E, called the microcanonical ensemble:

$$\Pr(X^{\text{eq}} = x) = \delta_{E(x),E} / \Omega(E)$$

where $\Omega(E)$ is the number of states with energy E.

Energy and Entropy

According to the first law and second law of thermodynamics together, the microstates evolve towards the maximum entropy distribution over a fixed energy E, called the microcanonical ensemble:

$$\Pr(X^{\text{eq}} = x) = \delta_{E(x),E} / \Omega(E)$$

where $\Omega(E)$ is the number of states with energy E.

This leads to an equilibrium estimate of entropy: $S[X^{eq}|E(x) = E'] = k_B \ln \Omega(E')$

Energy/Entropy vs.Temperature

What is temperature? -Average energy?

Energy/Entropy vs. Temperature

What is temperature? -Average energy?

Temperature of a system in equilibrium:

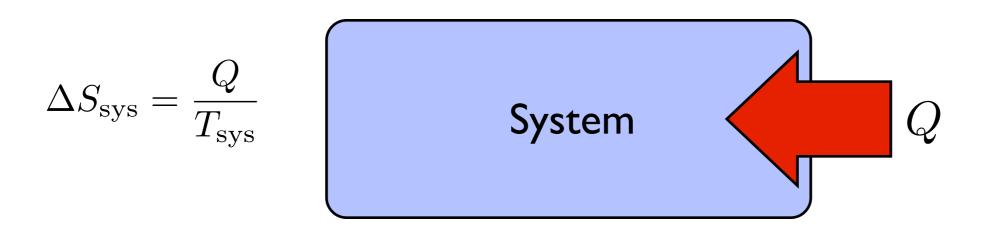
$$\frac{1}{T} = \frac{\partial S[X^{\rm eq}|E(x) = E']}{\partial E'}$$

Energy/Entropy vs.Temperature

What is temperature? -Average energy?

Temperature of a system in equilibrium:

$$\frac{1}{T} = \frac{\partial S[X^{\rm eq}|E(x) = E']}{\partial E'}$$

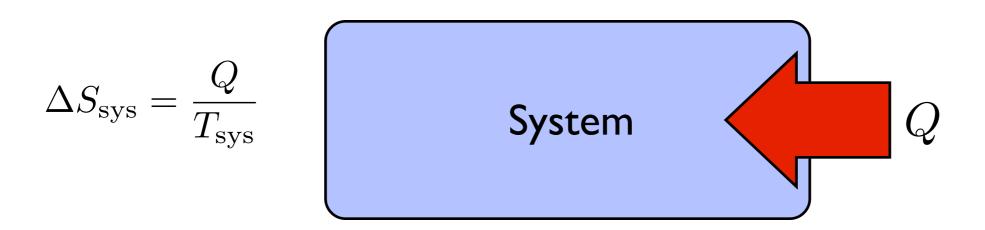


Energy/Entropy vs.Temperature

What is temperature? -Average energy?

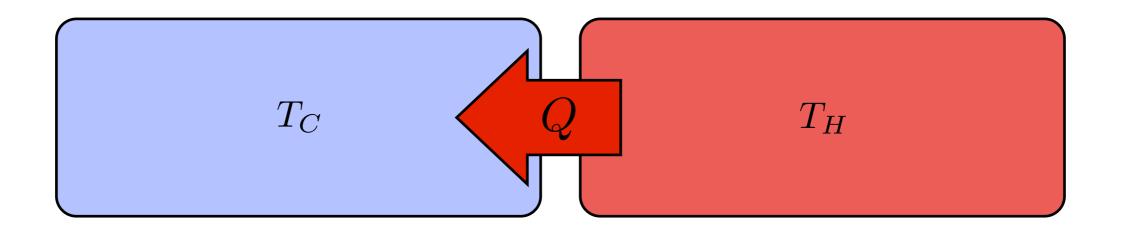
Temperature of a system in equilibrium:

$$\frac{1}{T} = \frac{\partial S[X^{\rm eq}|E(x) = E']}{\partial E'}$$

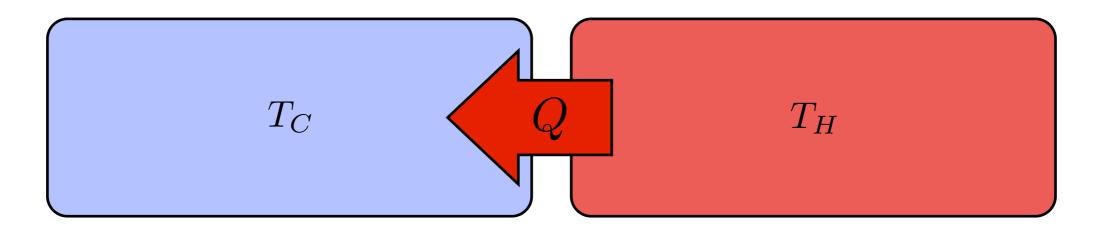


In most systems, every degree of freedom stores energy proportional to the temperature: $\langle E_{q_i} \rangle \approx \langle E_{p_i} \rangle \approx \frac{k_B T}{2}$

Entropy explains why heat diffuses from hot to cold: $T_H > T_C$

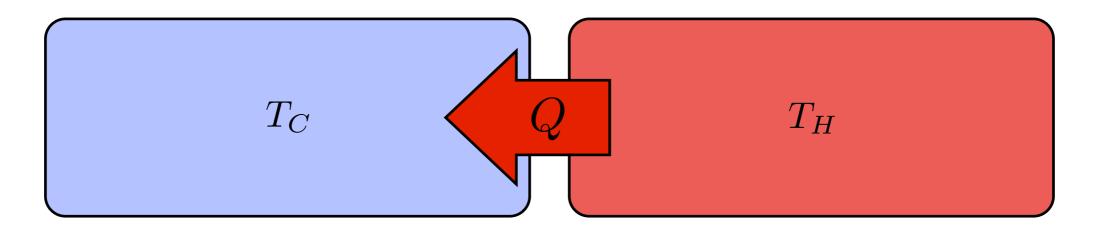


Entropy explains why heat diffuses from hot to cold: $T_H > T_C$



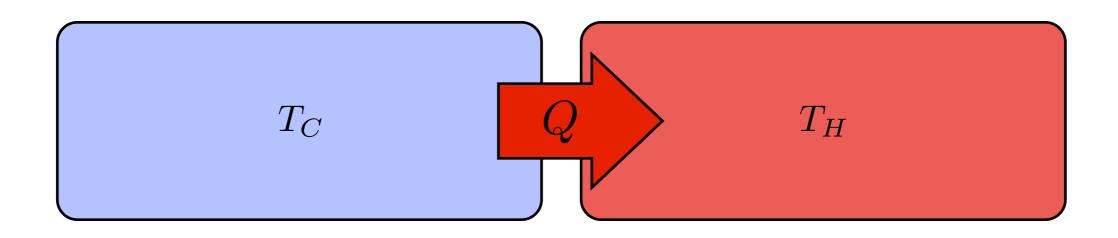
Total entropy increases by heat diffusion: $\Delta S_{\rm total} = Q/T_C - Q/T_H > 0$

Entropy explains why heat diffuses from hot to cold: $T_H > T_C$

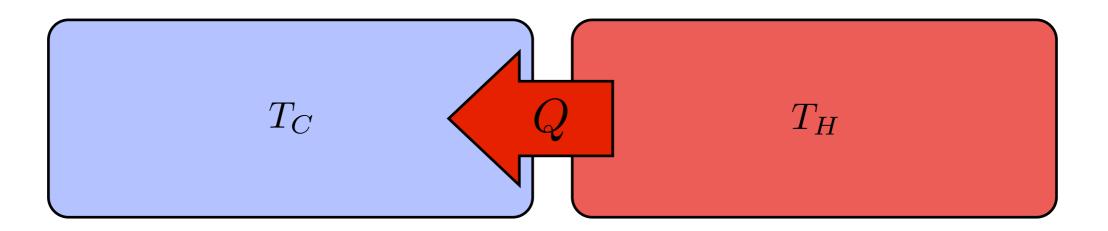


Total entropy increases by heat diffusion: $\Delta S_{\text{total}} = Q/T_C - Q/T_H > 0$

The reverse is impossible:



Entropy explains why heat diffuses from hot to cold: $T_H > T_C$



Total entropy increases by heat diffusion:

$$\Delta S_{\text{total}} = Q/T_C - Q/T_H > 0$$

The reverse is impossible:

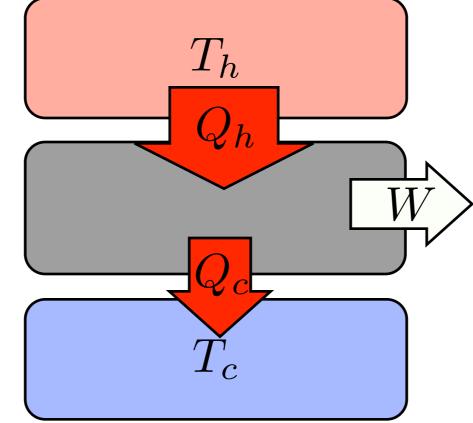
$$\Delta S_{\text{total}} = Q/T_H - Q/T_C < 0$$

$$T_C$$

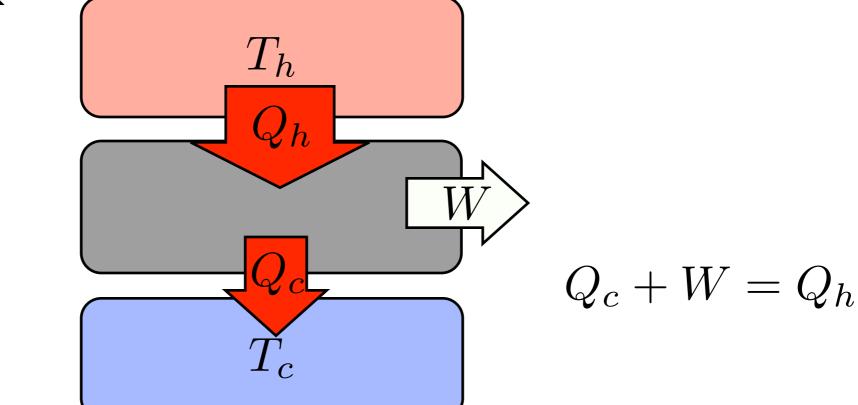
$$T_H$$

Some energy can be redirected to produce work

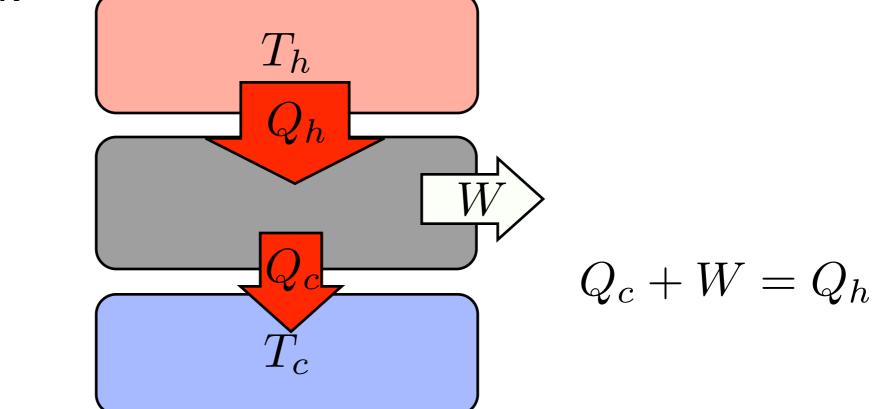
Some energy can be redirected to produce work



Some energy can be redirected to produce work

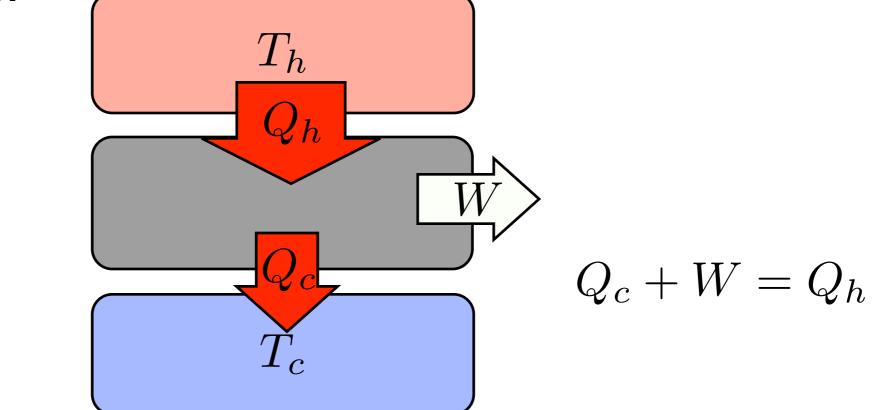


Some energy can be redirected to produce work



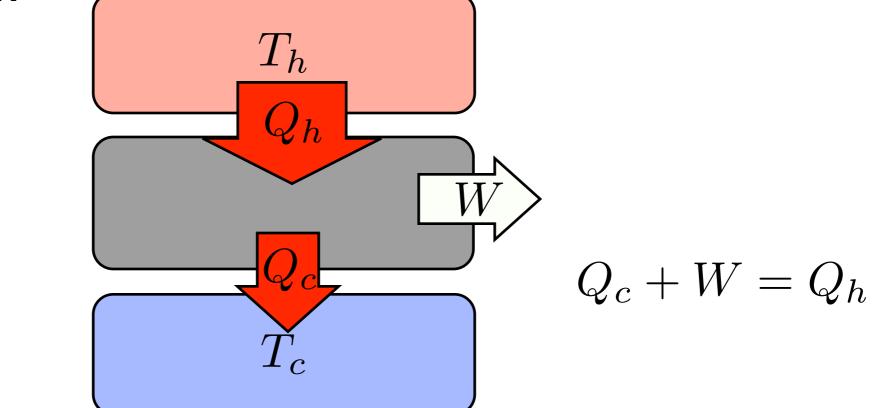
$$\Delta S_{total} = Q_c / T_c - Q_h / T_h \ge 0$$

Some energy can be redirected to produce work



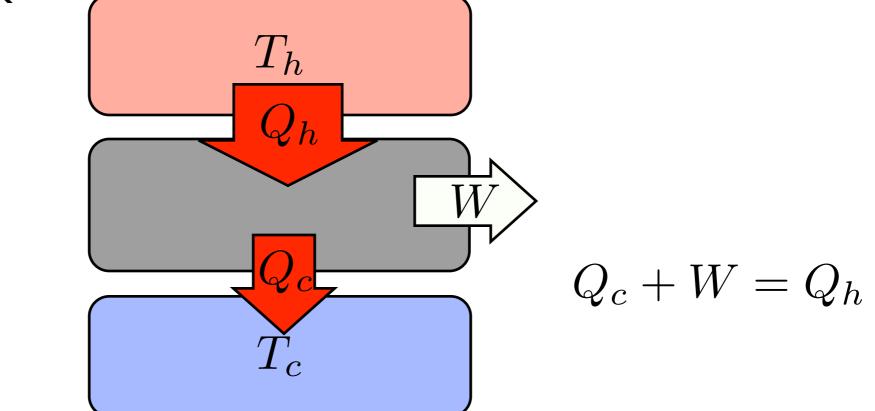
$$\Delta S_{total} = Q_c / T_c - Q_h / T_h \ge 0$$
$$\frac{Q_c}{Q_h} \ge \frac{T_c}{T_h}$$

Some energy can be redirected to produce work



$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$

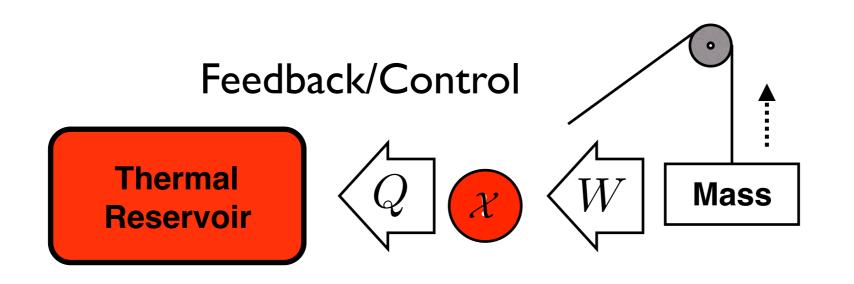
Some energy can be redirected to produce work



$$\begin{split} e &= \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h} \\ e &\leq 1 - \frac{T_c}{T_h} \end{split}$$

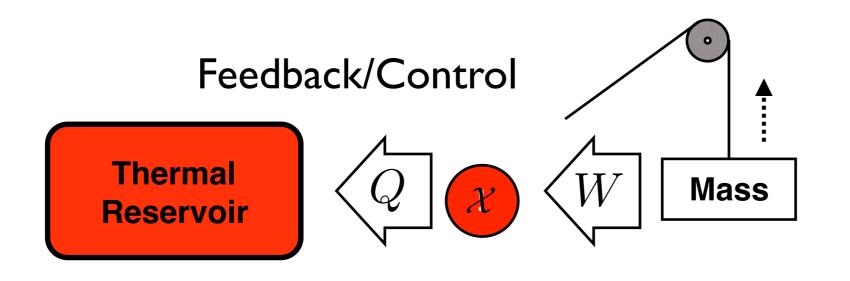
How do we extract work from a thermodynamic system?

How do we extract work from a thermodynamic system?



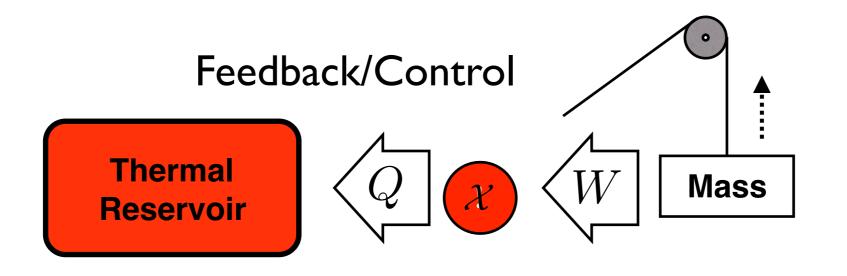
How do we extract work from a thermodynamic system?

Through control of the energy landscape $d\langle E \rangle = \sum_{x} \Pr(X_t = x) dE(x) + \sum_{x} E(x) d\Pr(X_t = x)$



How do we extract work from a thermodynamic system?

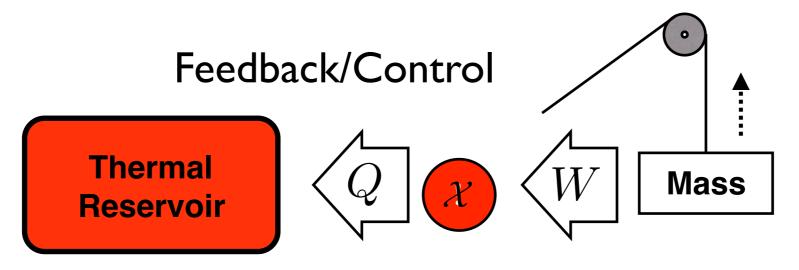
Through control of the energy landscape $d\langle E \rangle = \sum_{x} \Pr(X_t = x) dE(x) + \sum_{x} E(x) d\Pr(X_t = x)$ $= \langle dW \rangle + \langle dQ \rangle$



How do we extract work from a thermodynamic system?

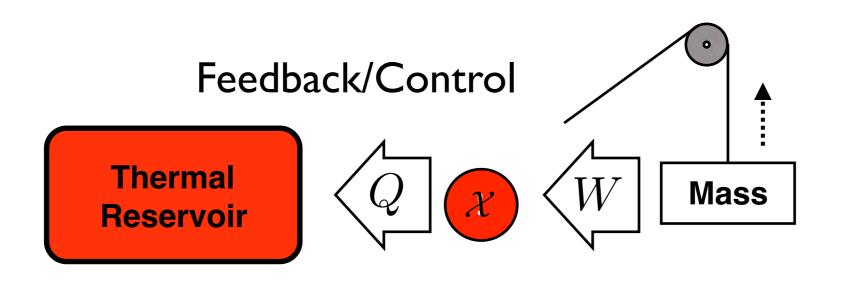
Through control of the energy landscape $d\langle E \rangle = \sum_{x} \Pr(X_t = x) dE(x) + \sum_{x} E(x) d\Pr(X_t = x)$ $= \langle dW \rangle + \langle dQ \rangle$

We consider control of a system X which interacts with an ideal thermal reservoir \mathcal{R} at temperature T to produce work



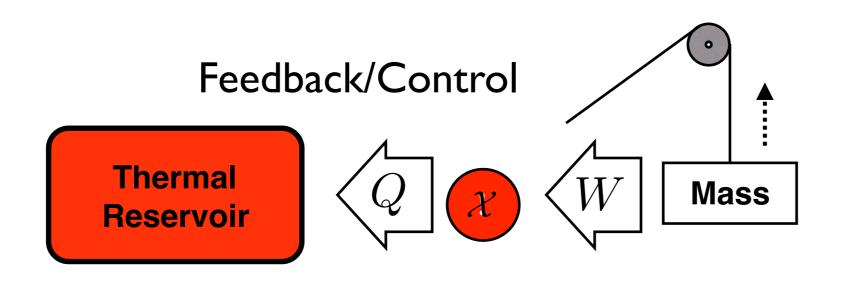
The total entropy production of the joint reservoir system

 $\Delta S_{\text{total}} = \Delta S[R] + \Delta S[X] - \Delta I[X;R]$



The total entropy production of the joint reservoir system

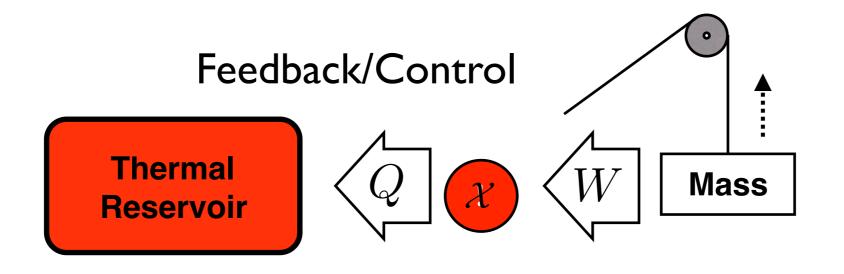
 $\Delta S_{\text{total}} = \Delta S[R] + \Delta S[X] - \Delta I[X;R]$ The fact that the reservoir is ideal means it has no memory of the ratchet, and so is uncorrelated with it: I[X;R] = 0



The total entropy production of the joint reservoir system

 $\Delta S_{\text{total}} = \Delta S[R] + \Delta S[X] - \Delta I[X;R]$ The fact that the reservoir is ideal means it has no memory of the ratchet, and so is uncorrelated with it: I[X;R] = 0

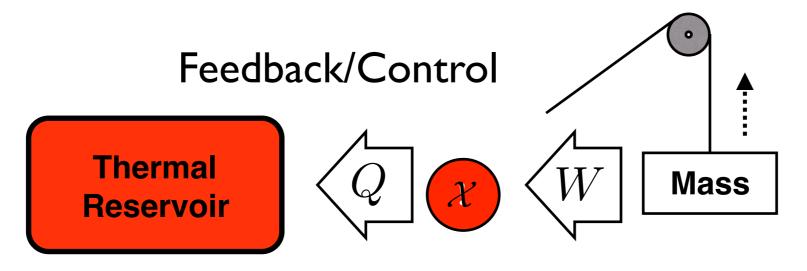
Heat production in control is bounded by change: $\frac{\langle Q \rangle}{T} + \Delta S[X] = \Delta S_{\text{total}} \ge 0$



The total entropy production of the joint reservoir system

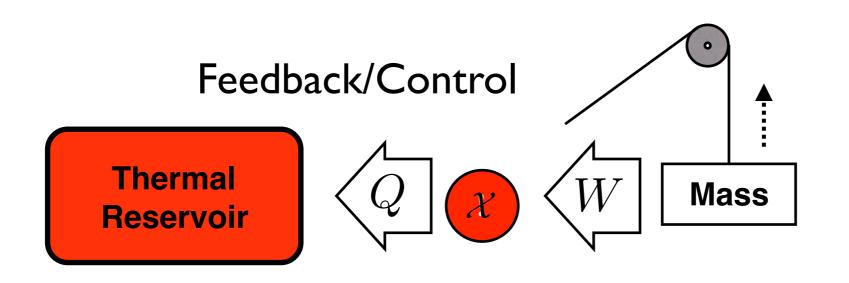
 $\Delta S_{\text{total}} = \Delta S[R] + \Delta S[X] - \Delta I[X;R]$ The fact that the reservoir is ideal means it has no memory of the ratchet, and so is uncorrelated with it: I[X;R] = 0

 $\begin{array}{l} \mbox{Heat production in control is bounded by} \\ \mbox{change:} \quad \frac{\langle Q \rangle}{T} + \Delta S[X] = \Delta S_{\rm total} \geq 0 \\ \\ \quad \langle Q \rangle \geq -T \Delta S[X] \end{array}$

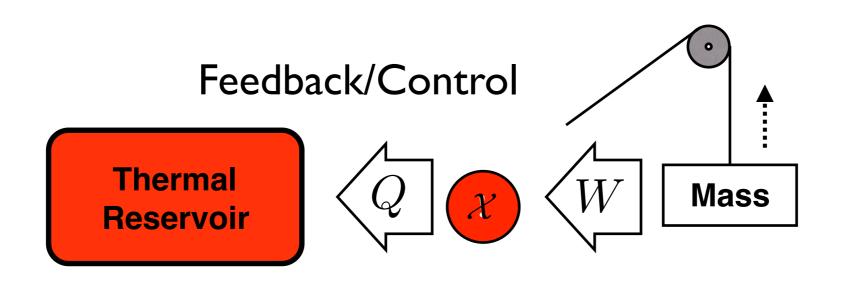


This can be re-expressed with work

 $T\Delta S_{\text{total}} = \langle W \rangle - \Delta \langle E[X] \rangle + T\Delta S[X]$

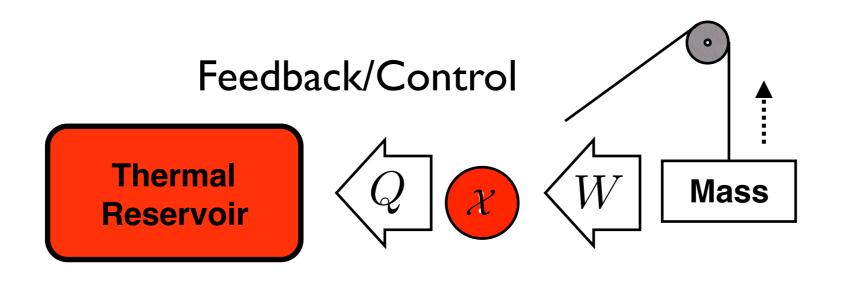


This can be re-expressed with work $T\Delta S_{\text{total}} = \langle W \rangle - \Delta \langle E[X] \rangle + T\Delta S[X]$ This can be interpreted as dissipated work $\langle W_{\text{diss}} \rangle = \langle W \rangle - \Delta F^{\text{NEQ}}$



This can be re-expressed with work $T\Delta S_{\text{total}} = \langle W \rangle - \Delta \langle E[X] \rangle + T\Delta S[X]$ This can be interpreted as dissipated work $\langle W_{\text{diss}} \rangle = \langle W \rangle - \Delta F^{\text{NEQ}}$

This includes a new quantity: the nonequilibrium free energy, which specifies the amount of work that could have been harvested: $F^{NEQ} = \langle E[X] \rangle - TS[X]$



What does entropy do?

What does entropy do?

I) Bounds function of heat engine

$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$
$$e \le 1 - \frac{T_c}{T_h}$$

What does entropy do?

I) Bounds function of heat engine with Carnot efficiency

$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$
$$e \le 1 - \frac{T_c}{T_h}$$

What does entropy do?

I) Bounds function of heat engine with Carnot efficiency

$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$
$$e \le 1 - \frac{T_c}{T_h}$$

2) Bounds work and heat production in controlled thermodynamic systems

What does entropy do?

I) Bounds function of heat engine with Carnot efficiency

$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$
$$e \le 1 - \frac{T_c}{T_h}$$

2) Bounds work and heat production in controlled thermodynamic systems

 $\langle Q \rangle \ge -k_B T \ln 2 \,\Delta H[X]$

What does entropy do?

I) Bounds function of heat engine with Carnot efficiency

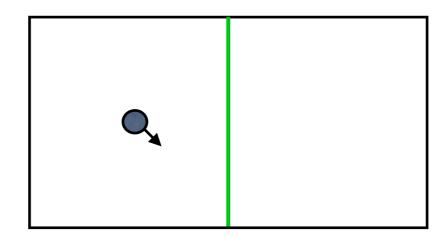
$$e = \frac{\text{output}}{\text{input}} = \frac{W}{Q_h} = 1 - \frac{Q_c}{Q_h}$$
$$e \le 1 - \frac{T_c}{T_h}$$

2) Bounds work and heat production in controlled thermodynamic systems

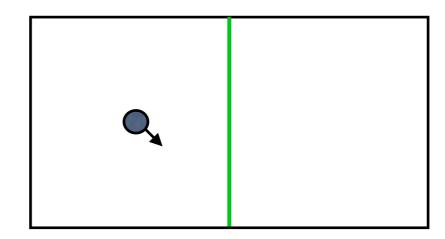
$$\langle Q \rangle \ge -k_B T \ln 2 \Delta H[X]$$

 $\langle W \rangle \ge \langle \Delta E[X] \rangle - k_B T \ln 2 \Delta H[X]$

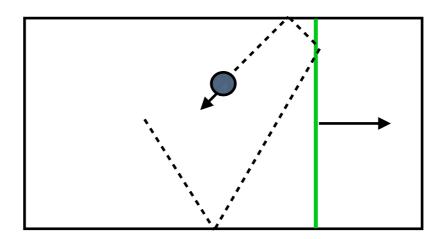
Single particle in a box



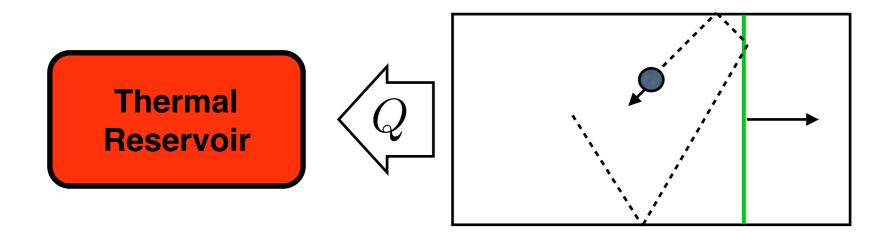
Single particle in a box, insert barrier



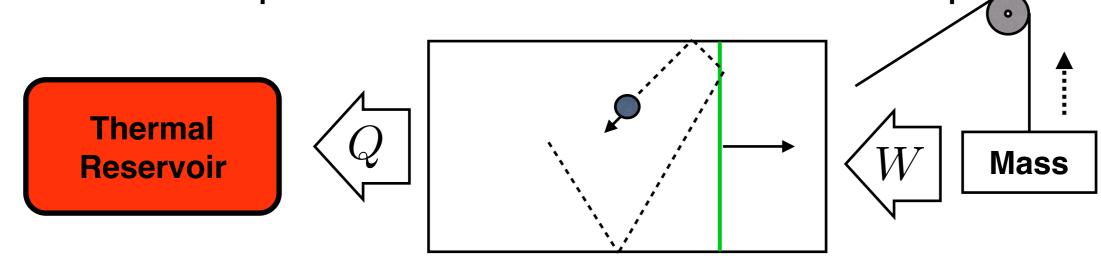
Single particle in a box, insert barrier, slide barrier away from particle



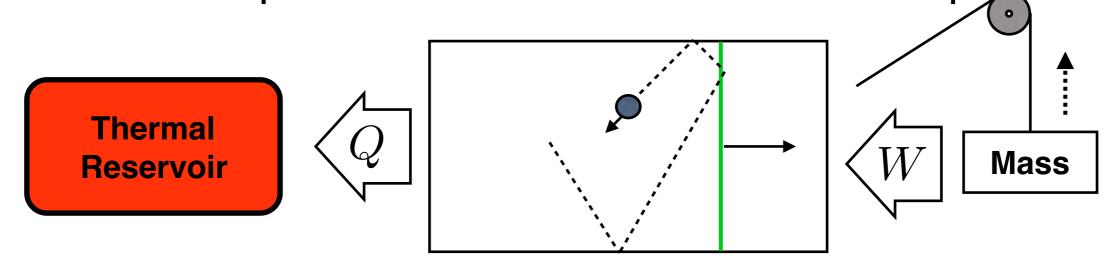
Single particle in a box, insert barrier, slide barrier away from particle, while in contact with heat reservoir at temperature T



Single particle in a box, insert barrier, slide barrier away from particle, while in contact with heat reservoir at temperature T, and extract work with the piston.



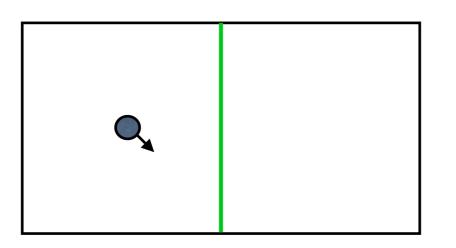
Single particle in a box, insert barrier, slide barrier away from particle, while in contact with heat reservoir at temperature T, and extract work with the piston.

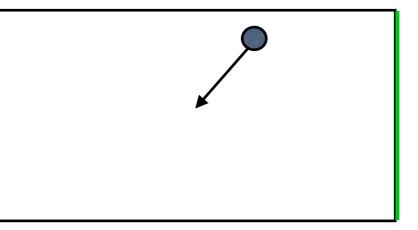


How much much work and heat flow?

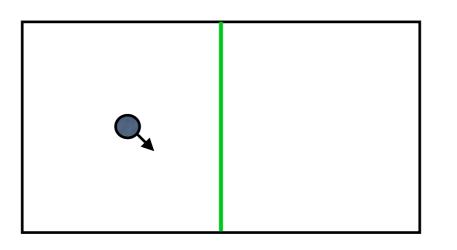
 $\langle Q \rangle \ge -k_B T \ln 2 \Delta H[X]$ $\langle W \rangle \ge \langle \Delta E[X] \rangle - k_B T \ln 2 \Delta H[X]$

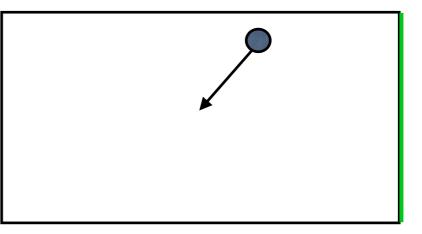
Initial vs. final energies and entropies:





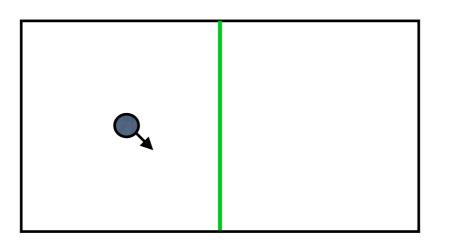
Initial vs. final energies and entropies:

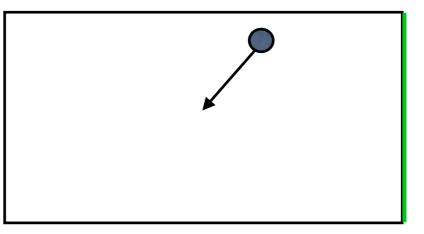




Energy of each position is the same, so average energy of positions is the same: $\langle E_{\vec{q}} \rangle_{\text{initial}} = \langle E_{\vec{q}} \rangle_{\text{final}}$

Initial vs. final energies and entropies:

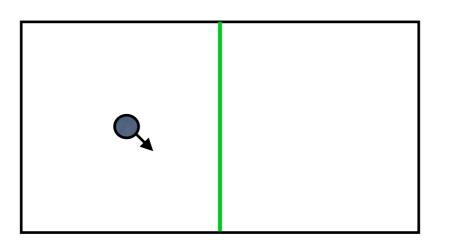


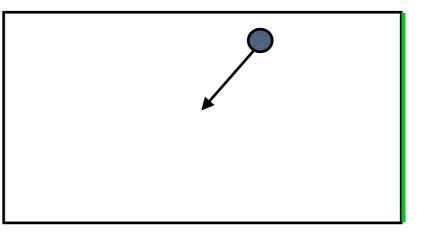


Energy of each position is the same, so average energy of positions is the same: $\langle E_{\vec{q}} \rangle_{\text{initial}} = \langle E_{\vec{q}} \rangle_{\text{final}}$ If in equilibrium, average kinetic energy of momentum variable is the same: k_BT

$$\langle E_{\vec{p}} \rangle_{\text{initial}} = \langle E_{\vec{p}} \rangle_{\text{final}} = \frac{\kappa_B I}{2} = \langle \text{Kinetic Energy} \rangle$$

Initial vs. final energies and entropies:

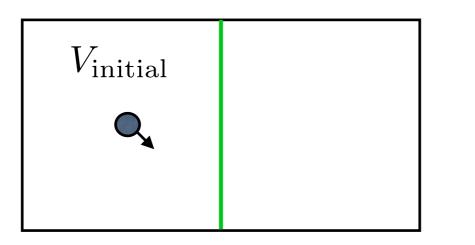


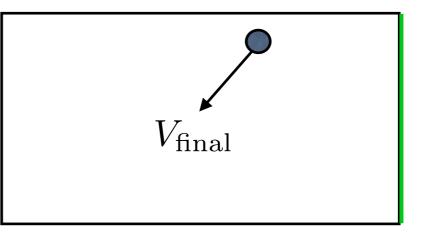


Energy of each position is the same, so average energy of positions is the same: $\langle E_{\vec{q}} \rangle_{\text{initial}} = \langle E_{\vec{q}} \rangle_{\text{final}}$ If in equilibrium, average kinetic energy of momentum variable is the same: $\langle E_{\vec{p}} \rangle_{\text{initial}} = \langle E_{\vec{p}} \rangle_{\text{final}} = \frac{k_B T}{2} = \langle \text{Kinetic Energy} \rangle$

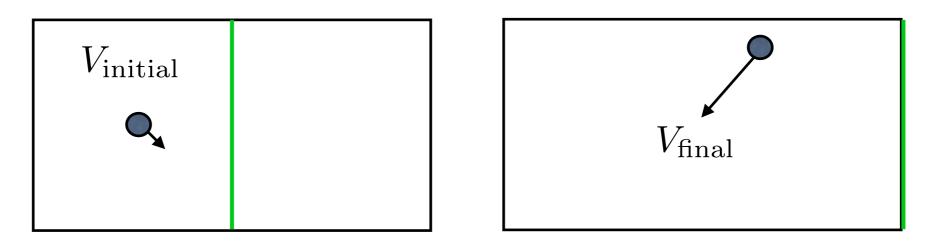
Average energy doesn't change: $\langle \Delta E[X] \rangle = 0$

Initial vs. final energies and entropies:





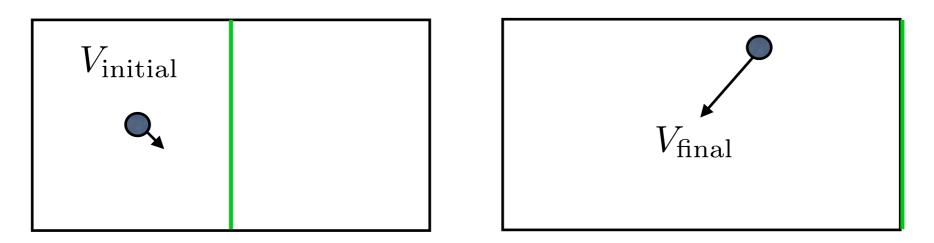
Initial vs. final energies and entropies:



Probability of each position is uniform and inversely proportional to the available volume:

$$\Pr(\vec{Q}_{\text{initial}} = \vec{q}) = \frac{c}{V_{\text{initial}}} \qquad \Pr(\vec{Q}_{\text{final}} = \vec{q}) = \frac{c}{V_{\text{final}}}$$

Initial vs. final energies and entropies:



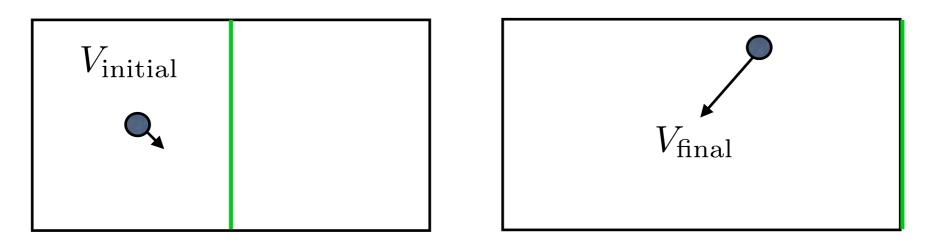
Probability of each position is uniform and inversely proportional to the available volume:

$$\Pr(\vec{Q}_{\text{initial}} = \vec{q}) = \frac{c}{V_{\text{initial}}} \qquad \Pr(\vec{Q}_{\text{final}} = \vec{q}) = \frac{c}{V_{\text{final}}}$$

At equilibrium, the probability of momentum is the same initially and finally

$$\Pr(\vec{P}_{\text{initial}} = \vec{p}) = \Pr(\vec{P}_{\text{final}} = \vec{p}) = \Pr(\vec{P}^{\text{eq}} = \vec{p})$$

Initial vs. final energies and entropies:



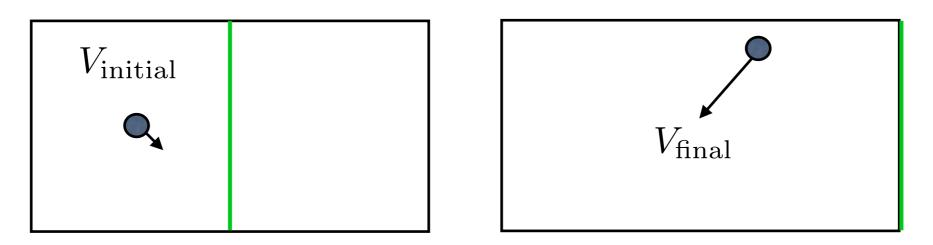
Probability of each position is uniform and inversely proportional to the available volume:

$$\Pr(\vec{Q}_{\text{initial}} = \vec{q}) = \frac{c}{V_{\text{initial}}} \qquad \Pr(\vec{Q}_{\text{final}} = \vec{q}) = \frac{c}{V_{\text{final}}}$$

At equilibrium, the probability of momentum is the same initially and finally, and independent of position distribution. $\Pr(\vec{P}_{\text{initial}} = \vec{p}) = \Pr(\vec{P}_{\text{final}} = \vec{p}) = \Pr(\vec{P}^{\text{eq}} = \vec{p})$

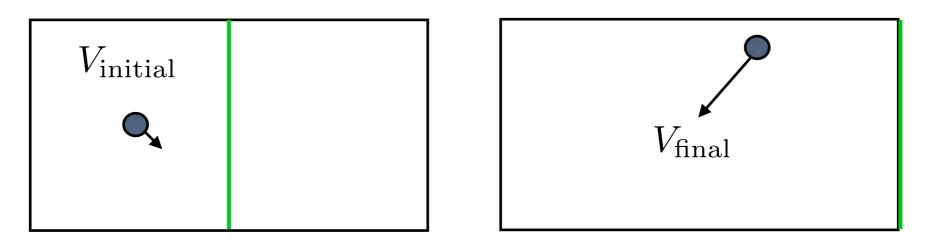
$$\Pr(X_{\text{initial}} = (\vec{q}, \vec{p})) = \frac{c \Pr(\vec{P}_{\text{initial}} = \vec{p})}{V_{\text{initial}}} \quad \Pr(X_{\text{final}} = (\vec{q}, \vec{p})) = \frac{c \Pr(\vec{P}_{\text{final}} = \vec{p})}{V_{\text{final}}}$$

Initial vs. final energies and entropies:



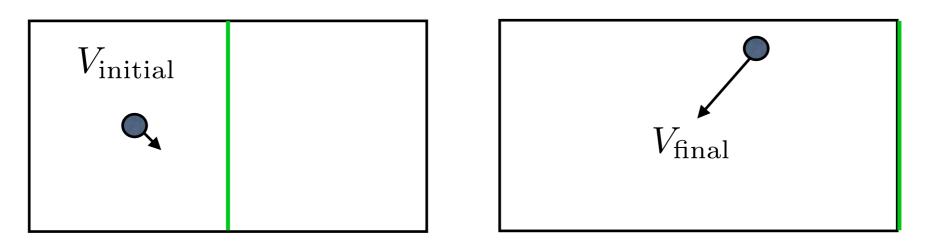
Thus, the change in Shannon entropy is $\Delta H[X] = H[X_{\text{final}}] - H[X_{\text{initial}}]$

Initial vs. final energies and entropies:



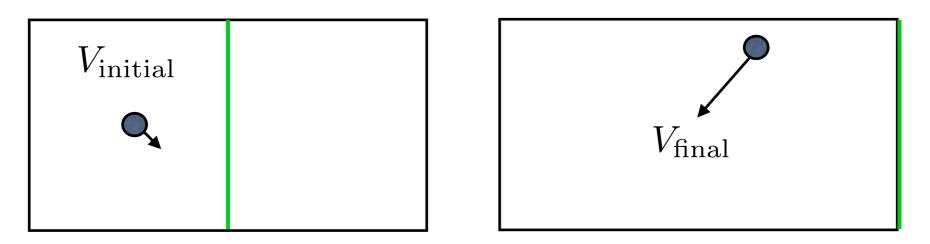
Thus, the change in Shannon entropy is $\Delta H[X] = H[X_{\text{final}}] - H[X_{\text{initial}}]$ $= H[\vec{P}^{\text{eq}}] - \sum_{x \in V_{\text{final}}} \frac{c}{V_{\text{final}}} \log_2 \frac{c}{V_{\text{final}}} - H[\vec{P}^{\text{eq}}] + \sum_{x \in V_{\text{initial}}} \frac{c}{V_{\text{initial}}} \log_2 \frac{c}{V_{\text{initial}}}$

Initial vs. final energies and entropies:



Thus, the change in Shannon entropy is $\Delta H[X] = H[X_{\text{final}}] - H[X_{\text{initial}}]$ $= H[\vec{P}^{\text{eq}}] - \sum_{x \in V_{\text{final}}} \frac{c}{V_{\text{final}}} \log_2 \frac{c}{V_{\text{final}}} - H[\vec{P}^{\text{eq}}] + \sum_{x \in V_{\text{initial}}} \frac{c}{V_{\text{initial}}} \log_2 \frac{c}{V_{\text{initial}}}$ $= \log_2 \frac{V_{\text{final}}}{V_{\text{initial}}}$

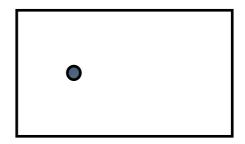
Initial vs. final energies and entropies:

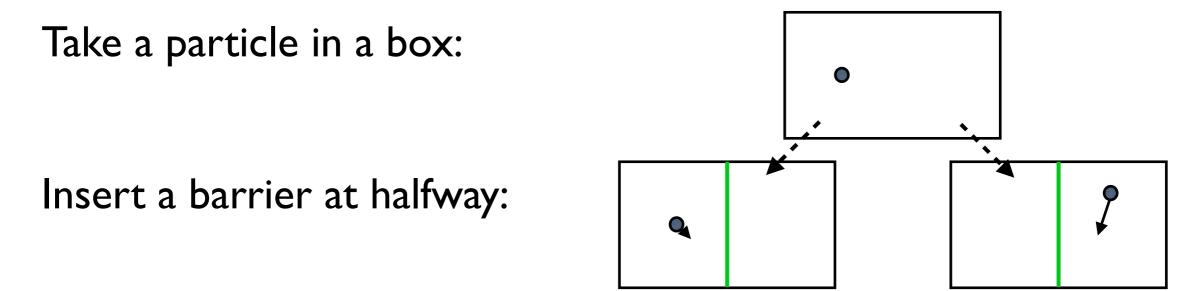


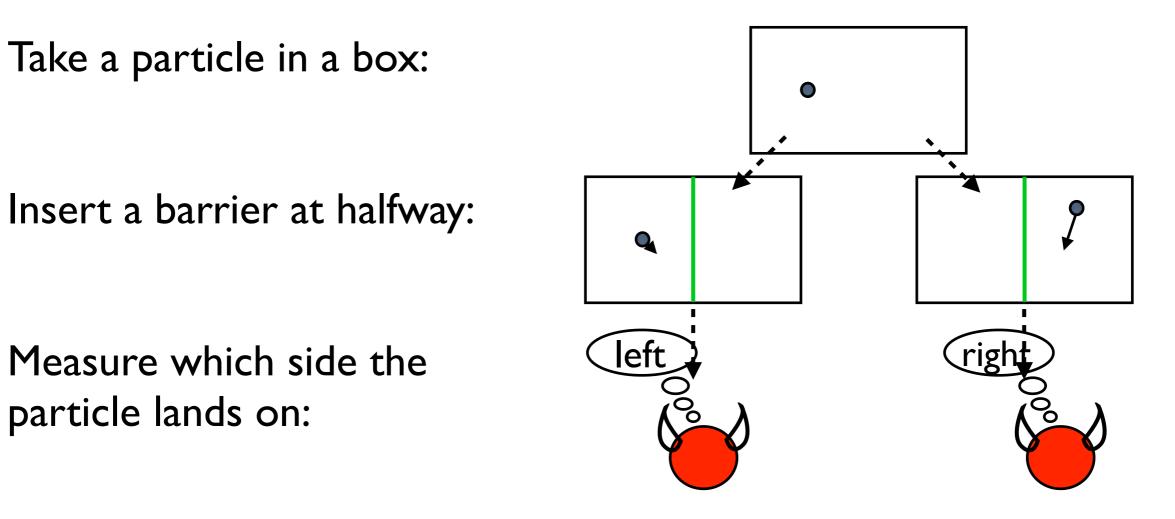
Thus, the change in Shannon entropy is $\Delta H[X] = H[X_{\text{final}}] - H[X_{\text{initial}}]$ $= H[\vec{P}^{\text{eq}}] - \sum_{x \in V_{\text{final}}} \frac{c}{V_{\text{final}}} \log_2 \frac{c}{V_{\text{final}}} - H[\vec{P}^{\text{eq}}] + \sum_{x \in V_{\text{initial}}} \frac{c}{V_{\text{initial}}} \log_2 \frac{c}{V_{\text{initial}}}$ $= \log_2 \frac{V_{\text{final}}}{V_{\text{initial}}}$ And the work and heat are bounded by the log ratio of

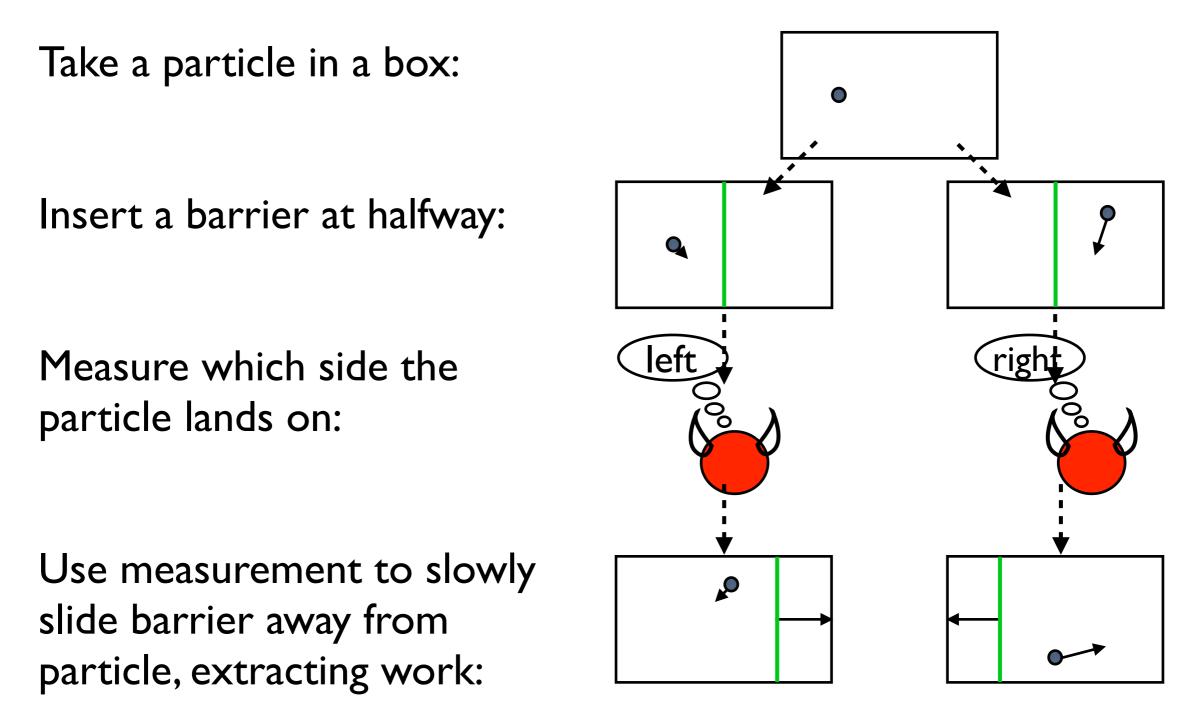
And the work and heat are bounded by the log ratio of the volumes (achievable if done very slowly): $\langle W \rangle \ge -k_B T \ln 2 \log_2 \frac{V_{\text{final}}}{V_{\text{initial}}} \qquad \langle Q \rangle \ge -k_B T \ln 2 \log_2 \frac{V_{\text{final}}}{V_{\text{initial}}}$

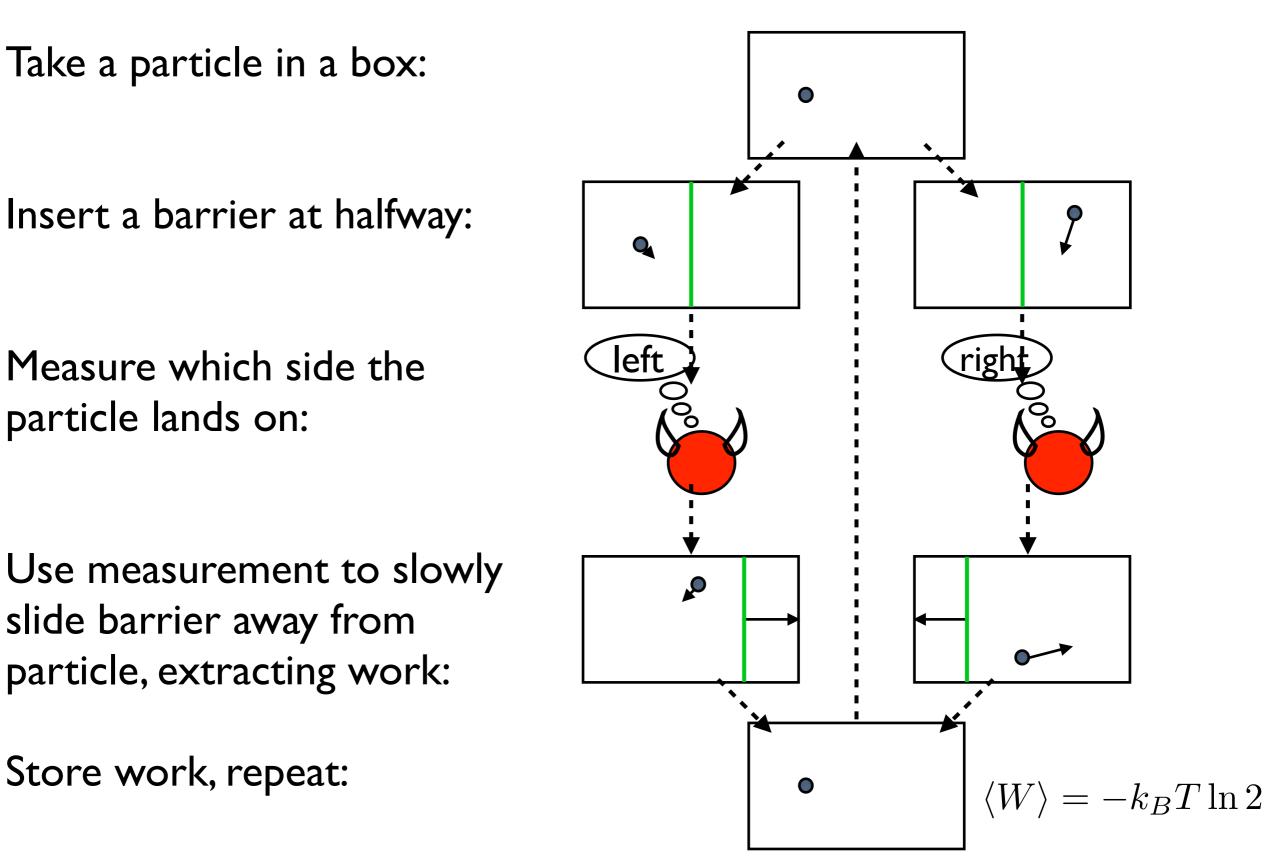
Take a particle in a box:





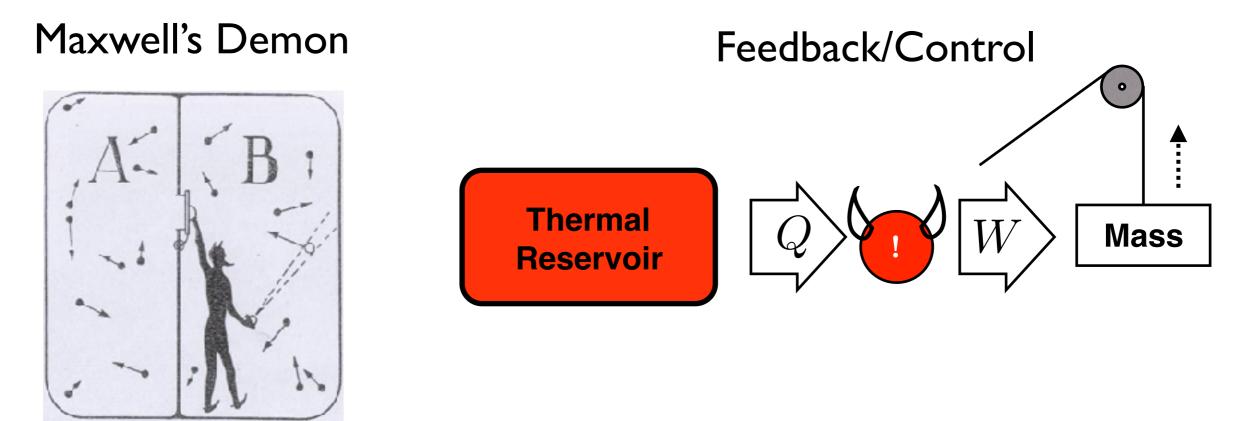


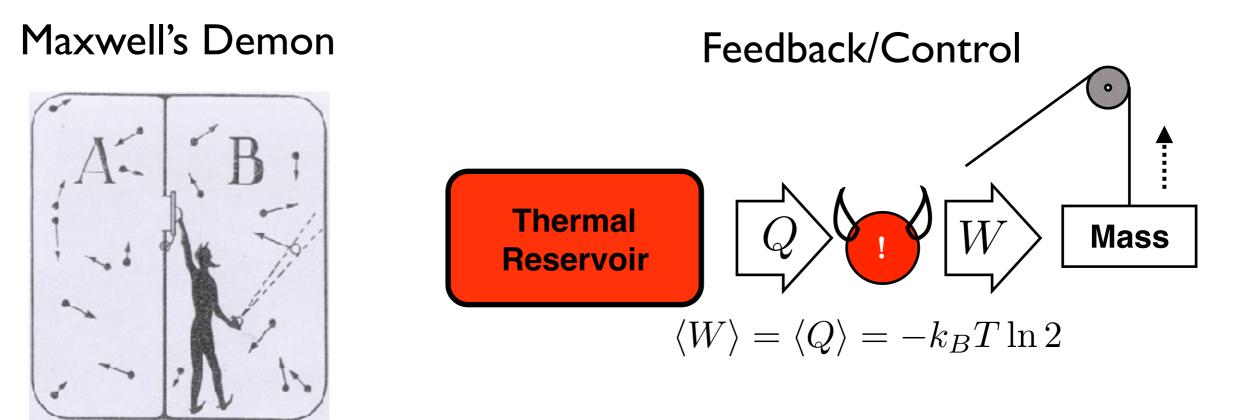


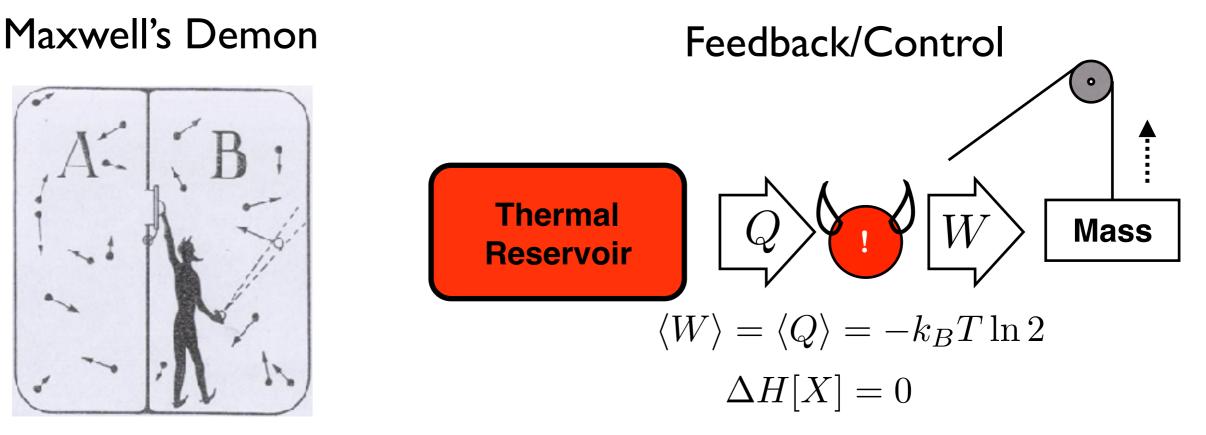


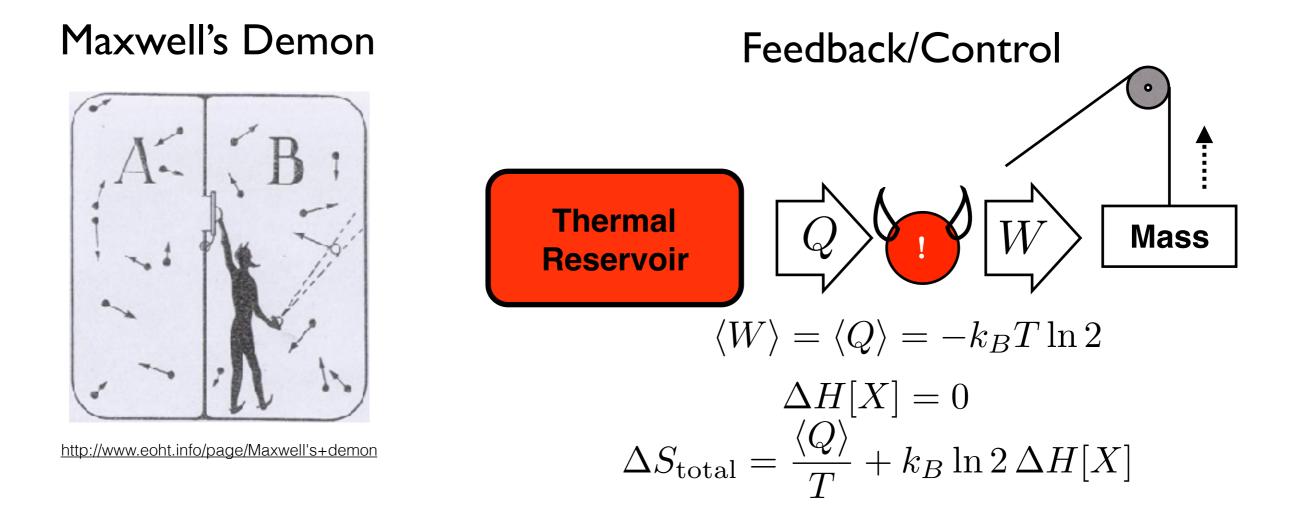
Maxwell's Demon

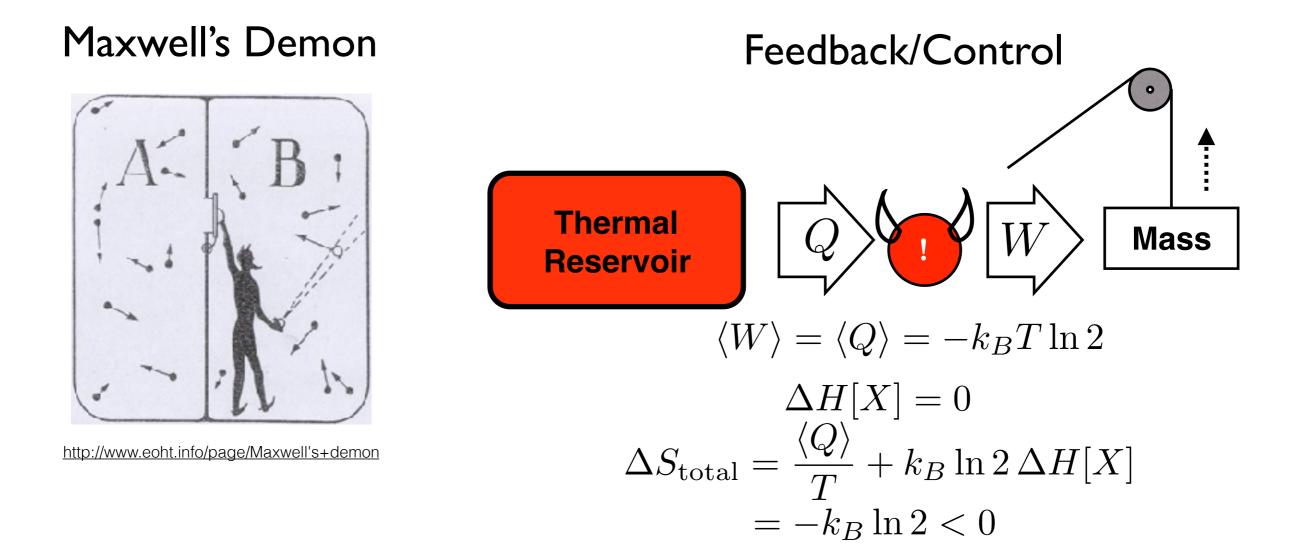


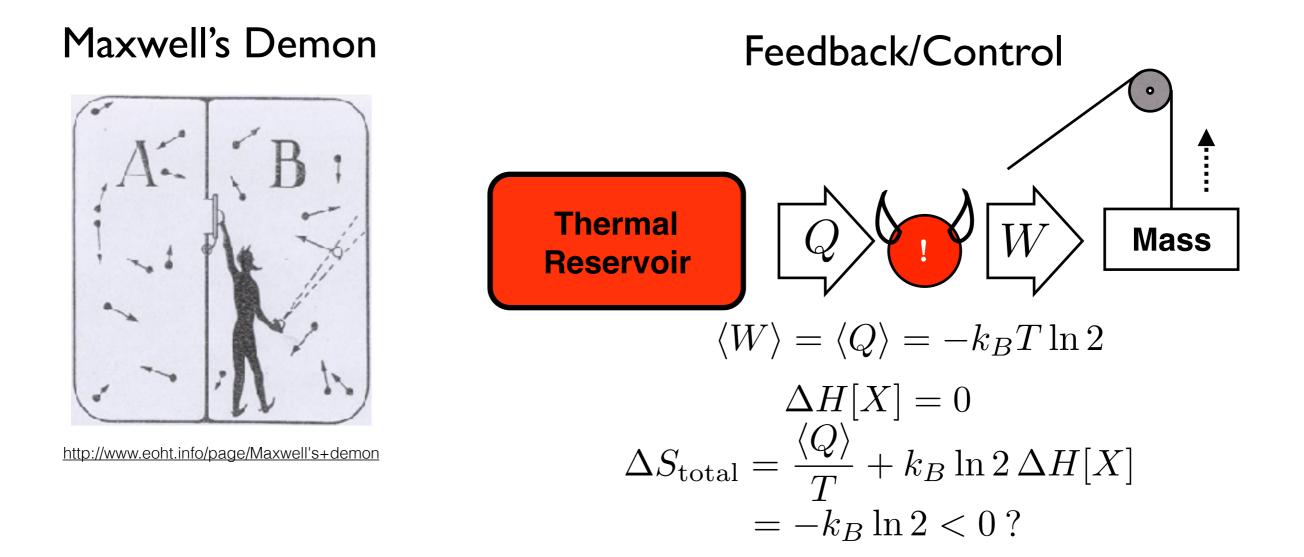


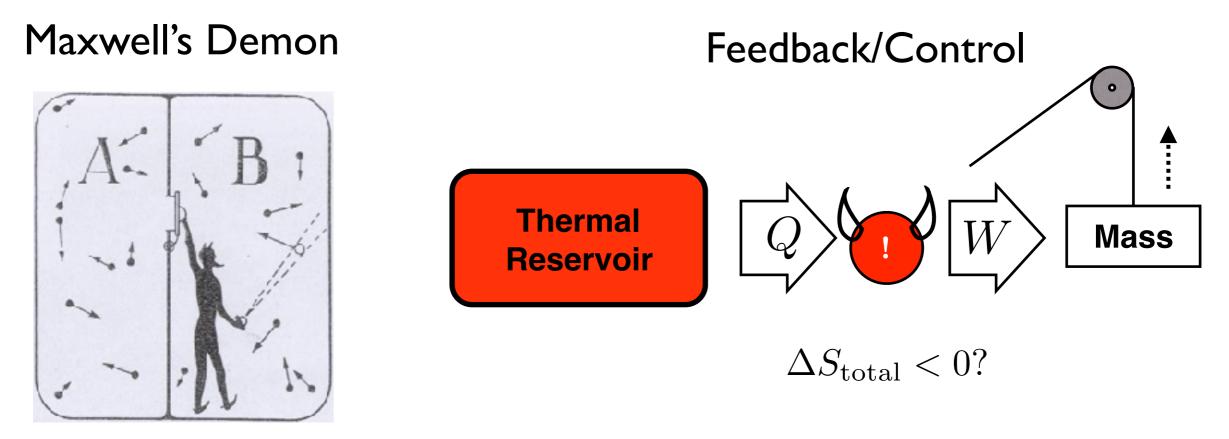




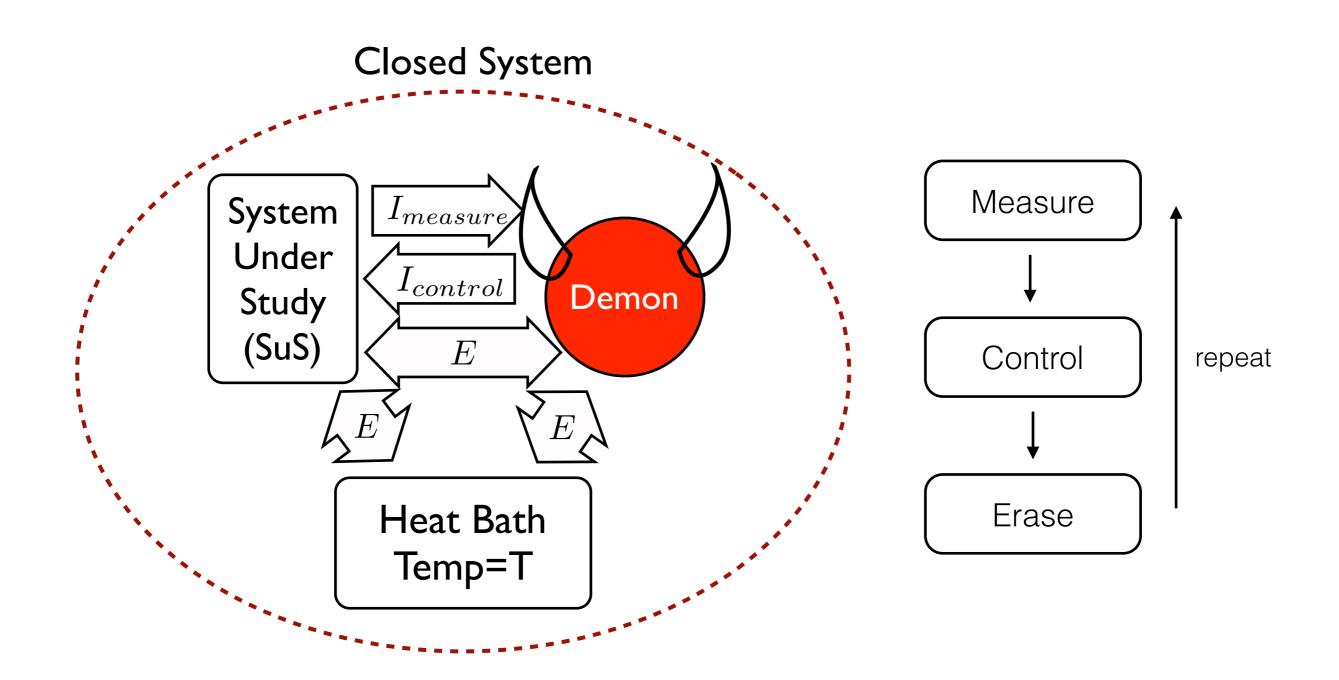


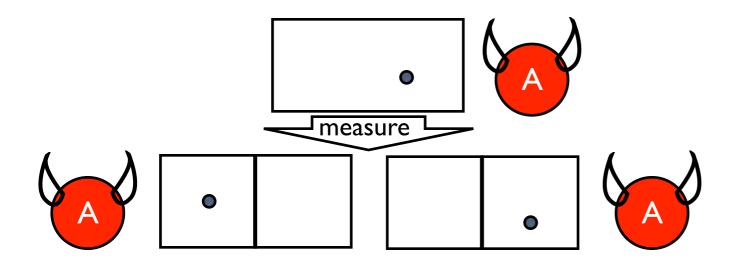


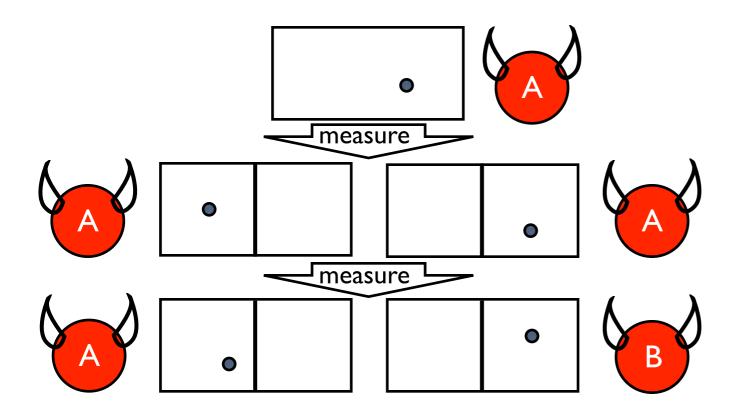


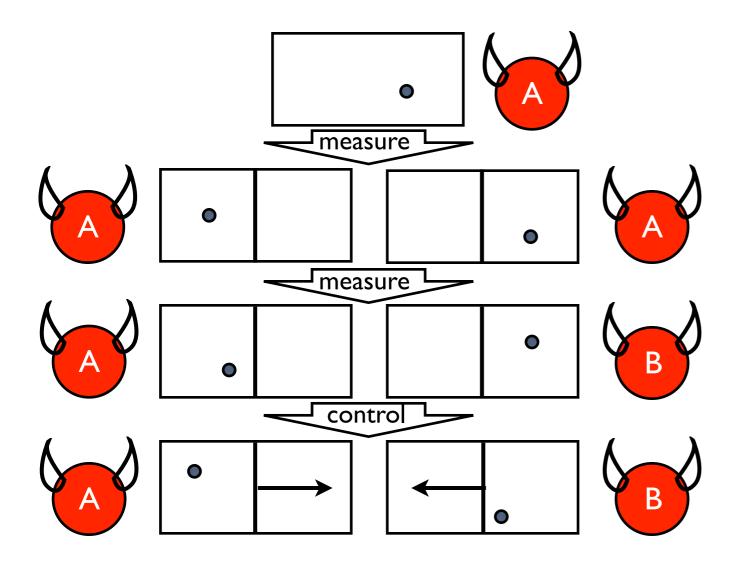


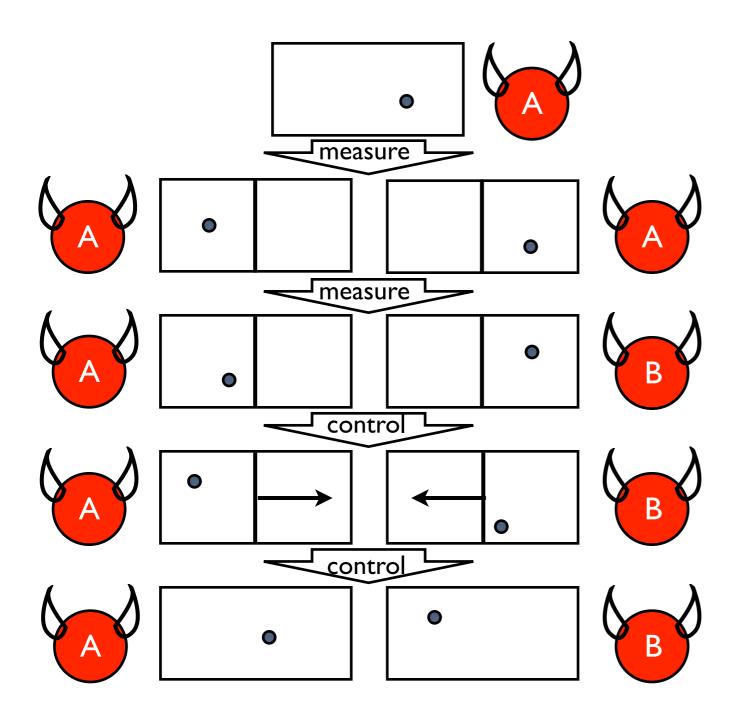
Maxwellian Demons



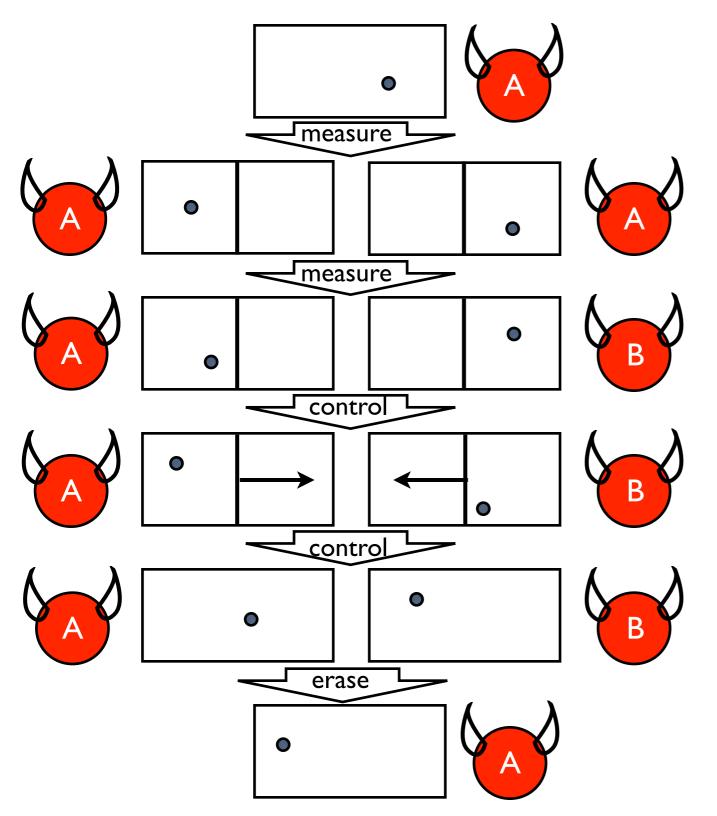




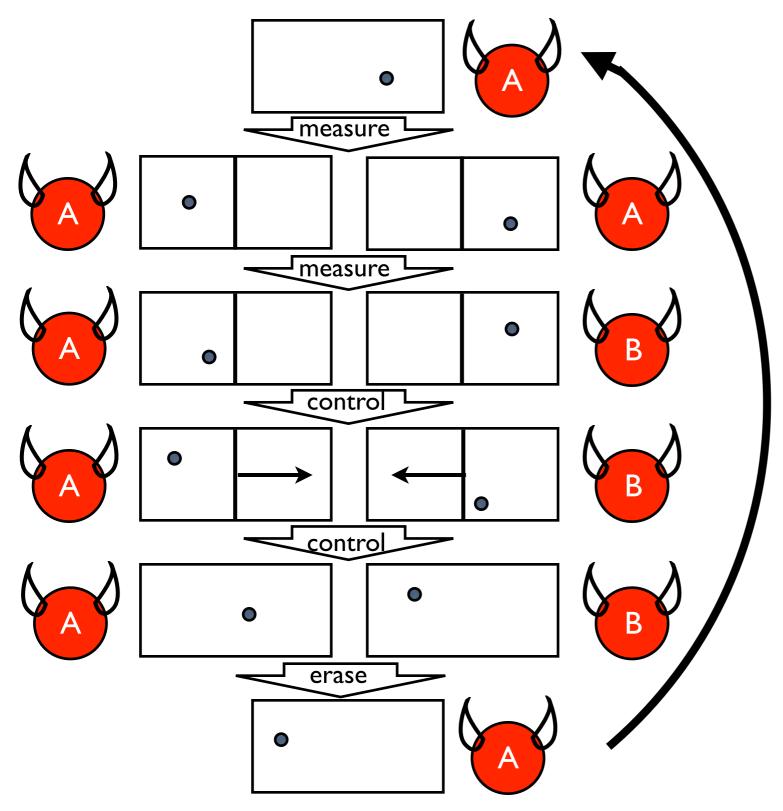




Demon is able to extract and store $k_B T \ln 2$ work every cycle.



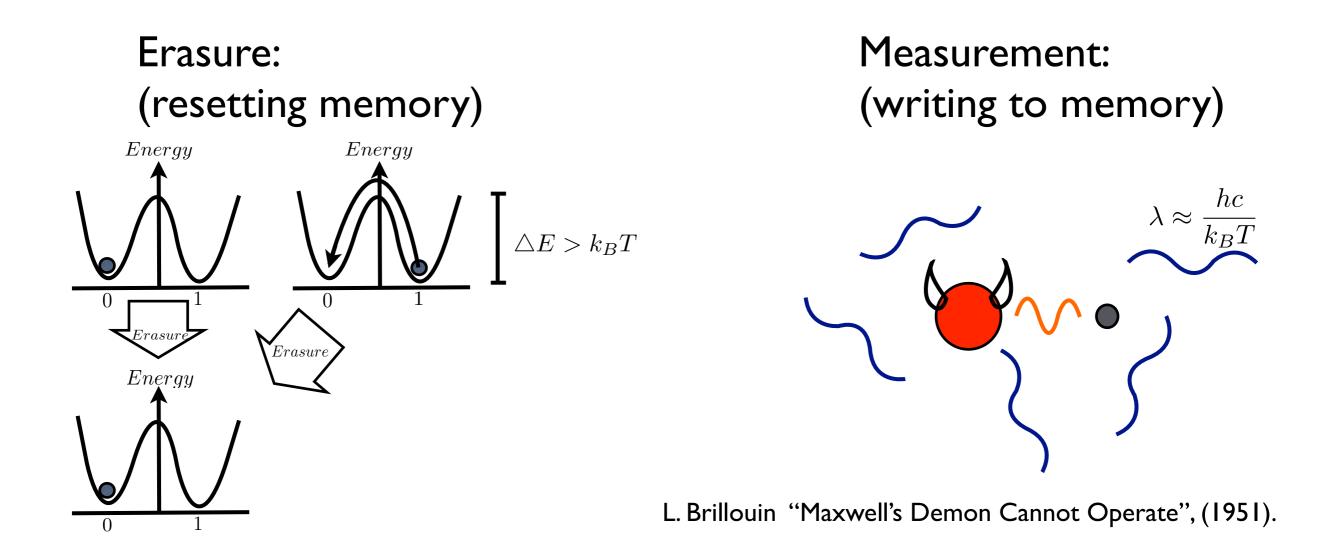
Demon is able to extract and store $k_B T \ln 2$ work every cycle.



Demon is able to extract and store $k_B T \ln 2$ work every cycle.

Violates the second law?

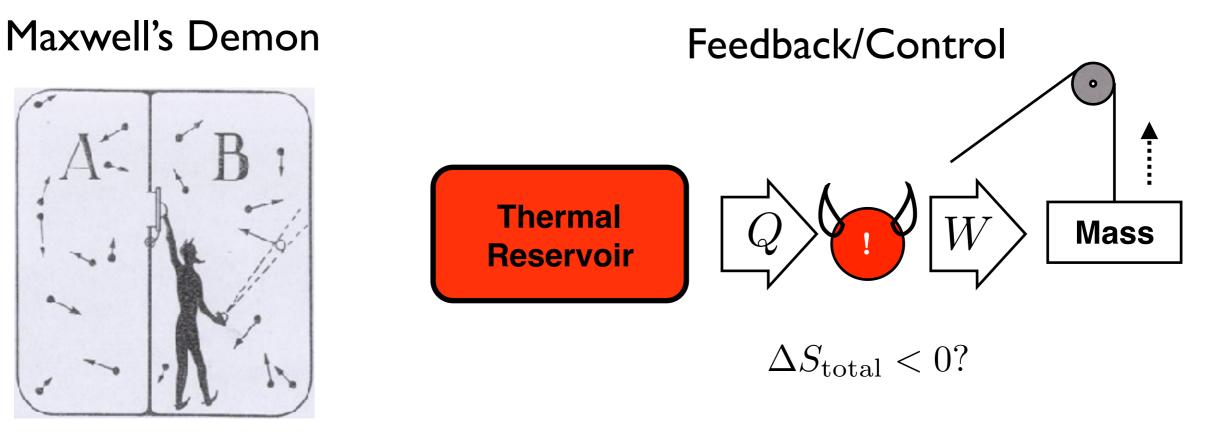
Energetic Costs



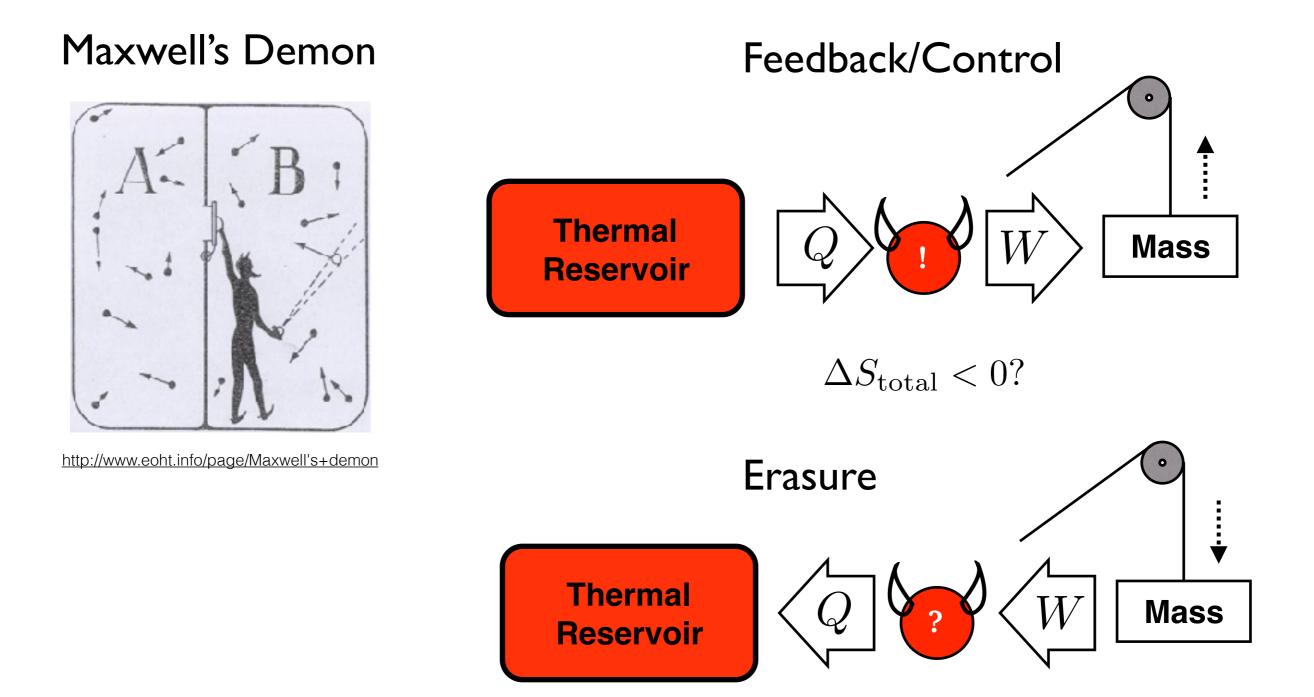
Landauer's principle: $Q_{\text{erase}} \ge k_B T \ln 2$

R. Landauer, "Irreversibility and heat generation in the computing process", (1961).

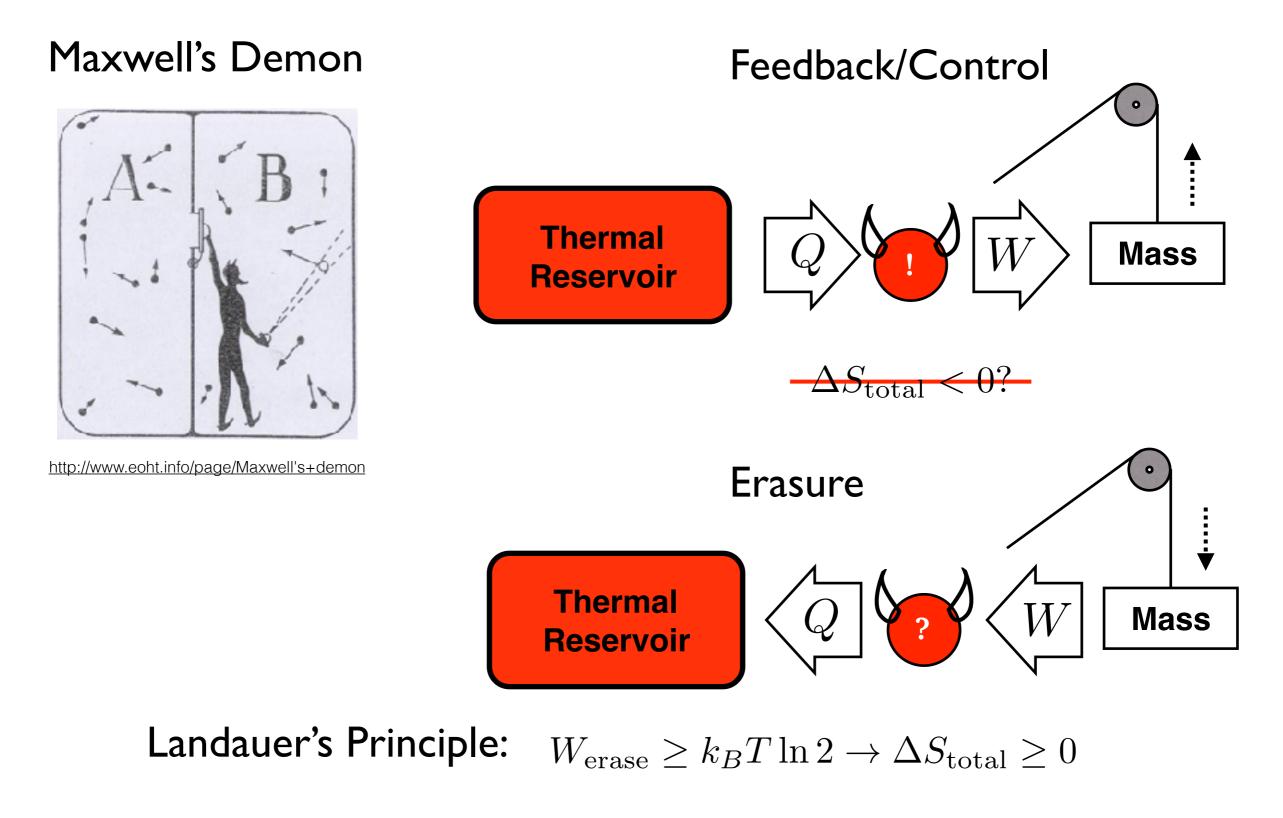
Information Bearing Degrees of Freedom



Information Bearing Degrees of Freedom

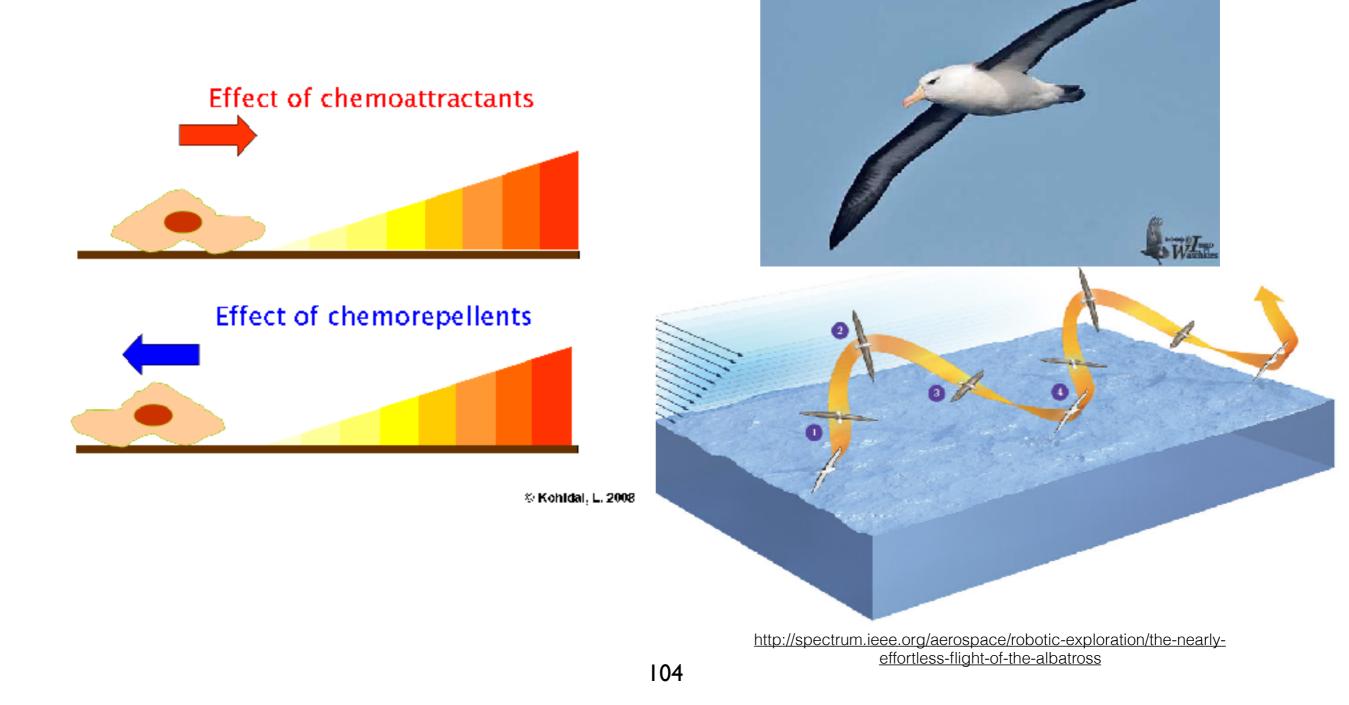


Information Bearing Degrees of Freedom



Benefit of Information Processing

Information processing allows systems to leverage structure of environment



Cost of Information Processing

Processing information costs work and dissipates heat.



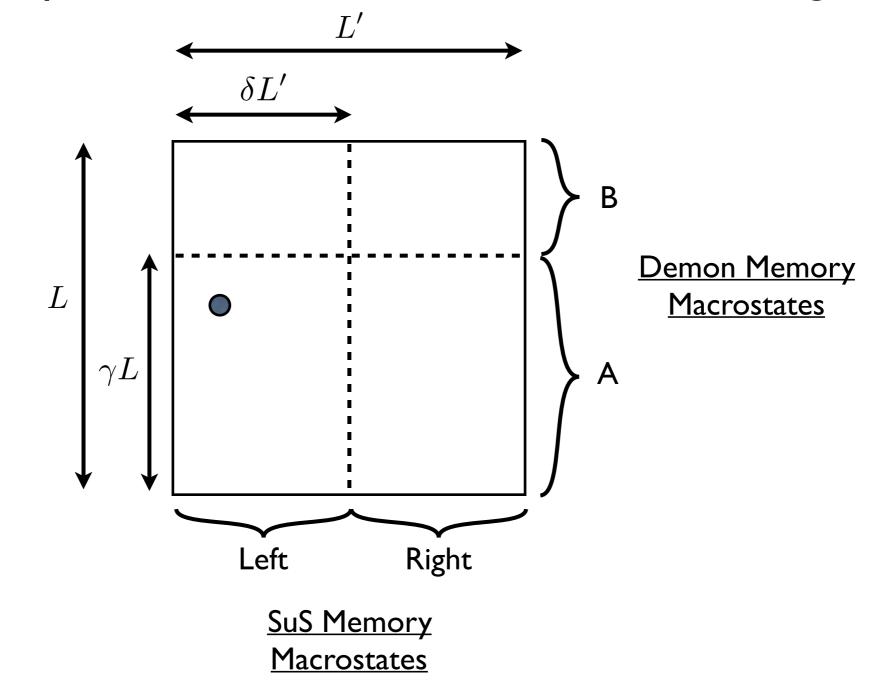
http://spectrum.ieee.org/computing/hardware/new-tech-keeps-datacenters-cool-in-warm-climates



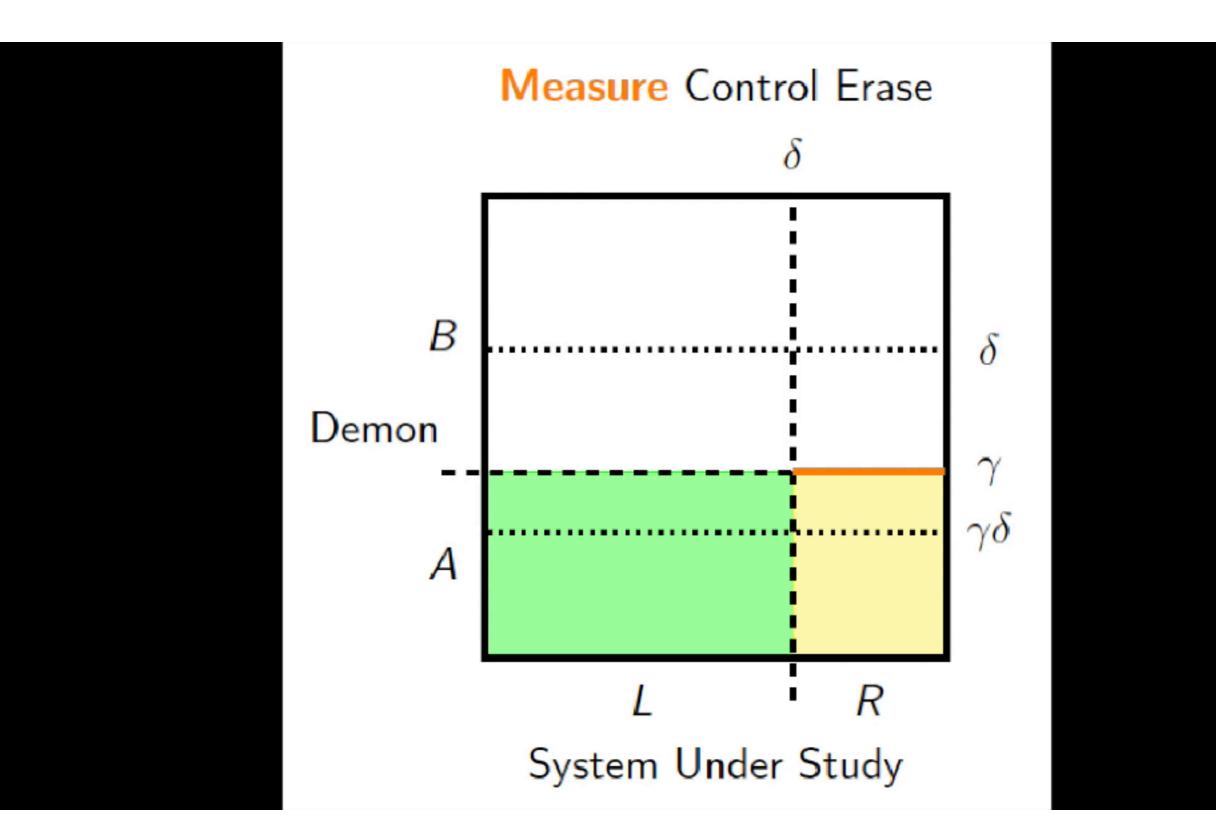
http://pooh.wikia.com/wiki/Think,_Think,_Think

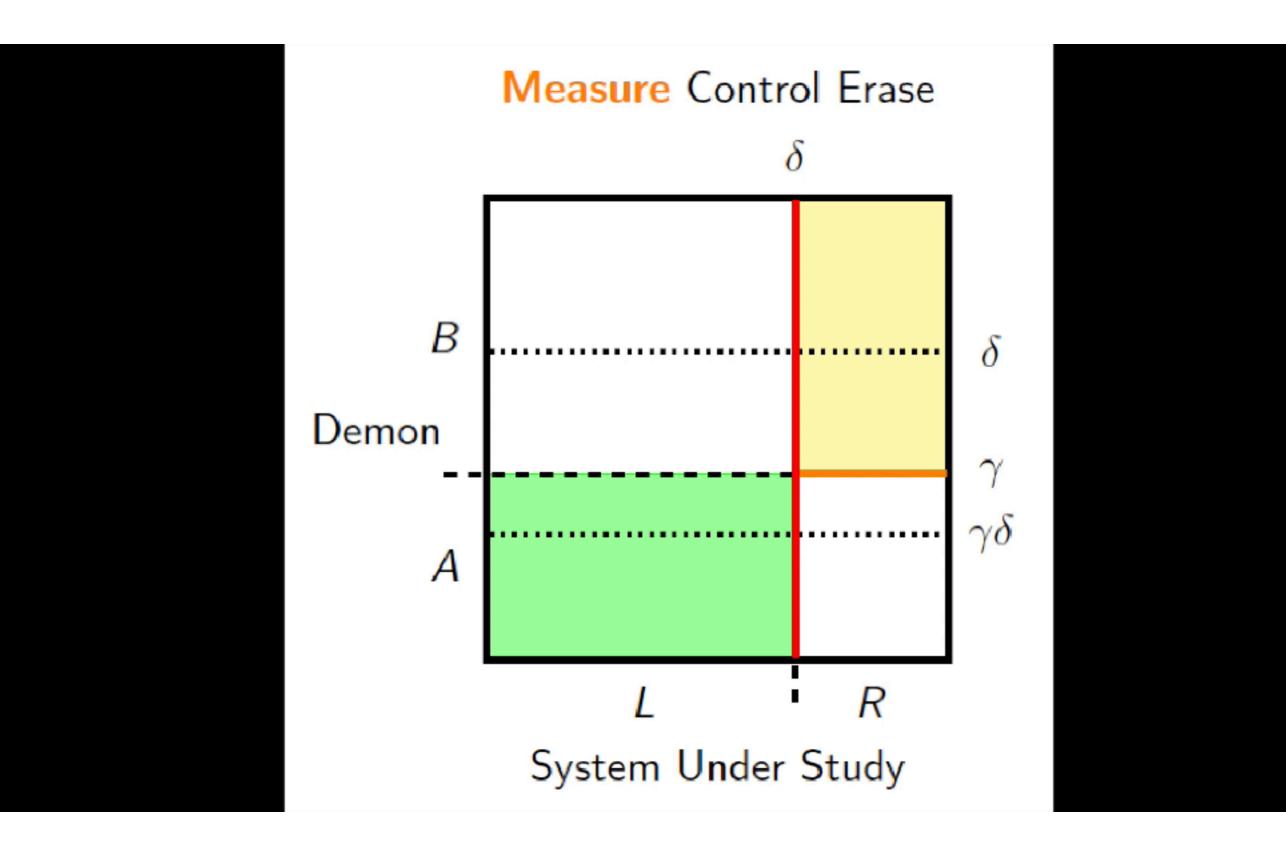
Szilard Example

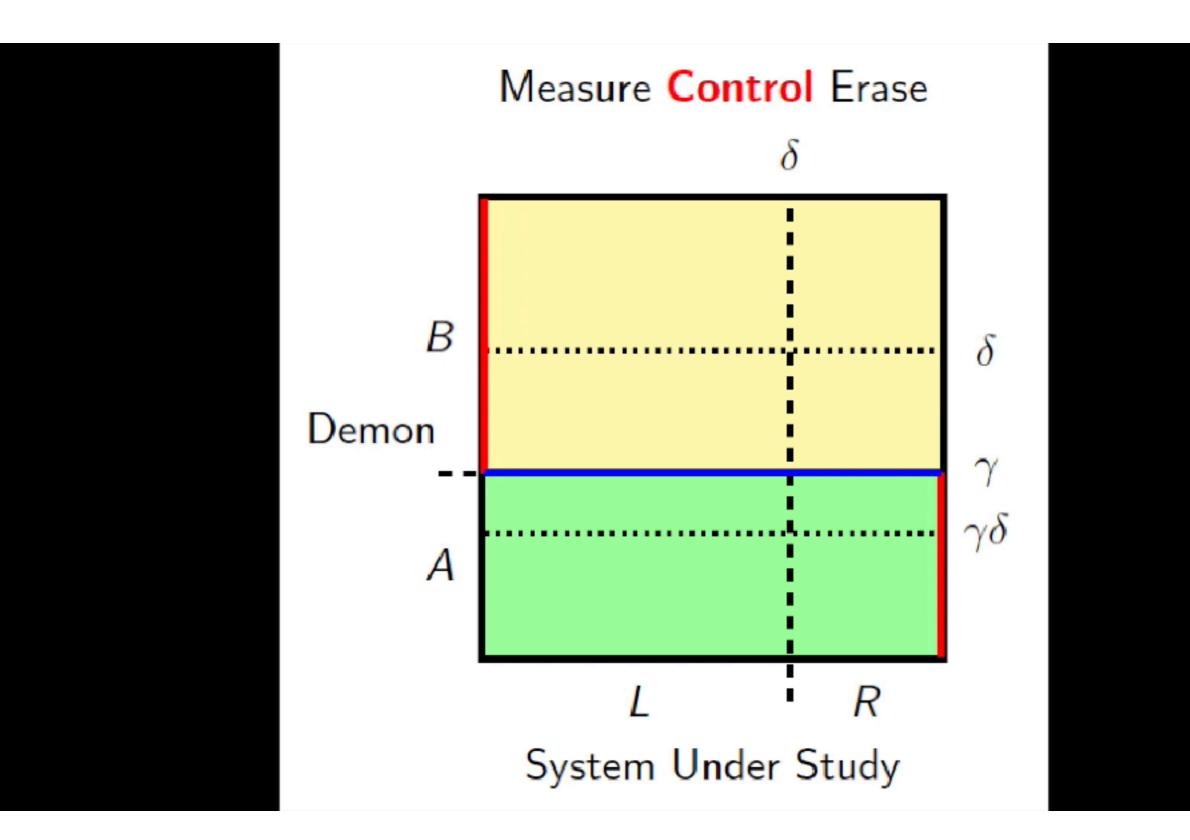
The Szilard system can be reformulated as a 2D box of gas.



C. Bennet, "Thermodynamics of Computation-A Review", (1981)







The average dissipated heat is exactly calculable as the work done to slide the barriers:

$$Q = W = -\int PdV$$

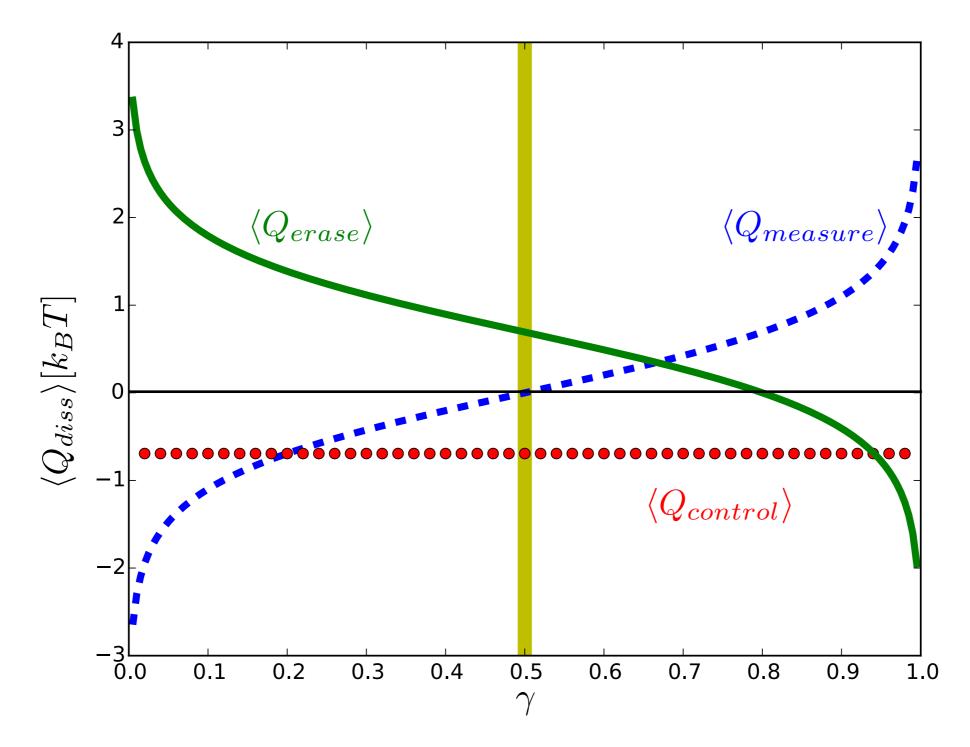
The average dissipated heat is exactly calculable as the work done to slide the barriers:

$$Q = W = -\int P dV$$
$$= -T\Delta S[X]$$

The average dissipated heat is exactly calculable as the work done to slide the barriers:

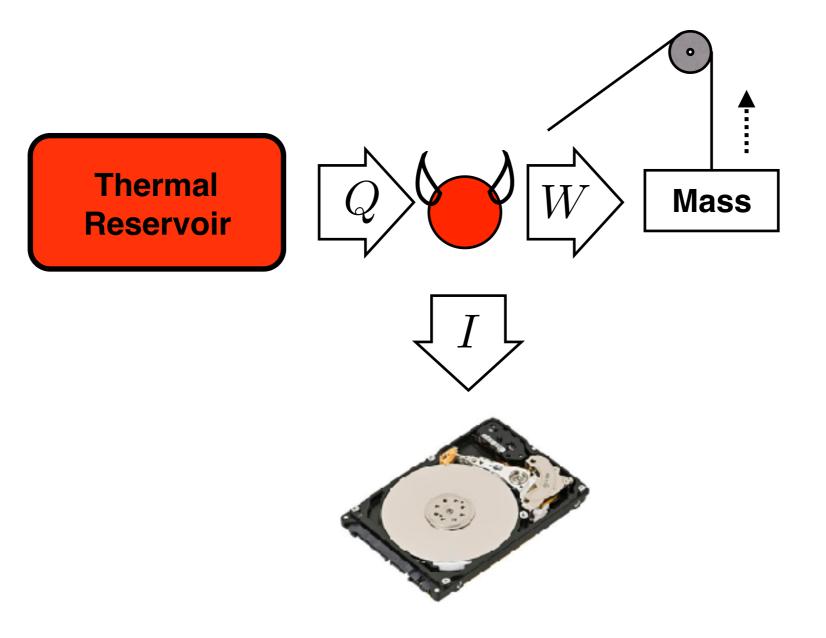
$$Q = W = -\int P dV$$
$$= -T\Delta S[X]$$

$$\Delta S_{\text{total}} = 0!$$



A. Boyd, J. Crutchfield, "Maxwell Demon Dynamics: Deterministic Chaos, the Szilard Map, and the Intelligence of Thermodynamic Systems", PRL, (2016)

Information is a Thermodynamic Fuel



Instead of erasing, write to a hard drive