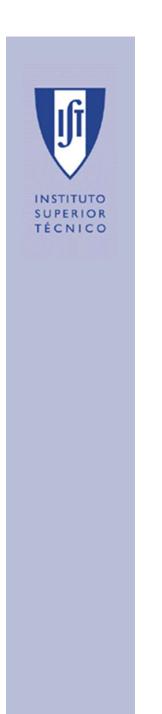


2020/2021

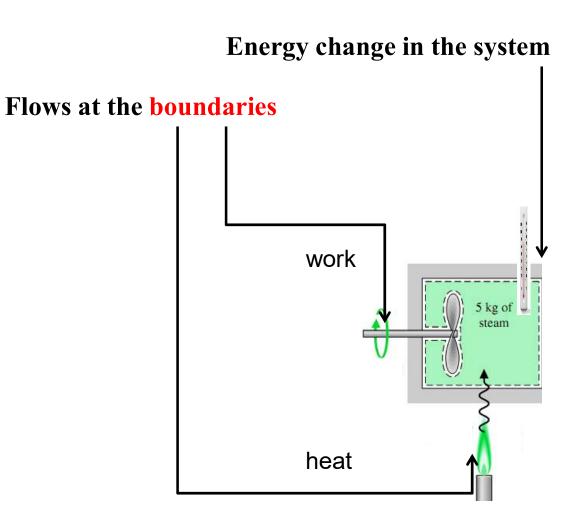
Thermodynamics

Tânia Sousataniasousa@tecnico.ulisboa.pt

Tiago Domingos tdomingos@tecnico.ulisboa.pt



Energy Balance in Closed Systems







Energy Balance in Closed Systems

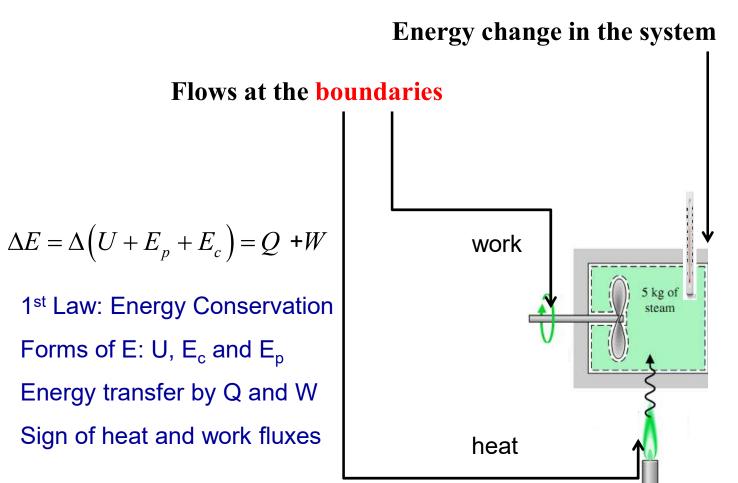
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Moran et al., 2014



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Energy Balance in Closed Systems



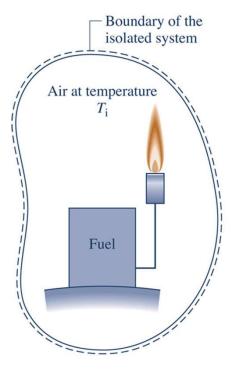




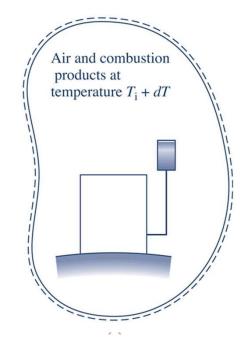


Energy Balance in isolated systems

• What happens to the total energy of the system?



Moran et al., 2014

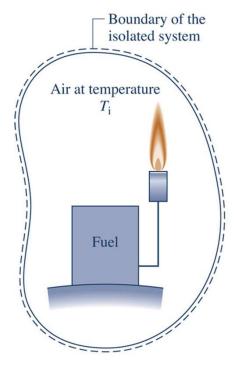


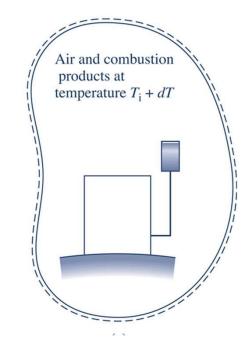


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Energy vs. Exergy

What happens to the total energy of the system?



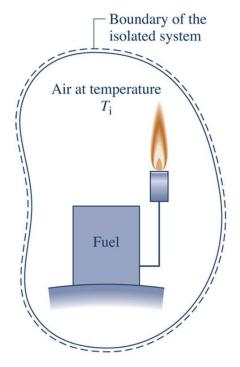


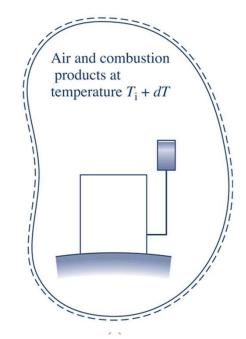
• It remains constant (system is isolated)



Energy vs. Exergy

• Can the process occur from the right to the left?







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Is the first law enough?

What happens if you put a dish with ice over a pan with boiling water?





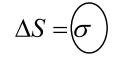
Entropy Balance in Adiabatic Systems & Isolated Systems



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Entropy Balance in Adiabatic Systems & Isolated Systems

Entropy change = Entropy production

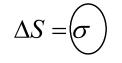


• 2nd Law: ?



INSTITUTO SUPERIOR TÉCNICO Entropy Balance in Adiabatic Systems & Isolated Systems

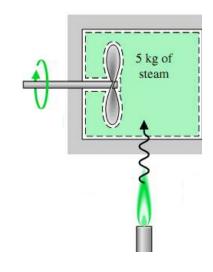
Entropy change = Entropy production



• 2nd Law: <u>In an adiabatic system the entropy</u> <u>must not decrease</u>



Entropy Balance in Closed Systems





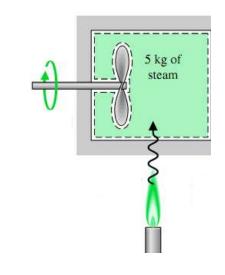
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Entropy Balance in Closed Systems

Entropy change = Entropy transfer in the form of heat + entropy production

$$\Delta S = \frac{Q}{T} + \sigma$$

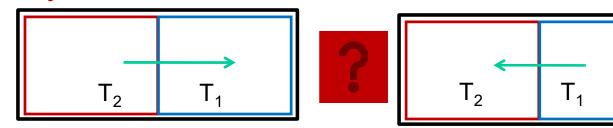
Entropy flows with heat but not with work





Entropy Balance in Adiabatic Systems

- Suppose the combined system (contained in the black boundary) is adiabatic and that $T_2 > T_1$
- What happens to the entropy of the combined system in each case?





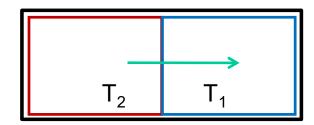
Entropy Balance in Adiabatic Systems

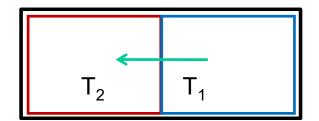
 $\Delta S = \sigma > 0$

$$\Delta S = \Delta S_1 + \Delta S_2 = \frac{Q}{T_1} - \frac{Q}{T_2}$$

$$\Delta S = \sigma > 0$$

$$\Delta S = \Delta S_1 + \Delta S_2 = \frac{Q}{T_2} - \frac{Q}{T_1}$$

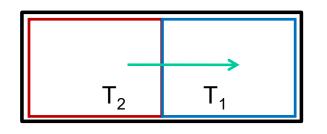


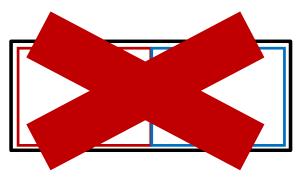




Entropy Balance in Adiabatic Systems

• <u>2nd Law: In an adiabatic system the entropy</u> <u>cannot decrease</u>



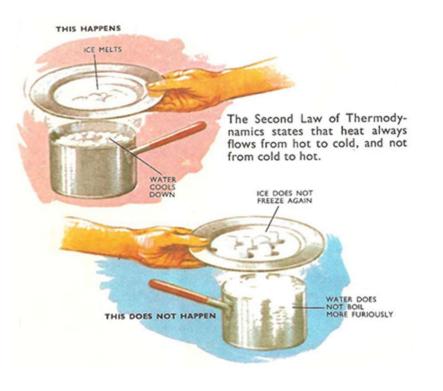


- 2nd Law: the arrow of time
- Entropy production in spontaneous processes



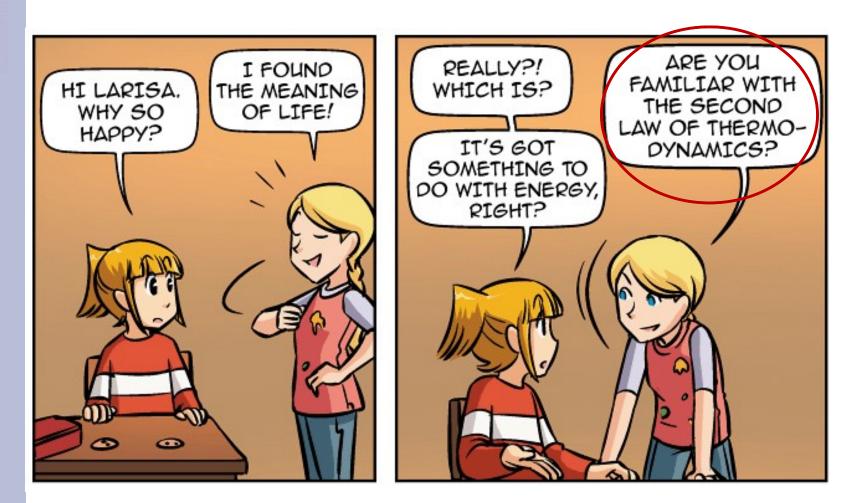
Is the first law enough?

• What happens if you put a dish with ice over a pan with boiling water?



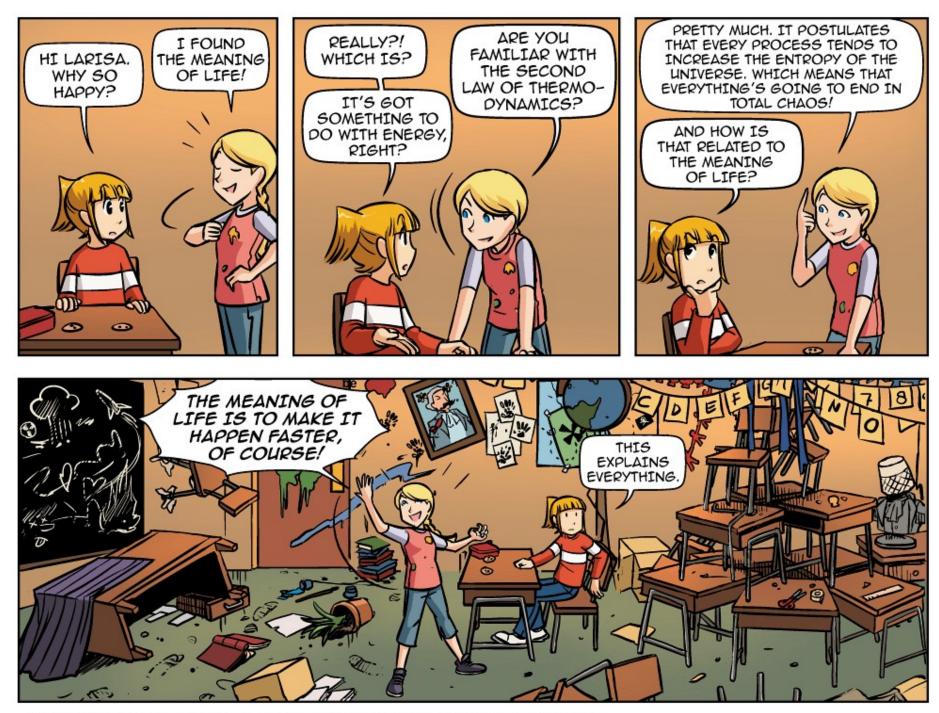


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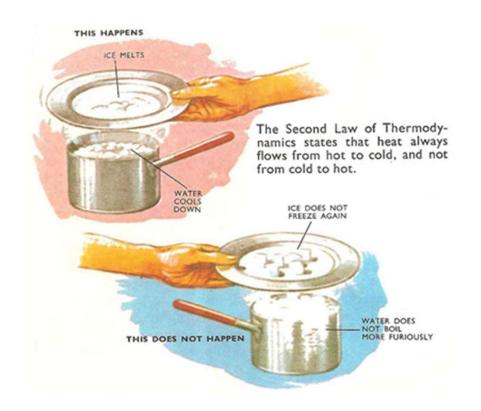


Sandra and Woo by Oliver Knörzer (writer), Powree (artist) and Lisa Moore (colorist) - www.sandraandwoo.com



The state variable: Entropy

• Entropy is the state variable that gives unidirectionality to time in physical processes ocurring in isolated & adiabatic systems.



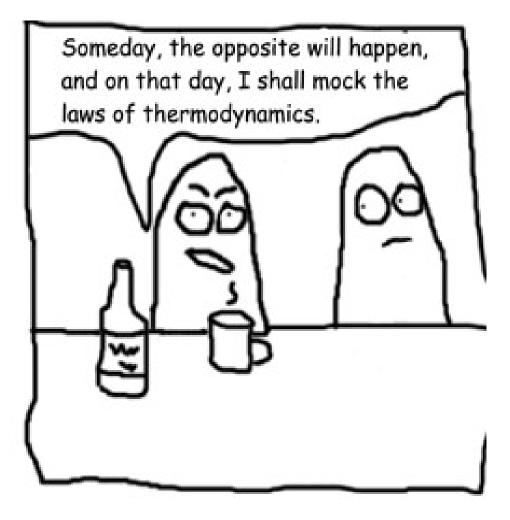














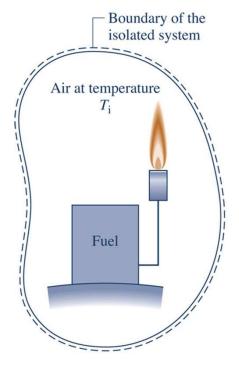


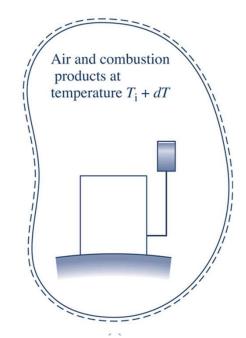


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Entropy

What happens to the total entropy of the system?

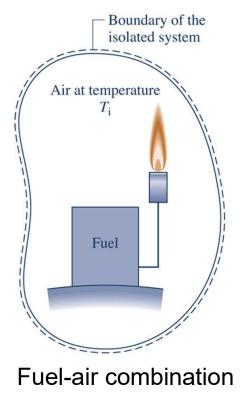


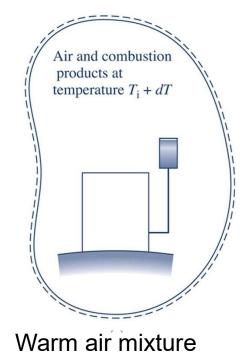


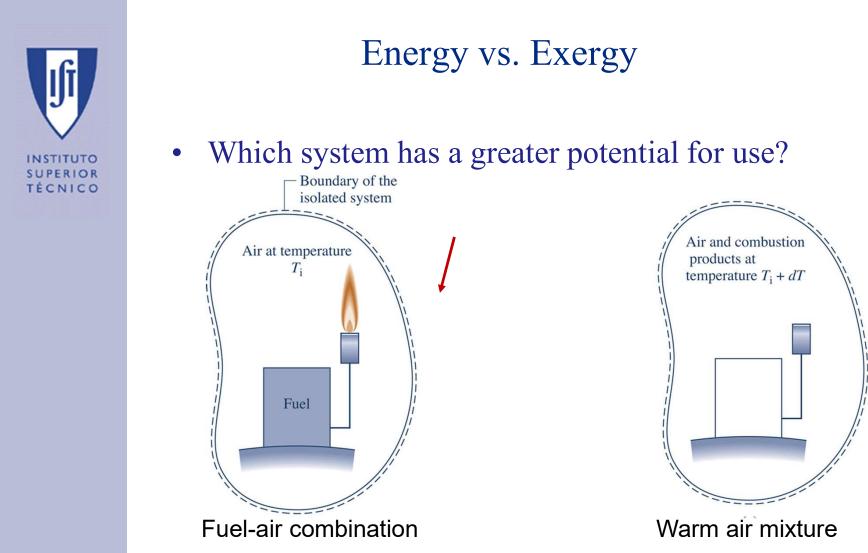


Energy vs. Exergy

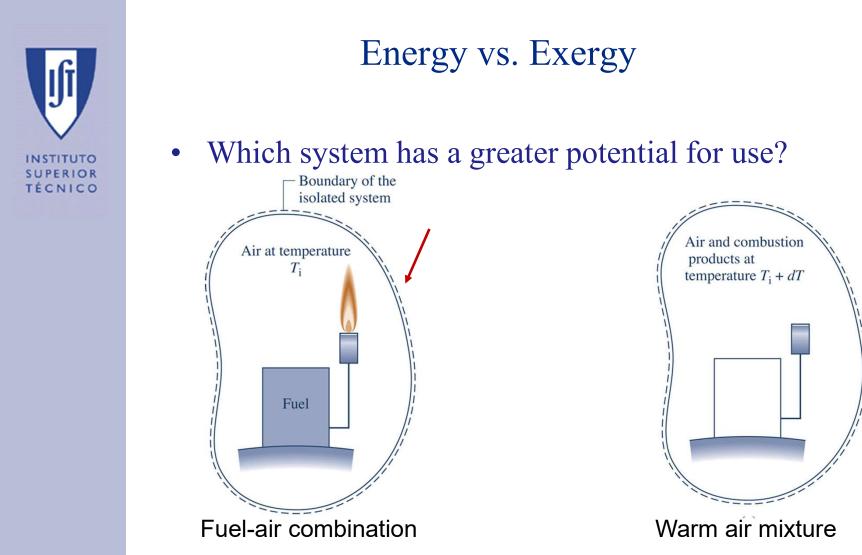
• Which system has a greater potential for use?







• "The fuel might be used to generate electricity, produce steam, or power a car whereas the final warm mixture is clearly unsuited for such applications."



• "During the process the initial potential for use (and economic value) is predominately *destroyed* owing to the irreversible nature of that process."



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Energy vs. Exergy

- Exergy is the property that quantifies the potential for use and it is exergy that has economic value.
- Exergy is a property that takes into account the second law of thermodynamics
- How can we make use of a body at $T_i > T_0$?

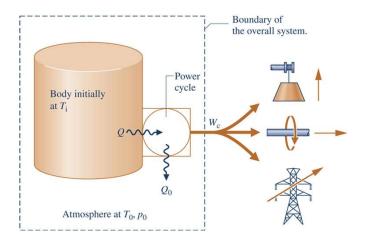




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Energy vs. Exergy

Controlled cooling to produce work



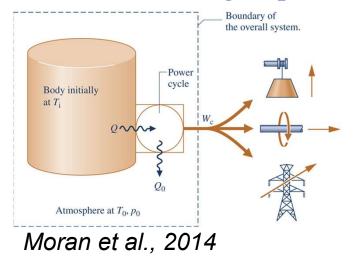
Moran et al., 2014



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Energy vs. Exergy

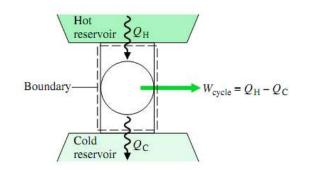
Controlled cooling to produce work



• The amount of maximum work depends on what?



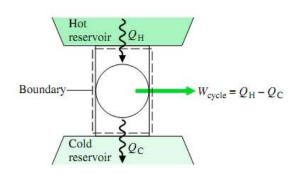
The Carnot Efficiency – Power Cycle







The Carnot Efficiency – Power Cycle





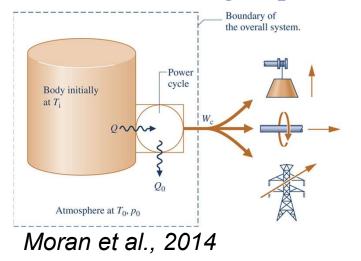
- First law: $\Delta E = Q_H Q_C W_{cycle} = 0 \Rightarrow W_{cycle} = Q_H Q_C$
- Second law: $\Delta S = \frac{Q_H}{T_H} \frac{Q_C}{T_C} + \sigma = 0 \Longrightarrow Q_C \neq 0$
- Ideal (Reversible) Cycle: $\Delta S = \frac{Q_H}{T_H} - \frac{Q_C}{T_C} = 0$ $\eta_{ideal} = \frac{W_{cycle}}{Q_H} = \frac{Q_H - Q_C}{Q_H} = 1 - \frac{Q_C}{Q_H} = 1 - \frac{T_C}{T_H}$



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Energy vs. Exergy

Controlled cooling to produce work



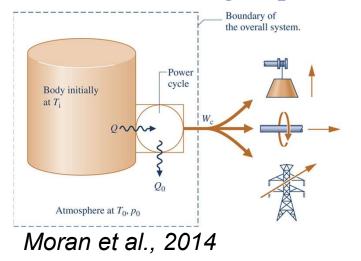
- Exergy is the maximum theoretical value for the work $W_{\rm c}$.
- What happens to the exergy of the body with time?



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Energy vs. Exergy

Controlled cooling to produce work

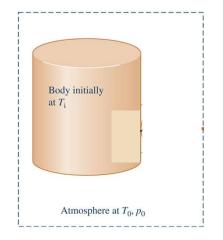


- Exergy is the maximum theoretical value for the work $W_{\rm c}$.
- What happens to the exergy of the body with time?



Energy vs. Exergy

• Could we produce work if $T_i < T_0$?

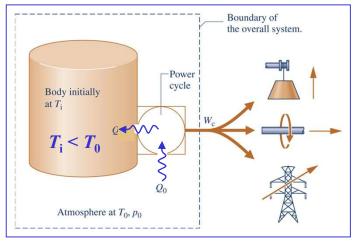




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Energy vs. Exergy

Could we produce work if $T_i < T_0$?



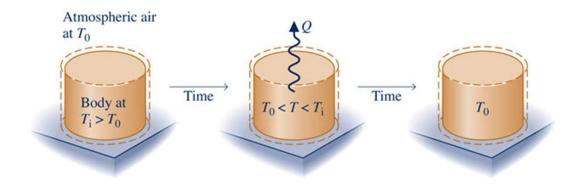
Moran et al., 2014

• The potential for developing work W_c exists because the initial state of the body differs from that of the environment, i.e., $T_i < T_0$.



Defining Exergy

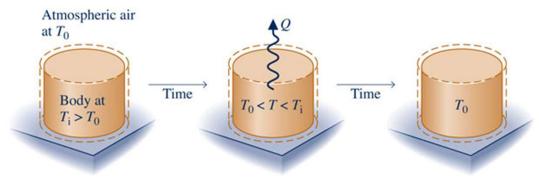
- Reference environment: T₀ and P₀
- Exergy is the maximum $W_{\rm c}$ obtainable from the system plus the environment as the system comes into equilibrium with the environment (goes to the dead state $T_0 \& P_0$).
- What happened in the case of spontaneous cooling?





Defining Exergy

- Reference environment: T₀ and P₀
- Exergy is the maximum W_c obtainable from the system plus the environment as the system comes into equilibrium with the environment (goes to the dead state $T_0 \& P_0$).
- W_{c} is null because exergy is destroyed by irreversibilities (spontaneous processes)





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Defining Exergy

If temperature and/or pressure of a system differ from that of the environment, the system has **thermomechanical** exergy. Another contribution, called **chemical exergy**, arises when there is a chemical composition difference between the system and environment.



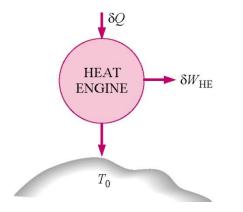




$$E_{2} - E_{1} = \underbrace{\int_{0}^{2} \left(1 - \frac{T_{0}}{T_{b}}\right) \delta Q}_{\text{exergy}} + \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma$$

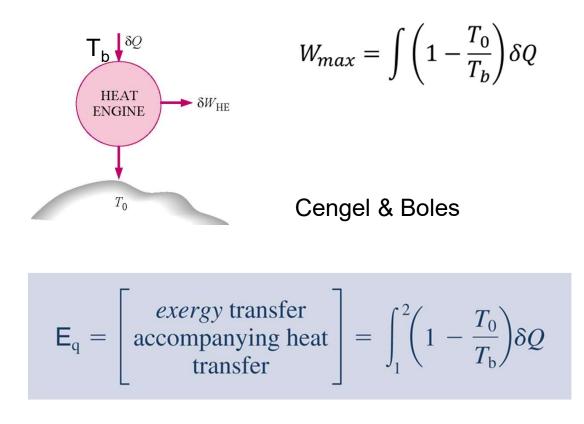


- Exergy transfer by heat
 - What is the maximum amount of work that can be produced with Q?





- Exergy transfer by heat
 - What is the maximum amount of work that can be produced with Q?

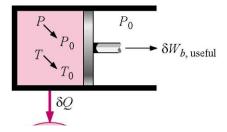




$$\mathbf{E}_{2} - \mathbf{E}_{1} = \frac{\int_{1}^{2} \left(1 - \frac{T_{0}}{T_{b}}\right) \delta Q \left(+ \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma \right) dV_{0} + \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma$$
exergy change



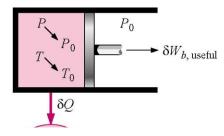
- Exergy transfer by work
 - What is the maximum amount of useful work that can be used from W?



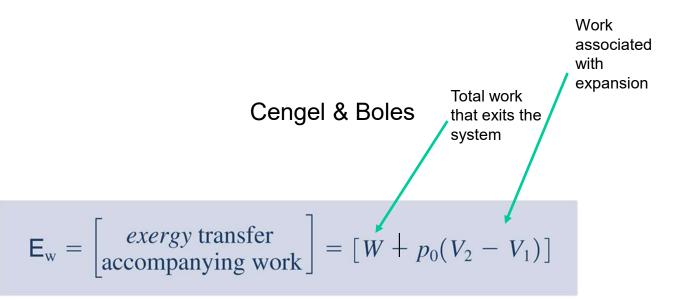


Exergy Balance in Closed Systems

• Exergy transfer by work



$$W = W_{b,useful} - \int P_0 \, dV$$
$$W_{b,useful} = W + P_0(V_2 - V_1)$$





$$\frac{\mathsf{E}_2 - \mathsf{E}_1}{\underset{\text{change}}{\text{exergy}}{\text{change}}} = \frac{\int_1^2 \left(1 - \frac{T_0}{T_b}\right) \delta Q + \left[W + p_0(V_2 - V_1)\right] - \underbrace{T_0 \sigma}$$



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Exergy Balance in Closed Systems

• Exergy destruction: irreversibilities destroy exergy

$$\mathsf{E}_{\mathrm{d}} = T_0 \sigma$$

• 2nd Law?



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Exergy Balance in Closed Systems

• Exergy destruction: irreversibilities destroy exergy

$$\mathsf{E}_{\mathrm{d}} = T_0 \sigma$$

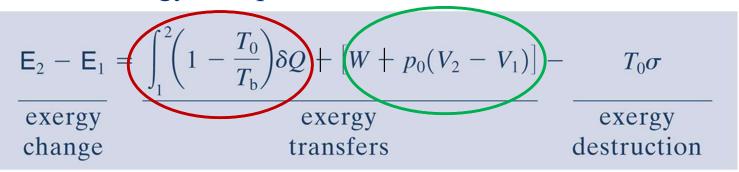
• 2nd Law?

 $E_{d}: \begin{cases} = 0 \text{ (no irreversibilities present within the system)} \\ > 0 \text{ (irreversibilities present within the system)} \\ < 0 \text{ (impossible)} \end{cases}$



Exergy Balance in Closed Systems

Exergy Change = exergy transfer by heat - exergy transfer by work - exergy dissipation



• What happens to the exergy of an isolated system?



Exergy Rate Balance for Closed Systems

• The exergy rate balance?

$$\frac{\mathsf{E}_{2} - \mathsf{E}_{1}}{\underset{\text{change}}{\text{exergy}}{\text{change}}} = \frac{\int_{1}^{2} \left(1 - \frac{T_{0}}{T_{b}}\right) \delta Q + \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma}{\underset{\text{exergy}}{\text{exergy}}} - \frac{T_{0}\sigma}{\underset{\text{destruction}}{\text{exergy}}}$$



Exergy Rate Balance for Closed Systems

• The exergy rate balance

$$\frac{\mathsf{E}_{2} - \mathsf{E}_{1}}{\frac{\mathsf{exergy}}{\mathsf{exergy}}} = \frac{\int_{1}^{2} \left(1 - \frac{T_{0}}{T_{b}}\right) \delta Q + \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma}{\frac{\mathsf{exergy}}{\mathsf{exergy}}} \frac{\mathsf{exergy}}{\mathsf{destruction}}$$

$$\frac{d\mathsf{E}}{dt} = \sum_{j} \left(1 - \frac{T_{0}}{T_{j}}\right) \dot{Q}_{j} + \left(\dot{W} + p_{0}\frac{dV}{dt}\right) - \dot{\mathsf{E}}_{d}$$

• Steady-state?



Exergy Rate Balance for Closed Systems

• The exergy rate balance

$$\frac{\mathsf{E}_{2} - \mathsf{E}_{1}}{\frac{\mathsf{exergy}}{\mathsf{exergy}}} = \frac{\int_{1}^{2} \left(1 - \frac{T_{0}}{T_{b}}\right) \delta Q + \left[W + p_{0}(V_{2} - V_{1})\right] - T_{0}\sigma}{\frac{\mathsf{exergy}}{\mathsf{exergy}}} = \frac{T_{0}\sigma}{\frac{\mathsf{exergy}}{\mathsf{destruction}}}$$

$$\frac{d\mathsf{E}}{dt} = \sum_{i} \left(1 - \frac{T_{0}}{T_{i}}\right) \dot{Q}_{i} + \left(\dot{W} + p_{0}\frac{dV}{dt}\right) - \dot{\mathsf{E}}_{d}$$

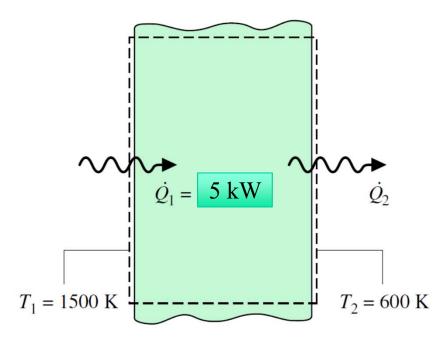
• Steady-state

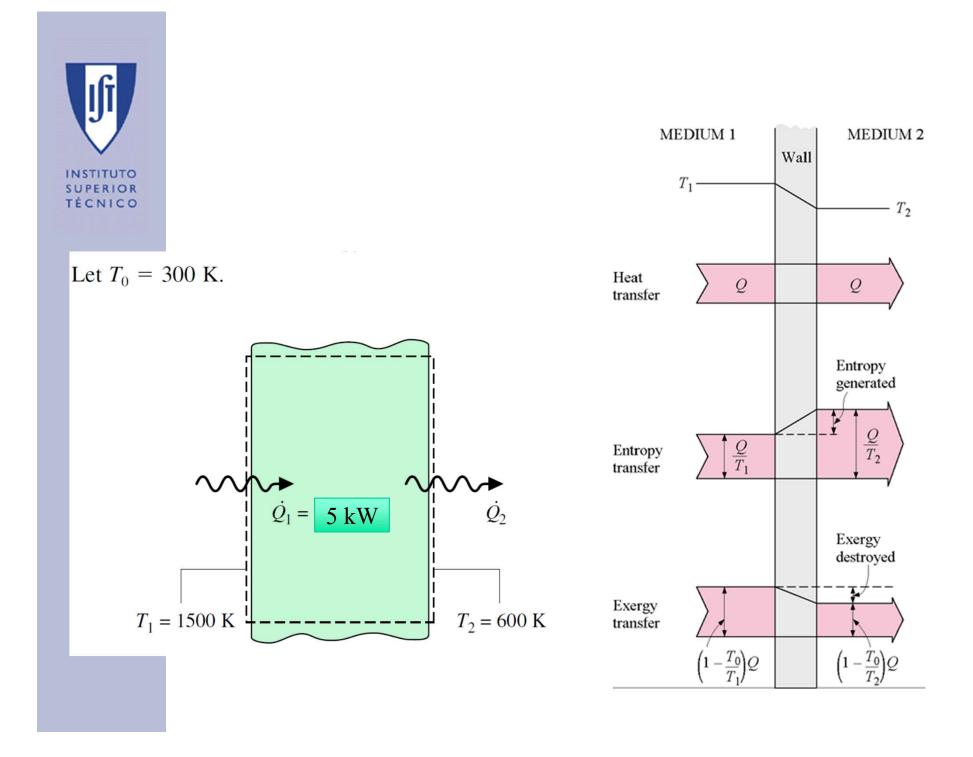
$$0 = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \dot{W} - \dot{\mathsf{E}}_{\mathsf{d}}$$



$$0 = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \dot{W} - \dot{\mathsf{E}}_{\mathsf{d}}$$

Let
$$T_0 = 300$$
 K.

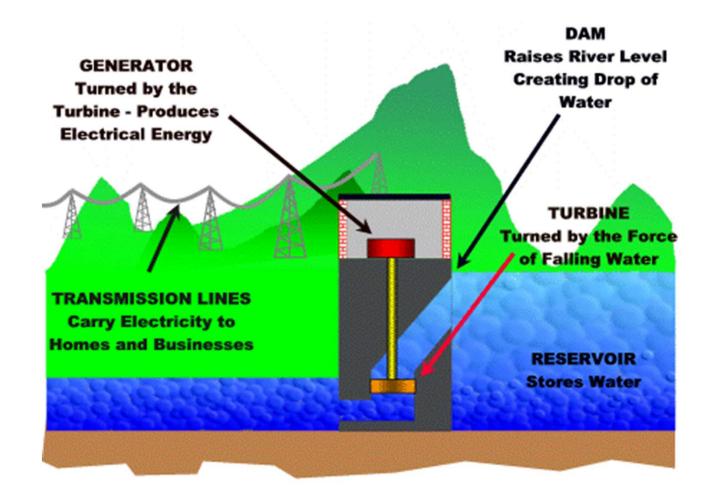






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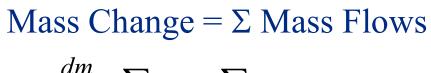
Energy Balance in Open Systems



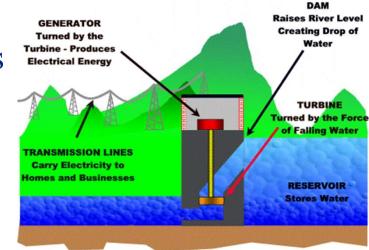


Energy Balance in Open Systems

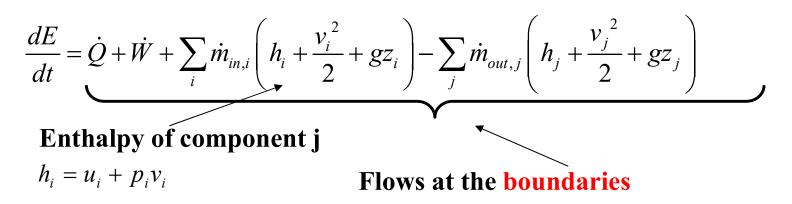
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$$\frac{dm}{dt} = \sum_{i} \dot{m}_{in,i} - \sum_{j} \dot{m}_{out,j}$$



Energy Change = Heat + Work + Energy in Mass Flow





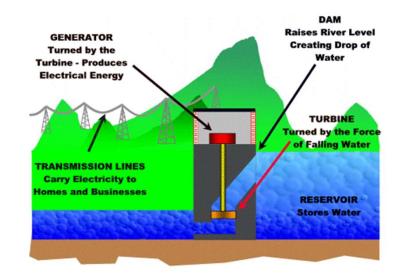
Energy Balance in Open Systems





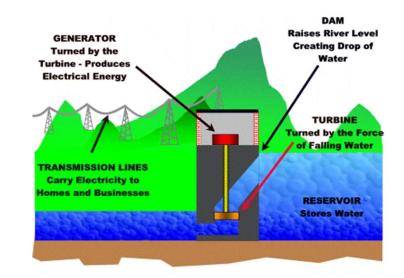
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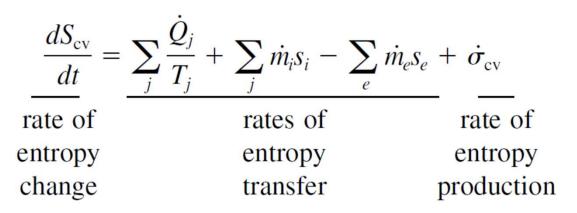
Entropy Balance in Open Systems





Entropy Balance in Open Systems







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Exergy Rate Balance for Control Volume Systems

The exergy rate balance for closed systems

$$\frac{d\mathsf{E}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \left(\dot{W} + p_0 \frac{dV}{dt} \right) - \dot{\mathsf{E}}_d$$

• The exergy rate balance for open systems



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Exergy Rate Balance for Control Volume Systems

The exergy rate balance for closed systems

$$\frac{d\mathsf{E}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \left(\dot{W} + p_0 \frac{dV}{dt} \right) - \dot{\mathsf{E}}_d$$

• The exergy rate balance for open systems

$$\frac{d\mathbf{E}_{cv}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \left(\dot{W}_{cv} \vdash p_0 \frac{dV_{cv}}{dt} \right) + \sum_{i} \dot{m}_i \mathbf{e}_{fi} - \sum_{e} \dot{m}_e \mathbf{e}_{fe} - \dot{\mathbf{E}}_d$$

• Steady-state?



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Exergy Rate Balance for Control Volume Systems

The exergy rate balance for closed systems

$$\frac{d\mathsf{E}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \left(\dot{W} + p_0 \frac{dV}{dt} \right) - \dot{\mathsf{E}}_d$$

• The exergy rate balance for open systems

$$\frac{d\mathbf{E}_{cv}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \left(\dot{W}_{cv} + p_0 \frac{dV_{cv}}{dt} \right) + \sum_{i} \dot{m}_i \mathbf{e}_{fi} - \sum_{e} \dot{m}_e \mathbf{e}_{fe} - \dot{\mathbf{E}}_d$$

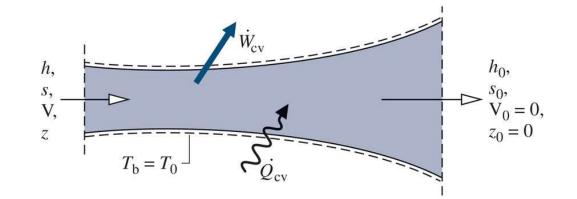
• Steady-state?

$$0 = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j + \dot{W}_{cv} + \dot{m} (\mathbf{e}_{f1} - \mathbf{e}_{f2}) - \dot{\mathsf{E}}_d$$



The specific flow exergy

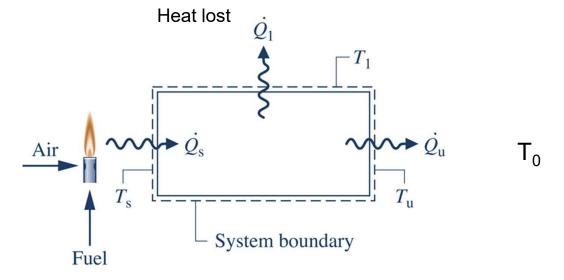
• What is the specific flow exergy?





Energy & Exergy Efficiencies

• Energy and exergy balances at steady-state?



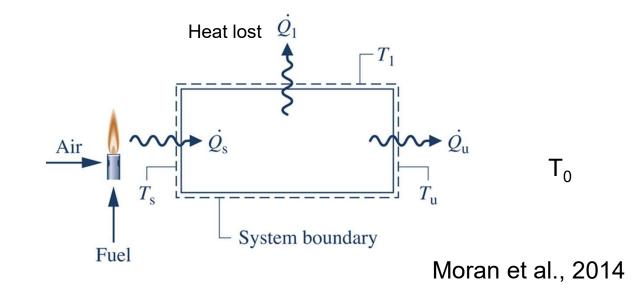
Moran et al., 2014



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Exergetic Efficiency

Energy and exergy balances at steady-state

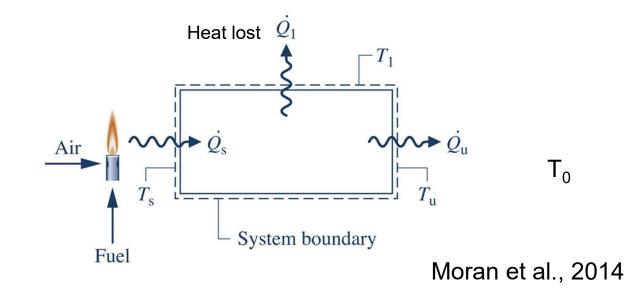


$$\frac{dE'}{dt}^{0} = (\dot{Q}_{s} - \dot{Q}_{u} - \dot{Q}_{1}) - \dot{W}^{0}$$

$$\frac{dE'}{dt}^{0} = \left[\left(1 - \frac{T_{0}}{T_{s}} \right) \dot{Q}_{s} - \left(1 - \frac{T_{0}}{T_{u}} \right) \dot{Q}_{u} - \left(1 - \frac{T_{0}}{T_{1}} \right) \dot{Q}_{1} \right] - \left[\dot{W}^{0} - p_{0} \frac{dV}{dt}^{0} \right] - \dot{\mathsf{E}}_{d}$$

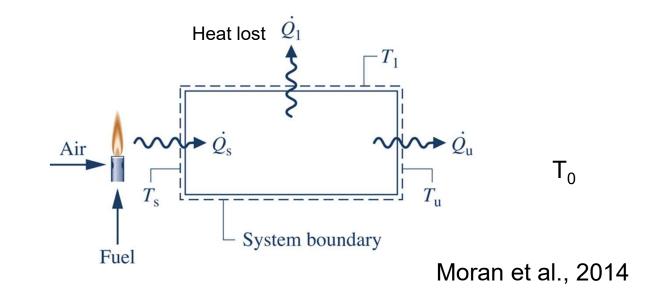


• Energy efficiency?





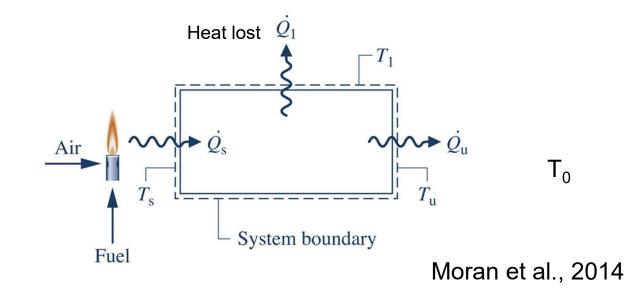
• Energy efficiency



$$\eta = rac{\dot{Q}_{u}}{\dot{Q}_{s}}$$

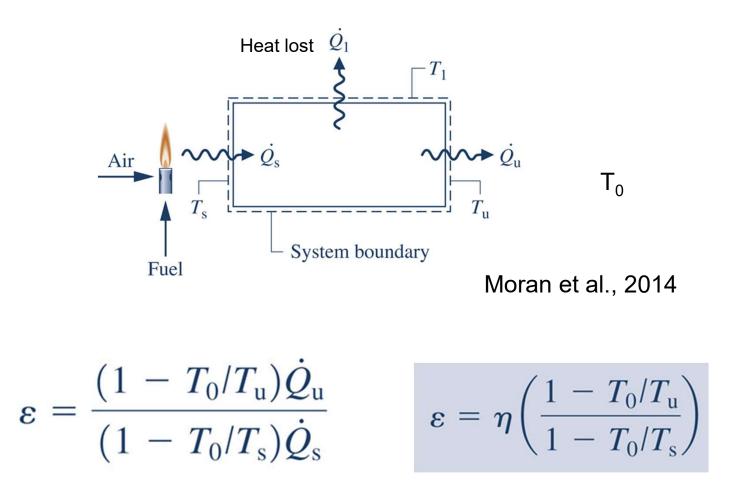


• Exergy efficiency?



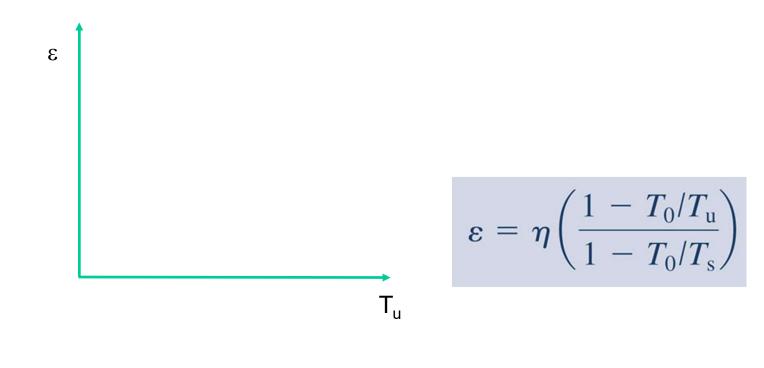


• Exergy efficiency





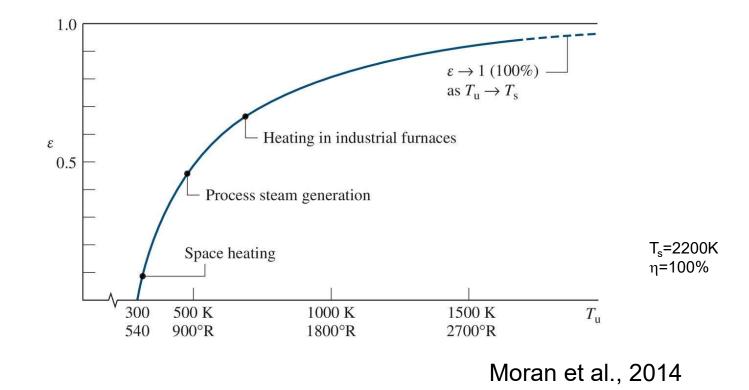
• How does exergy efficiency varies assuming $T_s=2200K$?



Moran et al., 2014



• Exergy efficiency



• Exergy analysis: (mis)match between energy used and end-use



Exergy efficiencies

• 2nd Law efficiency

$$\varepsilon = \frac{W_{max,output}}{W_{max,input}} \longleftarrow \quad \eta = \frac{E_{output}}{E_{input}}$$



Exergy efficiencies

• 2nd Law efficiency

$$\varepsilon = \frac{W_{max,output}}{W_{max,input}} \longleftarrow \quad \eta = \frac{E_{output}}{E_{input}}$$

• For a device which converts one form of mechanical energy to another $\varepsilon = \frac{W_{output}}{W_{input}} = \eta$





Household appliances with electric motors















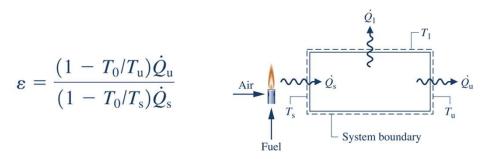


Exergy efficiencies

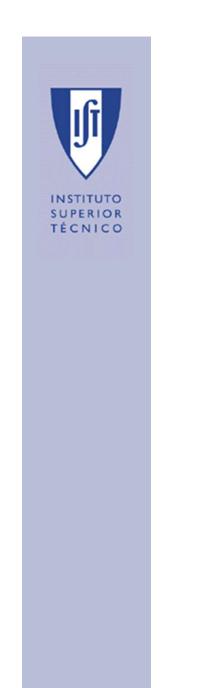
• 2nd Law efficiency

$$\varepsilon = \frac{W_{max,output}}{W_{max,input}} \longleftarrow \eta = \frac{E_{output}}{E_{input}}$$

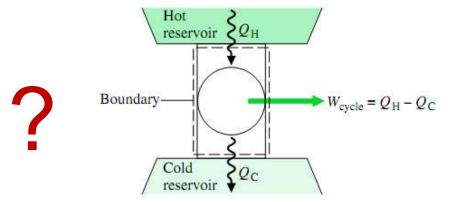
• When the input or output of a device is heat then heat must be downgraded into equivalent units of mechanical work (Cullen & Allwood, 2010)





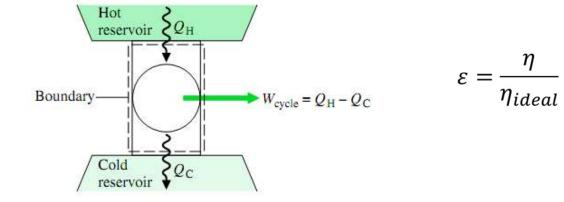


Exergy Efficiency – Power Cycle





Exergy Efficiency – Power Cycle





Exergy efficiencies

 Second law efficiencies are bounded (thay are less than or equal to 100%) - > they provide information on how much you can improve your efficiency.



Exergy efficiencies

• Exergy losses vs. Exergy destruction

$$\frac{d\mathsf{E}}{dt} = \sum_{j} \left(1 - \frac{T_0}{T_j} \right) \dot{Q}_j - \left(\dot{W} - p_0 \frac{dV}{dt} \right) - \dot{\mathsf{E}}_d$$

$$\varepsilon = \frac{\eta}{\gamma_{ideal}}$$





Exergy efficiencies

• 2nd Law efficiency

$$\varepsilon = \frac{W_{max,output}}{W_{max,input}} \quad \longleftarrow \quad \eta = \frac{E_{output}}{E_{input}}$$

• When the input is work (electricity) and the output is neither work nor heat (e.g. light or music then

$$\varepsilon = \frac{W_{output}}{W_{max,input}} =$$

$$=\frac{\eta}{\eta_{ideal}}=\frac{\eta}{683\,lm/W}$$

