




# Lecture 8 Chemistry 163B Winter 2020

## Introduction to 2<sup>nd</sup> Law




Chemistry 163B  
Winter 2020  
**Heuristic** Introduction  
Second Law,  
Statistics and Entropy

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lecture 8 objectives:   

### When and Why do things happen ??


(an overview of 2<sup>nd</sup> Law)

- exothermicity ( $q < 0$ ) often accompanies spontaneous processes, but not all; not a requirement
- can't find a repeatable (cyclic) process that fully converts heat (disorder) to work (order) 
- order and disorder in terms of microstates 
- the Universe meanders through the fields and meadows of microstates only to be observed in the 'state' corresponding to the maximum number of microstates!! 

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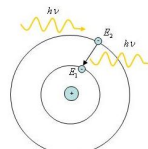
"Lower energy states are more stable"

Systems naturally go to lower energy ????



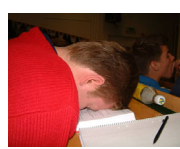
falling apples

[http://www.science museum.org.uk/online/real/falling\\_apples.html](http://www.science museum.org.uk/online/real/falling_apples.html)



excited state → ground state

[http://www.thespectroscopy.net/images/PLAS\\_EB\\_SIE.jpg](http://www.thespectroscopy.net/images/PLAS_EB_SIE.jpg)



chemistry 163B students

[http://www.moonbattery.com/sleeping\\_student.jpg](http://www.moonbattery.com/sleeping_student.jpg)

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But ..... ??? First Law of Thermodynamics

The energy lost has to go somewhere ???


Energy lost by system means energy gained by surroundings.

First Law of Thermodynamics  
 $\Delta U_{\text{sys}} = -\Delta U_{\text{surr}}$   
 $\Delta U_{\text{universe}} = 0$  (also consider  $E=mc^2$ )

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The second law of thermodynamics

- Why do things happen?
- What are the limitations on things that do occur spontaneously?
- exothermicity,  $\Delta H < 0$ , was considered by Berthelot to be the driving principle for spontaneity
- but this can't be true: some salts cool when dissolving and why do gasses diffuse if no energy difference



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SADI CARNOT

- born 1796, son of Lazare Carnot and antimonarchist
- named Sadi after Persian poet and mystic
- went to Polytechnique Ecole whose faculty included Lagrange, Fourier, Laplace, Berthelot, Ampere, duLong and had as classmates Cauchy, Coriolis, Poisson, Petit, Fresnel
- 1814 went into Corps of Engineers and when monarchy reestablished was sent to boondocks outpost
- 1824 wrote "Reflections on the Motive Power of Heat and Machines Adapted for Developing this Power"

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# Lecture 8 Chemistry 163B Winter 2020

## Introduction to 2<sup>nd</sup> Law

*Carnot cycle approach to 'what can happen': engines, work, efficiency*

- the 'Carnot Cycle' is a cyclic process (engine) of reversible (ideal) gas expansions and compressions ( $\Delta U=0$ )
- want process that does net w on surroundings i.e. convert heat to work [ $(W_{sys})_{total\ process} < 0; q_{sys} > 0$ ]
- we will employ Carnot cycle to show that
 

- "net disorder of universe" limits efficiency of the engine
  - analysis of the process will lead to a **NEW STATE FUNCTION** defined by  $\frac{dq_{rev}}{T}$  **an exact differential !!!**

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*Carnot cycle (E&R<sub>4th</sub>, pp. 124-129 and HW2 prob #10)*

**Problem #10**

1→2 isothermal reversible expansion at  $T_1$  (500K)  
 2→3 adiabatic reversible expansion  $T_1 \rightarrow T_2$  (457K)  
 3→4 isothermal compression at  $T_2$  (457K)  
 4→1 adiabatic compression  $T_2 \rightarrow T_1$  (500K)

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*Carnot cycle (graphical interpretation;  $w_{sys} = -\int P_{ext} dV$ ; review)*

expansions, steps I-II

$w_{sys} < 0, w_{surr} > 0 \quad T_{iso} = T_U$

compressions, steps III-IV

$w_{sys} > 0, w_{surr} < 0 \quad T_{iso} = T_L$

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*Carnot cycle: (graphical interpretation;  $w_{sys} = -\int P_{ext} dV$ ; review)*

expansions, steps I-II

$w_{sys} < 0, w_{surr} > 0$   
 $q_{in} \text{ at } T_{iso} = T_{Upper}$

compressions, steps III-IV

$w_{sys} > 0, w_{surr} < 0$   
 $q_{out} \text{ at } T_{iso} = T_{Lower}$

Carnot cyclic engine net work ON surroundings

$w_{net} > 0$   
 engine DOES work on surroundings

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*from Carnot cycle*

for system in complete cycle:  
 $\Delta U=0; q > 0; w < 0$  (work DONE on surr) (#10E)

$q > 0$  ( $q_{in}$ ) at higher  $T_1$ ;  $q < 0$  ( $q_{out}$ ) at lower  $T_2$

efficiency =  $-w/q_{1 \rightarrow 2}$   
 (how much net work out for heat in 1→2)

efficiency will depend on  $T_1$  and  $T_2$

HW prob #22  $\epsilon$  is efficiency

$$\epsilon = \frac{T_H - T_C}{T_H} \quad \text{or} \quad \epsilon = \frac{T_U - T_L}{T_U}$$

H=HOT C=COLD or U=UPPER L=LOWER

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*statements of the Second Law of Thermodynamics (also see handout)*

- Macroscopic properties of an isolated system eventually assume constant values (e.g. pressure in two bulbs of gas, becomes constant; two block of metal reach same T) [Andrews. p37]
- It is impossible to construct a device that operates in cycles and that converts heat into work without producing some other change in the surroundings. *Kelvin's Statement* [Raff p 157]; *Carnot Cycle*
- It is impossible to have a natural process which produces no other effect than absorption of heat from a colder body and discharge of heat to a warmer body. *Clausius's Statement, refrigerator*
- In the neighborhood of any prescribed initial state there are states which cannot be reached by any adiabatic process  
 ~ Carathéodory's statement [Andrews p. 58]

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## Introduction to 2<sup>nd</sup> Law

### Tolman's perpetual motion machine

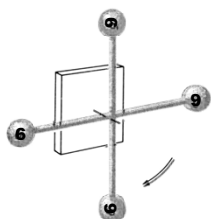


Figure 2-1 Tolman's perpetual motion machine. In his lectures on thermodynamics Professor R. C. Tolman used to delight in presenting this tongue-in-cheek example of a perpetuum mobile of the first kind. The numbers indicate weights, in arbitrary units.

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### first a look at disorder and its relation to entropy and 2<sup>nd</sup> Law

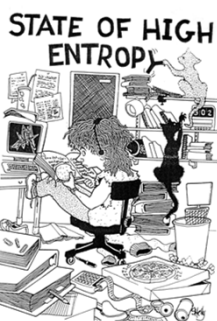


$$S = k \ln W$$

Boltzmann

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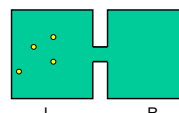
### disorder and entropy



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### microstates and macrostates

configuration 1 2 3 4	macrostate L R	W no. of microstates



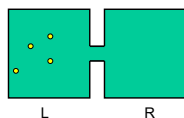
etc.

$$W = \frac{(n_{\text{total}})!}{(n_L)! (n_R)!} = \frac{24}{(n_L)! (n_R)!}$$

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### microstates and macrostates

configuration 1 2 3 4	macrostate L R	W no. of microstates
LLLL	4 0	1
LLLR LLRL LRLR RLLL	3 1	4
LLRR LRLR RLLR LRRL RLRL RRLR	2 2	6
~	1 3	4
~	0 4	1



etc.

$$W = \frac{(n_{\text{total}})!}{(n_L)! (n_R)!} = \frac{24}{(n_L)! (n_R)!}$$

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### macrostates and microstates

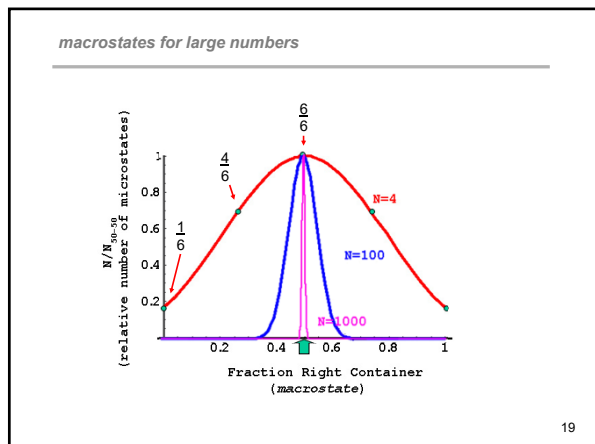
microstate: one of the **EQUALLY PROBABLE** configurations of molecules (e.g. LLLL vs LRLR)

macrostate: state with specific macroscopic properties  
e.g. L=2 R=2

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## Introduction to 2<sup>nd</sup> Law



moral of the story

although each allowed microstate (e.g. LLRR or LLLL) is equally probable

the overwhelming number of microstates correspond to macrostates with almost identical macroscopic properties (e.g. ~ 50-50 RvsL)

$W$ , the number of microstates corresponding to the macrostate, is a measure of the DISORDER of the system in that macrostate

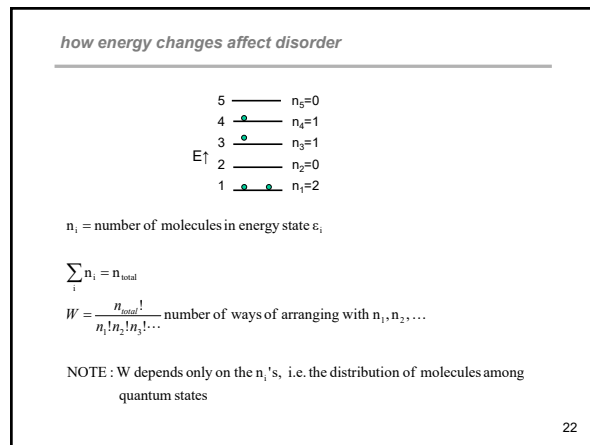
a system "meanders" through all available microstates; but you are only likely to observe it in one of the overwhelming number that correspond to the equilibrium macrostate

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famous quote

**“disorder happens”**

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how energy changes affect disorder

$$U = E_{\text{total}} = \sum_i n_i \epsilon_i$$

$$dE = \sum_i n_i d\epsilon_i + \sum_i \epsilon_i dn_i$$

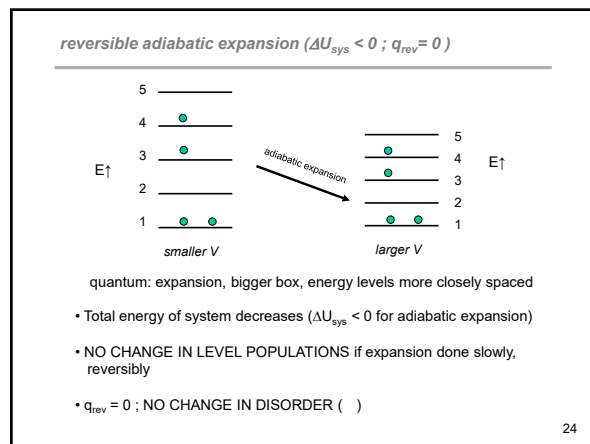
change in energy due to change in energy levels, e.g. 3D quantum p.i.b. change in energy levels as box changes size

**work**

change in energy due to change redistribution of molecules among energy levels, e.g. put in more total energy to fixed size 3D quantum p.i.b.

**heat**

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# Lecture 8 Chemistry 163B Winter 2020

## Introduction to 2<sup>nd</sup> Law

reversible isothermal expansion ideal gas ( $\Delta U_{sys} = 0$ ;  $q_{rev} > 0$ ;  $w = -q$ )

to maintain  $\Delta U = 0$ ,  $\Delta T = 0$  need to put in heat

- Levels get closer due to  $\Delta V > 0$ ;  $w < 0$
- To maintain  $\Delta U = 0$ ,  $q > 0$  and the level populations have to change and thus  $W$  changes
- $q_{rev} > 0$ ; INCREASE IN , INCREASE IN DISORDER

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reversible isothermal expansion ideal gas ( $\Delta U_{sys} = 0$ ;  $q_{rev} > 0$ ;  $w = -q$ )

to maintain  $\Delta U = 0$ ,  $\Delta T = 0$  need to put in heat

- Levels get closer due to  $\Delta V > 0$ ;  $w < 0$
- To maintain  $\Delta U = 0$ ,  $q > 0$  and the level populations have to change and thus  $W$  changes
- $q_{rev} > 0$ ; INCREASE IN , INCREASE IN DISORDER

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What happens to as thermal energy raised?

(disorder) increases

- $q_{rev} > 0$ ; INCREASE IN , INCREASE IN DISORDER

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What happens to as thermal energy raised?

(disorder) increases

- $q_{rev} > 0$ ; INCREASE IN , INCREASE IN DISORDER

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What happens to as thermal energy goes to zero ( $T \rightarrow 0K$ ) ?

decreases, in fact  $\rightarrow 1$   
only 'one way to distribute at 0°K

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What happens to as thermal energy goes to zero ( $T \rightarrow 0K$ ) ?

decreases, in fact  $\rightarrow 1$   
only 'one way to distribute at 0°K

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# Lecture 8 Chemistry 163B Winter 2020

## Introduction to 2<sup>nd</sup> Law

### Boltzmann and Entropy

S is entropy  
 $k = \text{Boltzmann's constant} = 1.3807 \times 10^{-23} \text{ J K}^{-1}$   
 disorder increases  $\Leftrightarrow$  entropy increases

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### take home messages

- Disorder,  $\Upsilon$ , did not change during an adiabatic reversible expansion ( $q_{\text{rev}} = 0$ )
- Disorder,  $\Upsilon$ , increased in isothermal reversible expansion ( $q_{\text{rev}} > 0$ )
- Disorder,  $\Upsilon$ , increased with T increase ( $q > 0$ )
- Disorder,  $\Upsilon$ , decreased with T decrease ( $q < 0$ )
- As  $T \rightarrow 0$ ,  $\Upsilon \rightarrow 1$

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### take home messages

A sample problem<sup>†</sup>:

Two bodies 500 molecules each  
 one at  $T_A = .63 \times T_f$   
 the other at  $T_B = 1.37 \times T_f$

bring into thermal contact

STAY COLD-HOT ??

or

EQUILIBRIUM ??

W (microstates)

$T_A = .63 T_f$  +  $T_B = 1.37 T_f \approx 10^{202} \times 10^{354}$

$.63 T_f$  |  $1.37 T_f \approx 10^{556} = 10^{202} \times 10^{354}$

or

$T_E = T_f \approx 10^{594}$

<sup>†</sup> adapted from Nash, *ChemThermo*, Addison Wesley, pp 175-176

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### take home message continued<sup>†</sup>

The equilibrium macrostate is  $\frac{10^{202}}{10^{354}} = 10^{38}$  time more likely than the hot-cold state, even though every (microstate)<sub>hot-cold</sub> has the same likelihood as a (microstate)<sub>equilibrium</sub>.

No more than one time in  $10^{38}$  a measurement will find the blocks in a half-hot and half-cold configuration.

If you had observed the microstate of the system  $10^6$  times a second constantly (without a msec of rest!) from the beginning of the universe until your midterm Friday ( $10^{10}$  years) **the odds against ever seeing a (microstate)<sub>hot-cold</sub> are 1:10<sup>15</sup> !!!**

<sup>†</sup> adapted from Nash, *ChemThermo*, Addison Wesley, pp 175-176

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### take home message continued<sup>†</sup>



... .. a progressive increase in "disorder" necessarily accompanies an approach to equilibrium characterized by the assumption of macrostates with ever-increasing values of W. And **what may at first appear to be a purposeful "drive" towards states of maximal disorder**, can now be seen to arise from **the operation of blind chance in an assembly** where all microstates remain equally probable, but where the overwhelming proportion of microstates is associated with the maximally disordered (nearly identical) macrostates corresponding to equilibrium macroscopic properties.

<sup>†</sup> adapted from Nash, *ChemThermo*, Addison Wesley, p 26.

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### MUCH MORE

**much more molecules, probability, statistical mechanics**

**CHEMISTRY 163C**

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# Lecture 8 Chemistry 163B Winter 2020

## Introduction to 2<sup>nd</sup> Law

**$q_{rev}$  VS  $q_{irrev}$  : some examples from 1<sup>st</sup> Law Calculations**

from lectures 2-3-4

Process 10 atm → 1atm 300K → $T_f$	$q_{rev}$	$q_{irrev}$	$(T_f)_{rev}$	$(T_f)_{irrev}$
isothermal				
adiabatic				

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**$q_{rev}$  VS  $q_{irrev}$  : some examples from 1<sup>st</sup> Law Calculations**

from lectures 2-3-4

Process 10 atm → 1atm 300K → $T_f$	$q_{rev}$	$q_{irrev}$	$(T_f)_{rev}$	$(T_f)_{irrev}$
isothermal	5743J	2244 J	300 K	300K
adiabatic	0	0 192K → 119K ( $C_v \Delta T$ ) -1517J	119K	192K ↓ 119K

for these expansions:  
same initial and final states  **$q_{rev} > q_{irrev}$**

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lecture 8 objectives:

### When and Why do things happen ??

(an overview of 2<sup>nd</sup> Law)

- ✓ exothermicity ( $q < 0$ ) often accompanies spontaneous processes, but not all; not a requirement
- ✓ can't find a repeatable (cyclic) process that fully converts heat (disorder) to work (order)
- ✓ order and disorder in terms of microstates
- ✓ the Universe meanders through the fields and meadows of microstates !!

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yes dear Juliet, you have nice eyes BUT  
--heat, work, efficiency, probability,  
disorder, entropy--  
the 2<sup>nd</sup> Law, it is so beautiful

oh Romeo my Romeo

### End Of Introductory Lecture on Second Law and Disorder

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بنا آدم اعضا یک یک یکند  
که در آفرینش یک کومند  
هر عضوی مدد آورده نگارد  
دیگر عضوانمانند قرار

**literal translation**  
(thank you Farzaneh):

Of one Essence is the human race  
thus has Creation put the base  
One Limb impacted is sufficient  
For all Others to feel the Mace

—Saadi (1184–1283)

**Persian Poet 13<sup>th</sup> Century**

Saadi

**Nicolas Léonard Sadi Carnot**  
(1796-1832) in the dress uniform  
of a student of the  
[Ecole Polytechnique](#)  $\epsilon = 1 - \frac{T_L}{T_U}$

**French Thermodynamicist**  
**Namesake 19<sup>th</sup> century**

Sadi

interpretative translation on UN building:  
Human beings are members of a whole,  
In creation of one essence and soul,  
If one member is afflicted with pain,  
Other members uneasy will remain.  
If you have no sympathy for human pain,  
The name of human you cannot retain.