# Entropy, Strings, and Partitions of Integers 

Jonathan Tannenhauser<br>ANOMALY<br>Wellesley College<br>November 4, 2010

## 1. Preliminaries

A partition of a positive integer $N$ is a way of writing $N$ as a sum of positive integers.

## 1. Preliminaries

A partition of a positive integer $N$ is a way of writing $N$ as a sum of positive integers.

How many partitions does the positive integer $N$ have?
Call this number $p(N)$.

## 1. Preliminaries

A partition of a positive integer $N$ is a way of writing $N$ as a sum of positive integers.

How many partitions does the positive integer $N$ have?
Call this number $p(N)$.
Example. $p(4)=5$ :

- 4
- $3+1$
- $2+2$
- $2+1+1$
- $1+1+1+1$

There's a formula for $p(N)$ :

$$
p(N)=\frac{1}{\pi \sqrt{2}} \sum_{k=1}^{\infty} A_{k}(N) \sqrt{k} \frac{d}{d N}\left(\frac{\sinh \left(\frac{\pi}{k} \sqrt{\frac{2}{3}\left(N-\frac{1}{24}\right)}\right)}{\sqrt{N-\frac{1}{24}}}\right)
$$

where

$$
A_{k}(N)=\sum_{0 \leq m<k,(m, k)=1} e^{\pi i(s(m, k)-2 N m / k)}
$$

and

$$
s(m, k)=\sum_{j=1}^{k-1} \frac{j}{k}\left(\frac{j m}{k}-\left\lfloor\frac{j m}{k}\right\rfloor-\frac{1}{2}\right) .
$$

(Rademacher 1937)

In 1918, Hardy and Ramanujan proved an asymptotic formula for $p(N)$ :

For large $N$,

$$
p(N) \sim \frac{1}{4 N \sqrt{3}} \exp \left(2 \pi \sqrt{\frac{N}{6}}\right)
$$

In 1918, Hardy and Ramanujan proved an asymptotic formula for $p(N)$ :

For large $N$,

$$
p(N) \sim \frac{1}{4 N \sqrt{3}} \exp \left(2 \pi \sqrt{\frac{N}{6}}\right)
$$

Our Goal: To prove (the red part of) this formula.

In 1918, Hardy and Ramanujan proved an asymptotic formula for $p(N)$ :

For large $N$,

$$
p(N) \sim \frac{1}{4 N \sqrt{3}} \exp \left(2 \pi \sqrt{\frac{N}{6}}\right)
$$

Our Goal: To prove (the red part of) this formula.

$$
\ln p(N) \sim 2 \pi \sqrt{\frac{N}{6}}
$$

## Proposition. (Euler)

$$
\sum_{N=0}^{\infty} p(N) x^{N}=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}
$$

## Proposition. (Euler)

$$
\sum_{N=0}^{\infty} p(N) x^{N}=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}
$$

Proof. Expand the right-hand side:

$$
\begin{gathered}
\left(1+x^{1}+\left(x^{1}\right)^{2}+\cdots\right)\left(1+x^{2}+\left(x^{2}\right)^{2}+\cdots\right)\left(1+x^{3}+\left(x^{3}\right)^{2} \cdots\right) \cdots . \\
\ell=1=2
\end{gathered}
$$

## Proposition. (Euler)

$$
\sum_{N=0}^{\infty} p(N) x^{N}=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}
$$

Proof. Expand the right-hand side:

$$
\begin{array}{cc}
\left(1+x^{1}+\left(x^{1}\right)^{2}+\cdots\right)\left(1+x^{2}+\left(x^{2}\right)^{2}+\cdots\right)\left(1+x^{3}+\left(x^{3}\right)^{2} \cdots\right) \cdots \\
\ell=1 & \ell=2
\end{array} \ell=3
$$

Each partition of $N$ gives one way of realizing $x^{N}$ in the product.

## Proposition. (Euler)

$$
\sum_{N=0}^{\infty} p(N) x^{N}=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}
$$

Proof. Expand the right-hand side:

$$
\begin{array}{cc}
\left(1+x^{1}+\left(x^{1}\right)^{2}+\cdots\right)\left(1+x^{2}+\left(x^{2}\right)^{2}+\cdots\right)\left(1+x^{3}+\left(x^{3}\right)^{2} \cdots\right) \cdots \\
\ell=1 & \ell=2
\end{array} \ell=3
$$

Each partition of $N$ gives one way of realizing $x^{N}$ in the product.
For example, the partition $7=1+1+2+3$ is realized by the blue terms.

## Proposition. (Euler)

$$
\sum_{N=0}^{\infty} p(N) x^{N}=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}
$$

Proof. Expand the right-hand side:

$$
\begin{gathered}
\left(1+x^{1}+\left(x^{1}\right)^{2}+\cdots\right)\left(1+x^{2}+\left(x^{2}\right)^{2}+\cdots\right)\left(1+x^{3}+\left(x^{3}\right)^{2} \cdots\right) \cdots \\
\ell=1 \\
\ell=2
\end{gathered}
$$

Each partition of $N$ gives one way of realizing $x^{N}$ in the product.
For example, the partition $7=1+1+2+3$ is realized by the blue terms.

Therefore, the coefficient of $x^{N}$ in the product is the number of partitions of $N$.

## 2. The Quantum String

## 2. The Quantum String

A harmonic oscillator is a system that vibrates back and forth at some frequency $\omega$, such as a mass on a spring.
$\oplus$


## 2. The Quantum String

A harmonic oscillator is a system that vibrates back and forth at some frequency $\omega$, such as a mass on a spring.
$\oplus$


In classical mechanics, the energy of an oscillator can be any non-negative number.

## 2. The Quantum String

A harmonic oscillator is a system that vibrates back and forth at some frequency $\omega$, such as a mass on a spring.


0

In classical mechanics, the energy of an oscillator can be any non-negative number.

In quantum mechanics, the only permitted energies are of the form

$$
E_{n}=n \hbar \omega,
$$

where $n$ is a non-negative integer and $\hbar=1.05 \times 10^{-34} \mathrm{~J} \cdot \mathrm{~s}$ is Planck's constant.

## 2. The Quantum String

A harmonic oscillator is a system that vibrates back and forth at some frequency $\omega$, such as a mass on a spring.


0

In classical mechanics, the energy of an oscillator can be any non-negative number.

In quantum mechanics, the only permitted energies are of the form

$$
E_{n}=n \hbar \omega,
$$

where $n$ is a non-negative integer and $\hbar=1.05 \times 10^{-34} \mathrm{~J} \cdot \mathrm{~s}$ is Planck's constant.

Terminology: An oscillator with energy $E_{n}$ "is in state $n$ " or "has $n$ excitations at frequency $\omega$."

A classical string has some fundamental frequency $\omega$, set by the requirement that the ends of the string are fixed.

A classical string has some fundamental frequency $\omega$, set by the requirement that the ends of the string are fixed.


The string's allowed frequencies of vibration are positive integer multiples of this fundamental frequency:

$$
\omega, 2 \omega, 3 \omega, \ldots
$$

So we can think of a string as being made up of an infinite collection of harmonic oscillators, with frequencies $\omega, 2 \omega, 3 \omega, \ldots$.

So we can think of a string as being made up of an infinite collection of harmonic oscillators, with frequencies $\omega, 2 \omega, 3 \omega, \ldots$.

Any pattern of vibration can be decomposed as a linear combination of these basic modes of vibration,

$$
y(x)=a_{1} \sin \omega x+a_{2} \sin 2 \omega x+a_{3} \sin 3 \omega x+\cdots
$$



A quantum string is a collection of oscillators with frequencies $\ell \omega$, where $\ell$ is a positive integer.

A quantum string is a collection of oscillators with frequencies $\ell \omega$, where $\ell$ is a positive integer.

Each oscillator has a discrete series of allowed energies

$$
E_{\ell, n_{\ell}}=n_{\ell} \hbar(\ell \omega) . \quad\left(n_{\ell}=0,1,2, \ldots\right)
$$

A quantum string is a collection of oscillators with frequencies $\ell \omega$, where $\ell$ is a positive integer.

Each oscillator has a discrete series of allowed energies

$$
E_{\ell, n_{\ell}}=n_{\ell} \hbar(\ell \omega) . \quad\left(n_{\ell}=0,1,2, \ldots\right)
$$

Therefore, the state of a quantum string is specified by a sequence $\psi$ of non-negative integers,

$$
\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)
$$

where $n_{\ell}$ is the state number of the oscillator with frequency $\ell \omega$.

A quantum string is a collection of oscillators with frequencies $\ell \omega$, where $\ell$ is a positive integer.

Each oscillator has a discrete series of allowed energies

$$
E_{\ell, n_{\ell}}=n_{\ell} \hbar(\ell \omega) . \quad\left(n_{\ell}=0,1,2, \ldots\right)
$$

Therefore, the state of a quantum string is specified by a sequence $\psi$ of non-negative integers,

$$
\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)
$$

where $n_{\ell}$ is the state number of the oscillator with frequency $\ell \omega$.
The total energy of a string in state $\psi$ is the sum of the energies in each of the individual oscillators,

$$
E=n_{1} \hbar \omega+n_{2} \hbar(2 \omega)+n_{3} \hbar(3 \omega)+\cdots \equiv N \hbar \omega,
$$

where

$$
N=n_{1}+2 n_{2}+3 n_{3}+\cdots .
$$

There is a correspondence between states $\psi$ of the string and partitions of $N=n_{1}+2 n_{2}+3 n_{3}+\cdots$ :

The state $\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)$ corresponds to the partition
$n_{1}$ summands
$n_{2}$ summands
$n_{3}$ summands
$N=\overbrace{1+1+\cdots+1}+\overbrace{2+2+\cdots+2}+\overbrace{3+3+\cdots+3}+\cdots$.

There is a correspondence between states $\psi$ of the string and partitions of $N=n_{1}+2 n_{2}+3 n_{3}+\cdots$ :

The state $\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)$ corresponds to the partition
$n_{1}$ summands
$N=\overbrace{1+1+\cdots+1}+\overbrace{2+2+\cdots+2}+\overbrace{3+3+\cdots+3}+\cdots$.

So if we can count how many string states there are with energy $N \hbar \omega$, we'll have solved our partition problem!

## 3. Statistical Mechanics

## 3. Statistical Mechanics

In classical physics, if we know the position and velocity of every particle in a system at a given time, then we can calculate the course of that system forever after.

## 3. Statistical Mechanics

In classical physics, if we know the position and velocity of every particle in a system at a given time, then we can calculate the course of that system forever after.

In practice, macroscopic systems contain too many particles to make such a calculation feasible.

## 3. Statistical Mechanics

In classical physics, if we know the position and velocity of every particle in a system at a given time, then we can calculate the course of that system forever after.

In practice, macroscopic systems contain too many particles to make such a calculation feasible.

But what we really care about is not the microscopic behavior of each of the constituent particles, but the macroscopic features of the system-its average energy, its temperature, its pressure.

## 3. Statistical Mechanics

In classical physics, if we know the position and velocity of every particle in a system at a given time, then we can calculate the course of that system forever after.

In practice, macroscopic systems contain too many particles to make such a calculation feasible.

But what we really care about is not the microscopic behavior of each of the constituent particles, but the macroscopic features of the system-its average energy, its temperature, its pressure.

These features depend only on the aggregate properties of the particles, so we can use statistical methods to study them.
A. Entropy

A. Entropy

Consider an isolated physical system in a particular macroscopic state.


## A. Entropy

Consider an isolated physical system in a particular macroscopic state.


There are many microstates-call their number $W$-that give rise to this macrostate.

## A. Entropy

Consider an isolated physical system in a particular macroscopic state.


There are many microstates-call their number $W$-that give rise to this macrostate.

The entropy of the system is defined as

$$
S=k \ln W,
$$

where $k=1.38 \times 10^{-23} \mathrm{~J} / \mathrm{K}$ is Boltzmann's constant.


In writing $S=k \ln W$, we have assumed each microstate is equally probable.

In writing $S=k \ln W$, we have assumed each microstate is equally probable.

In many systems, the microstates that give rise to the same macrostate are not equally probable.

In writing $S=k \ln W$, we have assumed each microstate is equally probable.

In many systems, the microstates that give rise to the same macrostate are not equally probable.

If the probability of microstate $\psi$ is $p_{\psi}$, then we define the system's entropy to be

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

In writing $S=k \ln W$, we have assumed each microstate is equally probable.

In many systems, the microstates that give rise to the same macrostate are not equally probable.

If the probability of microstate $\psi$ is $p_{\psi}$, then we define the system's entropy to be

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Note. For a system with $W$ equally probable microstates, $p_{\psi}=1 / W$ for all $\psi$, so that

$$
S=-k \sum_{\psi=1}^{W} \frac{1}{W} \ln \left(\frac{1}{W}\right)=-k W\left(\frac{1}{W}\right)(-\ln W)=k \ln W
$$

B. The Partition Function
B. The Partition Function

Consider a non-isolated system, at fixed temperature $T$, that is free to exchange energy with its surroundings, so that its energy is not fixed.

Consider a non-isolated system, at fixed temperature $T$, that is free to exchange energy with its surroundings, so that its energy is not fixed.

We can achieve these conditions by keeping our system in contact with a heat bath at temperature $T$.


Consider a non-isolated system, at fixed temperature $T$, that is free to exchange energy with its surroundings, so that its energy is not fixed.

We can achieve these conditions by keeping our system in contact with a heat bath at temperature $T$.


What is the probability $p_{\psi}$ of finding our system in microstate $\psi$ with energy $E_{\psi}$ ?

Imagine an ensemble consisting of many copies of our system in contact with the heat bath.

Imagine an ensemble consisting of many copies of our system in contact with the heat bath.


Let
be the number of copies in state
$\psi_{1}$ with energy
$E_{\psi_{1}}$, $a_{\psi_{2}}$
$a_{\psi_{3}}$
$\psi_{3}$

Let $a_{\psi}$ $a_{\psi_{1}}$ be the number of copies in state $\quad \psi_{1}$ with energy

Then

$$
\frac{a_{\psi_{2}}}{a_{\psi_{1}}}=f\left(E_{\psi_{1}}, E_{\psi_{2}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) .
$$

Let $\quad a_{\psi_{1}}$ be the number of copies in state $\psi_{1}$ with energy
$a_{\psi_{2}}$
$a_{\psi_{3}}$
$E_{\psi_{1}}$, $E_{\psi_{2}}$, $E_{\psi_{3}}$.

Then

$$
\frac{a_{\psi_{2}}}{a_{\psi_{1}}}=f\left(E_{\psi_{1}}, E_{\psi_{2}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) .
$$

Similarly, $\quad \frac{a_{\psi_{3}}}{a_{\psi_{2}}}=f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)$
and

$$
\frac{a_{\psi_{3}}}{a_{\psi_{1}}}=f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)
$$

Let $\quad a_{\psi_{1}}$ be the number of copies in state $\quad \psi_{1}$ with energy
$E_{\psi_{1}}$, $a_{\psi_{2}}$
$a_{\psi_{3}}$ $\psi_{2}$
$\psi_{3}$

Then

$$
\frac{a_{\psi_{2}}}{a_{\psi_{1}}}=f\left(E_{\psi_{1}}, E_{\psi_{2}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right)
$$

Similarly, $\quad \frac{a_{\psi_{3}}}{a_{\psi_{2}}}=f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)$
and

$$
\frac{a_{\psi_{3}}}{a_{\psi_{1}}}=f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)
$$

Since

$$
\begin{aligned}
& \quad \frac{a_{\psi_{3}}}{a_{\psi_{1}}}=\frac{a_{\psi_{2}}}{a_{\psi_{1}}} \cdot \frac{a_{\psi_{3}}}{a_{\psi_{2}}} \\
& f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)
\end{aligned}
$$

The only function for which

$$
f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)
$$

holds for all $E_{\psi_{i}}$ is an exponential function.

The only function for which

$$
f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)
$$

holds for all $E_{\psi_{i}}$ is an exponential function.
Therefore, for each state $\psi$,

$$
f\left(E_{\psi}\right) \propto \exp \left(-\beta E_{\psi}\right)
$$

for some constant $\beta$, and likewise

$$
a_{\psi} \propto \exp \left(-\beta E_{\psi}\right)
$$

The only function for which

$$
f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)
$$

holds for all $E_{\psi_{i}}$ is an exponential function.
Therefore, for each state $\psi$,

$$
f\left(E_{\psi}\right) \propto \exp \left(-\beta E_{\psi}\right)
$$

for some constant $\beta$, and likewise

$$
a_{\psi} \propto \exp \left(-\beta E_{\psi}\right)
$$

But $a_{\psi} \propto p_{\psi}$, the probability of finding the system in state $\psi$.

The only function for which

$$
f\left(E_{\psi_{3}}-E_{\psi_{1}}\right)=f\left(E_{\psi_{2}}-E_{\psi_{1}}\right) f\left(E_{\psi_{3}}-E_{\psi_{2}}\right)
$$

holds for all $E_{\psi_{i}}$ is an exponential function.
Therefore, for each state $\psi$,

$$
f\left(E_{\psi}\right) \propto \exp \left(-\beta E_{\psi}\right)
$$

for some constant $\beta$, and likewise

$$
a_{\psi} \propto \exp \left(-\beta E_{\psi}\right)
$$

But $a_{\psi} \propto p_{\psi}$, the probability of finding the system in state $\psi$.
Therefore, $p_{\psi} \propto \exp \left(-\beta E_{\psi}\right)$.

Fact. $\beta=1 / k T$ : that is,

$$
p_{\psi} \propto \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

Fact. $\beta=1 / k T$ : that is,

$$
p_{\psi} \propto \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

Intuition. $k T$ is the characteristic energy scale of the system.
States with energy $\lesssim k T$ are probable; states with energy $\gtrsim k T$ are improbable.

Fact. $\beta=1 / k T$ : that is,

$$
p_{\psi} \propto \exp \left(-\frac{E_{\psi}}{k T}\right) .
$$

Intuition. $k T$ is the characteristic energy scale of the system.
States with energy $\lesssim k T$ are probable; states with energy $\gtrsim k T$ are improbable.

If we call the constant of proportionality $1 / Z$, then

Fact. $\beta=1 / k T$ : that is,

$$
p_{\psi} \propto \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

Intuition. $k T$ is the characteristic energy scale of the system.
States with energy $\lesssim k T$ are probable; states with energy $\gtrsim k T$ are improbable.

If we call the constant of proportionality $1 / Z$, then

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

Fact. $\beta=1 / k T$ : that is,

$$
p_{\psi} \propto \exp \left(-\frac{E_{\psi}}{k T}\right) .
$$

Intuition. $k T$ is the characteristic energy scale of the system.
States with energy $\lesssim k T$ are probable; states with energy $\gtrsim k T$ are improbable.

If we call the constant of proportionality $1 / Z$, then

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

This probability distribution is called the Boltzmann distribution.

In the expression

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right),
$$

we can determine $Z$ by noting that our system must be in some microstate, so

$$
\sum_{\psi} p_{\psi}=\frac{1}{Z} \sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)=1
$$

In the expression

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right),
$$

we can determine $Z$ by noting that our system must be in some microstate, so

$$
\sum_{\psi} p_{\psi}=\frac{1}{Z} \sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)=1
$$

So

$$
Z=\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

In the expression

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right),
$$

we can determine $Z$ by noting that our system must be in some microstate, so

$$
\sum_{\psi} p_{\psi}=\frac{1}{Z} \sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)=1
$$

So

$$
Z=\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

The factor $Z$ is called the partition function. It enables us to compute any macroscopic property of our system.
C. Average Energy, Entropy, and Free Energy

## C. Average Energy, Entropy, and Free Energy

We can now compute various physically important macroscopic quantities, starting from

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

(probability of microstate $\psi$ )
and

$$
Z=\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

(the partition function)

## C. Average Energy, Entropy, and Free Energy

We can now compute various physically important macroscopic quantities, starting from

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right) \quad \text { (probability of microstate } \psi \text { ) }
$$

and

$$
Z=\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

(the partition function)

The system's average energy $\langle E\rangle$ is the expected value of $E_{\psi}$,

$$
\langle E\rangle=\sum_{\psi} p_{\psi} E_{\psi}=\frac{1}{Z} \sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right) E_{\psi} .
$$

Let us compute the entropy

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Let us compute the entropy

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Since

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right),
$$

we find

Let us compute the entropy

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Since

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

we find

$$
S=-k \sum_{\psi} p_{\psi}\left(-\frac{E_{\psi}}{k T}-\ln Z\right)=\frac{\langle E\rangle}{T}+k \ln Z
$$

Let us compute the entropy

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Since

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

we find

$$
S=-k \sum_{\psi} p_{\psi}\left(-\frac{E_{\psi}}{k T}-\ln Z\right)=\frac{\langle E\rangle}{T}+k \ln Z
$$

Rearranging, we get

$$
\langle E\rangle-T S=-k T \ln Z \equiv F
$$

which defines the free energy $F$.

Let us compute the entropy

$$
S=-k \sum_{\psi} p_{\psi} \ln p_{\psi}
$$

Since

$$
p_{\psi}=\frac{1}{Z} \exp \left(-\frac{E_{\psi}}{k T}\right),
$$

we find

$$
S=-k \sum_{\psi} p_{\psi}\left(-\frac{E_{\psi}}{k T}-\ln Z\right)=\frac{\langle E\rangle}{T}+k \ln Z
$$

Rearranging, we get

$$
\langle E\rangle-T S=-k T \ln Z \equiv F,
$$

which defines the free energy $F$.
Fact. $S=-\frac{d F}{d T}$
4. The Main Calculation

## 4. The Main Calculation

A quantum string in state $\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)$ with
$n_{1}$ excitations at frequency $\omega$, $n_{2}$ excitations at frequency $2 \omega$, $n_{3}$ excitations at frequency $3 \omega$, etc.,
has energy

$$
E_{\psi}=\hbar \omega N
$$

where

$$
N=n_{1}+2 n_{2}+3 n_{3}+\cdots=\sum_{\ell=1}^{\infty} \ell n_{\ell} .
$$

## 4. The Main Calculation

A quantum string in state $\psi=\left(n_{1}, n_{2}, n_{3}, \ldots\right)$ with
$n_{1}$ excitations at frequency $\omega$, $n_{2}$ excitations at frequency $2 \omega$, $n_{3}$ excitations at frequency $3 \omega$, etc.,
has energy

$$
E_{\psi}=\hbar \omega N
$$

where

$$
N=n_{1}+2 n_{2}+3 n_{3}+\cdots=\sum_{\ell=1}^{\infty} \ell n_{\ell} .
$$

This string state corresponds to the partition

$$
N=\overbrace{1+1+\cdots+1}^{n_{1} \text { summands }}+\overbrace{2+2+\cdots+2}^{n_{2} \text { summands }}+\overbrace{3+3+\cdots+3}^{n_{3} \text { summands }}+\cdots .
$$

Therefore, counting the number of partitions of $N$ is equivalent to counting the number of string states with energy $N \hbar \omega$.

Therefore, counting the number of partitions of $N$ is equivalent to counting the number of string states with energy $N \hbar \omega$.

But ( $k$ times the logarithm of) the number of the string states is the entropy.

Therefore, counting the number of partitions of $N$ is equivalent to counting the number of string states with energy $N \hbar \omega$.

But ( $k$ times the logarithm of) the number of the string states is the entropy.

Strategy: Use statistical mechanics to calculate the entropy of the string, and thereby to count the number of partitions of $N$.

Everything begins with the partition function

$$
Z=\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right)
$$

Everything begins with the partition function

$$
\begin{aligned}
Z & =\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right) \\
& =\sum_{n_{1}=0}^{\infty} \sum_{n_{2}=0}^{\infty} \cdots \exp \left(-\frac{\hbar \omega}{k T}\left(n_{1}+2 n_{2}+\cdots\right)\right)
\end{aligned}
$$

Everything begins with the partition function

$$
\begin{aligned}
Z & =\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right) \\
& =\sum_{n_{1}=0}^{\infty} \sum_{n_{2}=0}^{\infty} \cdots \exp \left(-\frac{\hbar \omega}{k T}\left(n_{1}+2 n_{2}+\cdots\right)\right) \\
& =\left[\sum_{n_{1}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot n_{1}\right)\right]\left[\sum_{n_{2}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot 2 n_{2}\right)\right] \cdots \\
& =\prod_{\ell=1}^{\infty}\left[\sum_{n_{\ell}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot \ell n_{\ell}\right)\right]
\end{aligned}
$$

Everything begins with the partition function

$$
\begin{aligned}
Z & =\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right) \\
& =\sum_{n_{1}=0}^{\infty} \sum_{n_{2}=0}^{\infty} \cdots \exp \left(-\frac{\hbar \omega}{k T}\left(n_{1}+2 n_{2}+\cdots\right)\right) \\
& =\left[\sum_{n_{1}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot n_{1}\right)\right]\left[\sum_{n_{2}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot 2 n_{2}\right)\right] \cdots \\
& =\prod_{\ell=1}^{\infty}\left[\sum_{n_{\ell}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot \ell n_{\ell}\right)\right] \\
& =\prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1}
\end{aligned}
$$

Everything begins with the partition function

$$
\begin{aligned}
Z & =\sum_{\psi} \exp \left(-\frac{E_{\psi}}{k T}\right) \\
& =\sum_{n_{1}=0}^{\infty} \sum_{n_{2}=0}^{\infty} \cdots \exp \left(-\frac{\hbar \omega}{k T}\left(n_{1}+2 n_{2}+\cdots\right)\right) \\
& =\left[\sum_{n_{1}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot n_{1}\right)\right]\left[\sum_{n_{2}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot 2 n_{2}\right)\right] \cdots \\
& =\prod_{\ell=1}^{\infty}\left[\sum_{n_{\ell}=0}^{\infty} \exp \left(-\frac{\hbar \omega}{k T} \cdot \ell n_{\ell}\right)\right] \\
& =\prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1}
\end{aligned}
$$

Note. If we define $x=e^{-\frac{\hbar \omega}{k T}}$, then $Z=\prod_{\ell=1}^{\infty} \frac{1}{1-x^{\ell}}=\sum_{N=0}^{\infty} p(N) x^{N}$ is exactly the generating function for partitions!

The free energy is

$$
F=-k T \ln Z
$$

The free energy is

$$
\begin{aligned}
F & =-k T \ln Z \\
& =k T \ln \prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1}
\end{aligned}
$$

The free energy is

$$
\begin{aligned}
F & =-k T \ln Z \\
& =k T \ln \prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1} \\
& =k T \sum_{\ell=1}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right]
\end{aligned}
$$

The free energy is

$$
\begin{aligned}
F & =-k T \ln Z \\
& =k T \ln \prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1} \\
& =k T \sum_{\ell=1}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right]
\end{aligned}
$$

We now take the limit $T \rightarrow \infty$, or equivalently, set $\frac{\hbar \omega}{k T} \ll 1$.
We'll see later that this is tantamount to assuming $N \gg 1$.

The free energy is

$$
\begin{aligned}
F & =-k T \ln Z \\
& =k T \ln \prod_{\ell=1}^{\infty}\left[1-\exp \left(-\frac{\hbar \omega}{k T} \cdot \ell\right)\right]^{-1} \\
& =k T \sum_{\ell=1}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right]
\end{aligned}
$$

We now take the limit $T \rightarrow \infty$, or equivalently, set $\frac{\hbar \omega}{k T} \ll 1$. We'll see later that this is tantamount to assuming $N \gg 1$. In this limit, the summand $\ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right]$ changes slowly with $\ell$.


So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

Changing variables to $q=\frac{\hbar \omega l}{k T}$,

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

Changing variables to $q=\frac{\hbar \omega l}{k T}$,

$$
F \approx \frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty} \ln \left(1-e^{-q}\right) d q
$$

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

Changing variables to $q=\frac{\hbar \omega \ell}{k T}$,

$$
F \approx \frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty} \ln \left(1-e^{-q}\right) d q
$$

Taylor-expanding the logarithm via

$$
\ln (1-y)=-\left(y+\frac{y^{2}}{2}+\frac{y^{3}}{3}+\cdots\right)
$$

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell .
$$

Changing variables to $q=\frac{\hbar \omega \ell}{k T}$,

$$
F \approx \frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty} \ln \left(1-e^{-q}\right) d q
$$

Taylor-expanding the logarithm via

$$
\begin{gathered}
\ln (1-y)=-\left(y+\frac{y^{2}}{2}+\frac{y^{3}}{3}+\cdots\right) \\
F \approx-\frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty}\left(e^{-q}+\frac{1}{2} e^{-2 q}+\frac{1}{3} e^{-3 q}+\cdots\right) d q
\end{gathered}
$$

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

Changing variables to $q=\frac{\hbar \omega \ell}{k T}$,

$$
F \approx \frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty} \ln \left(1-e^{-q}\right) d q
$$

Taylor-expanding the logarithm via

$$
\begin{gathered}
\ln (1-y)=-\left(y+\frac{y^{2}}{2}+\frac{y^{3}}{3}+\cdots\right) \\
F \approx-\frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty}\left(e^{-q}+\frac{1}{2} e^{-2 q}+\frac{1}{3} e^{-3 q}+\cdots\right) d q \\
=-\frac{(k T)^{2}}{\hbar \omega}\left(1+\frac{1}{2^{2}}+\frac{1}{3^{2}}+\cdots\right)
\end{gathered}
$$

So we may approximate the sum by an integral,

$$
F \approx k T \int_{0}^{\infty} \ln \left[1-\exp \left(-\frac{\hbar \omega \ell}{k T}\right)\right] d \ell
$$

Changing variables to $q=\frac{\hbar \omega \ell}{k T}$,

$$
F \approx \frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty} \ln \left(1-e^{-q}\right) d q
$$

Taylor-expanding the logarithm via

$$
\begin{aligned}
& \ln (1-y)=-\left(y+\frac{y^{2}}{2}+\frac{y^{3}}{3}+\cdots\right) \\
F \approx & -\frac{(k T)^{2}}{\hbar \omega} \int_{0}^{\infty}\left(e^{-q}+\frac{1}{2} e^{-2 q}+\frac{1}{3} e^{-3 q}+\cdots\right) d q \\
= & -\frac{(k T)^{2}}{\hbar \omega}\left(1+\frac{1}{2^{2}}+\frac{1}{3^{2}}+\cdots\right) \\
= & -\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6} .
\end{aligned}
$$

The entropy is

$$
S=-\frac{d F}{d T}=-\frac{d}{d T}\left(-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}\right)=\frac{k^{2} T}{\hbar \omega} \cdot \frac{\pi^{2}}{3} .
$$

The entropy is

$$
S=-\frac{d F}{d T}=-\frac{d}{d T}\left(-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}\right)=\frac{k^{2} T}{\hbar \omega} \cdot \frac{\pi^{2}}{3} .
$$

The average energy is

$$
\begin{aligned}
\langle E\rangle & =F+T S \\
& =-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}+\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{3} \\
& =\frac{\pi^{2}}{6}\left(\frac{k T}{\hbar \omega}\right)^{2} \cdot \hbar \omega
\end{aligned}
$$

The entropy is

$$
S=-\frac{d F}{d T}=-\frac{d}{d T}\left(-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}\right)=\frac{k^{2} T}{\hbar \omega} \cdot \frac{\pi^{2}}{3} .
$$

The average energy is

$$
\begin{aligned}
\langle E\rangle & =F+T S \\
& =-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}+\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{3} \\
& =\frac{\pi^{2}}{6}\left(\frac{k T}{\hbar \omega}\right)^{2} \cdot \hbar \omega
\end{aligned}
$$

Since $E=N \hbar \omega$, we can now see that the large- $T$ limit is the same as the large- $N$ limit.

The entropy is

$$
S=-\frac{d F}{d T}=-\frac{d}{d T}\left(-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}\right)=\frac{k^{2} T}{\hbar \omega} \cdot \frac{\pi^{2}}{3} .
$$

The average energy is

$$
\begin{aligned}
\langle E\rangle & =F+T S \\
& =-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}+\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{3} \\
& =\frac{\pi^{2}}{6}\left(\frac{k T}{\hbar \omega}\right)^{2} \cdot \hbar \omega
\end{aligned}
$$

Since $E=N \hbar \omega$, we can now see that the large- $T$ limit is the same as the large- $N$ limit.

In this limit, the string is overwhelmingly likely to be in a state near its average energy, so we may freely interchange $E$ and $\langle E\rangle$.

The entropy is

$$
S=-\frac{d F}{d T}=-\frac{d}{d T}\left(-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}\right)=\frac{k^{2} T}{\hbar \omega} \cdot \frac{\pi^{2}}{3} .
$$

The average energy is

$$
\begin{aligned}
\langle E\rangle & =F+T S \\
& =-\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{6}+\frac{(k T)^{2}}{\hbar \omega} \cdot \frac{\pi^{2}}{3} \\
& =\frac{\pi^{2}}{6}\left(\frac{k T}{\hbar \omega}\right)^{2} \cdot \hbar \omega
\end{aligned}
$$

Since $E=N \hbar \omega$, we can now see that the large- $T$ limit is the same as the large- $N$ limit.

In this limit, the string is overwhelmingly likely to be in a state near its average energy, so we may freely interchange $E$ and $\langle E\rangle$.

Written as a function of the energy, the entropy is

$$
S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}} .
$$

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

$$
S=k \cdot 2 \pi \sqrt{\frac{N}{6}} .
$$

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

$$
S=k \cdot 2 \pi \sqrt{\frac{N}{6}} .
$$

On the other hand, $S=k \ln W$, where $W$ is the number of states of the string;

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

$$
S=k \cdot 2 \pi \sqrt{\frac{N}{6}} .
$$

On the other hand, $S=k \ln W$, where $W$ is the number of states of the string;
and $W=p(N)$;

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

$$
S=k \cdot 2 \pi \sqrt{\frac{N}{6}} .
$$

On the other hand, $S=k \ln W$, where $W$ is the number of states of the string;
and $W=p(N)$;

SO

$$
k \ln p(N)=k \cdot 2 \pi \sqrt{\frac{N}{6}}
$$

Substituting $E=N \hbar \omega$ into $S(E)=k \pi \sqrt{\frac{2 E}{3 \hbar \omega}}$, we get

$$
S=k \cdot 2 \pi \sqrt{\frac{N}{6}} .
$$

On the other hand, $S=k \ln W$, where $W$ is the number of states of the string;
and $W=p(N)$;
so

$$
k \ln p(N)=k \cdot 2 \pi \sqrt{\frac{N}{6}}
$$

or

$$
\ln p(N)=2 \pi \sqrt{\frac{N}{6}}, \quad \text { Q.E.D.! }
$$

