A Concise History of the Black-body Radiation Problem

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The way the topic of black-body radiation is presented in standard textbooks (i.e. from Rayleigh-Jeans to Max Planck) does not follow the actual historical timeline of the understanding of the black-body radiation problem. Authors believe that a presentation which follows an actual timeline of the ideas (although not a logical presentation of the field) would be of interest not only from the history of science perspective but also from a pedagogical perspective. Therefore, we attempt a concise history of this very interesting field of science.

Introduction

Generally, we understand Planck’s distribution function for the black-body radiation as a modification of Rayleigh-Jeans law using the canonical ensemble of quantized energy. However, as the foundation theory of quantum mechanics, it is worth understanding the actual development of the ideas starting from the pioneering work of Gustav Kirchhoff in this field. Kirchhoff’s theoretical analysis gave scientists a way to characterize the black-body spectrum. Josef Stefan’s empirical deduction of the $T^4$ law initiated further theoretical investigations. Ludwig Boltzmann’s thermodynamic derivation of Josef Stefan’s law verifies the idea of radiation pressure. Next, the fascinating derivation of Wilhelm Wien’s scaling law (but lesser known) is discussed, which led Max Planck to conclude that the entropy of an oscillator is a function of the ratio of the average energy and frequency of an oscillator. Next, we review Planck’s original derivation of his distribution function to show the historical importance of Kirchhoff’s law, Stefan-Boltzmann’s $T^4$ law, and Wien’s scaling law.
Finally, the classical distribution function of Lord Rayleigh and James Jeans is discussed. In the following paragraphs, we briefly sketch this historical development. In the following sections, a more detailed analysis is given.

The theory of radiative heat exchange was first initiated by Genevan physicist Pierre Prevost in 1791 \[20, 4\]. He defined the thermal equilibrium in the context of radiative heat transfer and explained that each body radiates and receives heat independently of the presence of the other bodies. The thermal radiation is in the infrared frequency range when the temperature of the body is equal to room temperature, so we can not see that. As we increase the temperature, the body starts glowing from red to white in colour. In 1830, Leopoldo Nobili and Macedonio Melloni made a thermopile device, which converts thermal energy into electrical current. In 1831, they made the first radiometer using thermopile and galvanometer \[23\]. Their device showed that the amount of radiation emitted from different surfaces at the same temperature is not the same. In 1847 John William Draper observed that \(525^\circ\)C is an average temperature where emission radiation becomes visible, and it is known as Draper point \[7\]. In 1858, Scottish physicist Balfour Stewart experimentally measured the radiation using thermopile and compared thermal emission and absorptive power of different materials with lamp-black \[35, 33, 31\]. Stewart wrote, "Lamp-black which absorbs all the rays that fall upon it, therefore, possesses the greatest possible absorbing power and posses the greatest possible radiating power".

The notion of a black-body was first defined in a concrete way by Gustav Kirchhoff: A body which reflects no light at all nor allows light to pass. No ideal black-body exists in nature. Lamp-black and Platinum-black are good approximations. Kirchhoff theoretically explained that the emission spectrum (energy density) is independent of the shape, size and material of the black-body, and it is the only function of radiation wavelength and temperature. Thus, if two different black bodies are in equilibrium, then their
radiation field is identical.

John Draper, John Tyndall and many other physicists have studied the temperature dependence of energy density of thermal radiation. Draper also plotted the data on total energy emission vs temperature in 1847 [7]. Josef Stefan empirically described the dependence of temperature on the total energy emitted by a black-body. Then five years later, Ludwig Boltzmann theoretically proved Stefan’s law using purely thermodynamic arguments. This law is known as Stefan-Boltzmann $T^4$ law.

Wilhelm Wien is known for finding the relation between temperature and wavelength, where the energy density is maximum. Apart from this Wien’s displacement law, he was the first who gave the simplified form of the Kirchhoff universal function using rigorous thermodynamical arguments in 1893. He also gave the empirical energy distribution for the black-body in 1896, but it turned out to be just an approximation.

In June 1900, Lord Rayleigh considered black-body radiation as electromagnetic standing wave vibrations in cavity enclosure, and he used the equipartition theorem of statistical mechanics. He found that the energy distribution function of black-body radiation is proportional to the $T \lambda^{-4}$. Rayleigh and English physicist James Jeans found a complete form of the energy distribution function five years later, but it disagreed with the experimental observations.

German physicist Max Planck has been working on the black-body problem for more than five years. He assumed that cavity walls are made of a collection of oscillators of electric dipoles. He published his exact solution for the black-body problem in two papers (October 1900 and December 1900), where he introduced the idea of energy quantization. Planck’s implication on the quantum of energy and historical debates on it is discussed by Draper’s data was on the Fahrenheit scale. The absolute thermometric scale (Kelvin scale) was invented by William Lord Kelvin in 1848.
In the subsequent section, we consider the contributions of the leading investigators in this field one by one.

1. Enter Gustav Kirchhoff

In 1859, German physicist Gustav Robert Kirchhoff gave the first theoretical argument on the back-body radiation problem. Along with Robert Bunsen, Kirchhoff studied the coincidence of the emission lines of sodium with particular absorption lines in the solar spectrum using a prism. From his experiments, he considered that at thermal equilibrium, the ratio of emissive power to absorptive power is the same for all bodies and asked the question: does this the case for each wavelength separately? He concludes that for any thermally radiating body, the ratio of emissive power $e$ and the absorptive power $a$ is a universal function that only depends upon the wavelength and temperature [12, 13].

Kirchhoff considered two thin plates $C$ and $c$ of emissive power $E$ and $e$ and absorptive power $A$ and $a$ respectively [12, 33]. One of the plate surfaces is a perfectly reflecting mirror, and the plates are arranged parallel to each other, as shown in Figure (2). Plate $c$
is made of special material that only absorbs and emits the wavelength radiation $\Lambda$. All other wavelengths pass through. Kirchhoff considered a quantity of radiation $E$ of wavelength $\Lambda$ emitted by $C$. When it comes to plate $c$, it absorbs $aE$ radiation and throws $(1 - a)E$ radiation back. When this $(1 - a)E$ radiation reach to plate $C$, it absorbs $A(1 - a)E$ radiation and throws $(1 - A)(1 - a)E$ then again plate $c$ absorbs $a(1 - A)(1 - a)E$ radiation and throws $(1 - a)(1 - A)(1 - a)E$ radiation and so on. This infinite series of bounces back and forth of radiation between two mirrors gives the total absorption by plate $c$ as

$$\frac{aE}{1 - (1 - A)(1 - a)}.$$  \hspace{1cm} (1)

Similarly, if the plate $c$ emits the quantity of radiation $e$ of wavelength $\Lambda$, then the plate $c$ absorbs radiation of quantity

$$\frac{a(1 - A)e}{1 - (1 - A)(1 - a)}. \hspace{1cm} (2)$$

So at thermal equilibrium, total absorption by the plate $c$ should be equal to $e$.

$$e = \frac{aE}{1 - (1 - A)(1 - a)} + \frac{a(1 - A)e}{1 - (1 - A)(1 - a)}.$$ \hspace{1cm} (3)

By rearranging the terms, we get

$$\frac{e}{a} = \frac{E}{A}.$$ \hspace{1cm} (4)
and this is true for any plate $C'$ with emissive power $E'$ and absorptive power $A'$.

\[
\frac{e}{a} = \frac{E'}{A'} \quad (5)
\]

This shows that for the same temperature and wavelength, the ratio of emissive power $e$ and the absorptive power $a$ is the same for all bodies. i.e. it is universal function $f(\lambda, T)$.

\[
\frac{e}{a} = f(\lambda, T) \quad (6)
\]

After one month, in Jan-1860, Kirchhoff published a second paper where he proved this theorem in general context[13, 11, 33, 32]. In the second paper, he introduced the term black-body, which absorbs all of the radiation falling upon it, so its absorptive power is one. Then the emission power of the black-body is a universal function of wavelength and temperature. Collecting contributions from the whole spectrum (integrating over all wavelengths), we can see that the total energy emitted per unit volume by the black-body is the only function of its temperature, which is given by the Stefan-Boltzmann $T^4$ law.

2. Enter Josef Stefan

Figure 3. Josef Stefan (1835-1893) [Photo: Wikipedia Commons]

In 1864, Irish physicist John Tyndall did measurements of the infrared emission by the platinum filament, and the corresponding
John Tyndall was a pioneering experimental physicist in the field of atmospheric physics. He is known for discovering the greenhouse effect and the Tyndall effect.

3. Enter Ludwig Boltzmann

In 1884, Stefan’s formal doctoral student Ludwig Boltzmann published a thermodynamic derivation of Stephen’s empirical law. Using the idea of Adolfo Bartoli on radiation pressure \([1]\), Boltzmann considered radiation particles as the ideal gas in the piston-cylinder system.

The pressure exerted on the piston by the radiation particles when the piston moves slowly is

\[
P = \frac{1}{3}\rho_T \tag{7}
\]

where \(\rho_T\) is the total energy density (Appendix A).
Putting total internal energy $U = \rho_T V$ and radiation pressure into the fundamental thermodynamics relation $dU = TdS - PdV$, we get

$$TdS = \frac{4}{3}\rho_T dV + V\frac{d\rho_T}{dT}dT$$  \hspace{1cm} (8)

We know $S \equiv S(T, V)$ so

$$dS = \frac{\partial S}{\partial T}dT + \frac{\partial S}{\partial V}dV$$  \hspace{1cm} (9)

By comparing equation (8) and (9), we get

$$\frac{\partial S}{\partial T} = \frac{V}{T} \frac{d\rho_T}{dT}, \quad \frac{\partial S}{\partial V} = \frac{4}{3} \frac{\rho_T}{T}.$$  \hspace{1cm} (10)

So we have

$$\frac{\partial^2 S}{\partial V \partial T} = \frac{1}{T} \frac{d\rho_T}{dT}, \quad \frac{\partial^2 S}{\partial T \partial V} = -\frac{4}{3} \frac{\rho_T}{T^2} + \frac{4}{3} \frac{1}{T} \frac{d\rho_T}{dT}.$$  \hspace{1cm} (11)

By symmetry of partial derivative, equating above terms, we get

$$\frac{d\rho_T}{dT} = a\frac{\rho_T}{T}.$$  \hspace{1cm} (12)

Integrating the above equation, we get

$$\rho_T = aT^4.$$  \hspace{1cm} (13)

Using Lambert’s relation $\rho_T = \frac{4}{5}E$ (where $E$ is the total emissive power over the hemisphere) we get the well-known Stefan-Boltzmann law $E = \sigma T^4$. The constant $\sigma = \frac{ac}{T^4}$ is known as Stefan-Boltzmann constant. The exact form of a constant can not be determined from purely classical mechanics.

4. Enter Wilhelm Wien

The German physicist Wilhelm Franz Wien’s work has been the most important work in studying black-body radiation. In 1893, Wien used thermodynamical arguments and the Doppler effect and derived important relation between the energy density of radiation with frequency and temperature.

$$\rho(v) = v^3f(v/T)$$  \hspace{1cm} (14)
This should be called Wien’s scaling law. Using this, he also derived that the product of the wavelength where the emission spectrum of black-body has maximum and temperature is constant, known as “Wein’s displacement law”.

Now we present a concise derivation of Wien’s scaling law which is hard to find in the books. As before, suppose radiation is enclosed in a piston-cylinder system and is in thermodynamical equilibrium at temperature T.

Suppose that the inner surface of the piston is perfectly reflected. Radiation exerts radiation pressure on the piston. A piston is allowed to move out quasi-statically, and this process is made adiabatic, so no external heat is allowed to enter into the cylinder.

Suppose at time $t = 0$ a radiation beam coming from point $S$ is incident on a mirror at angle $\theta$ with the normal to the piston. The observer at $O$ receives the piston reflected beam from the moving piston.

\footnote{Only Max Born discussed this briefly in his book "Atomic Physics".}

Figure 5. Wilhelm Wien (1864-1928) [Photo: Wikipedia Commons]

Figure 6. The radiation inside the piston-cylinder system. $S$ denotes a radiation source, $O$ denotes an observer, and $\mu$ is the velocity of the piston.
mirror. Let \( u \) be a velocity of the piston. Now, the Doppler effect with velocity \( u \cos \theta \) since it is relevant to observer \( O \) gives the frequency observed by \( O \).

\[
\nu' = \nu - \frac{2u}{c} \nu \cos \theta
\]  

(15)

Let us denote the \( I(\nu)d\nu \) as intensity of radiation whose frequency is in range \( \nu \) and \( \nu + d\nu \). Then the energy in the range \( \nu \) and \( \nu + d\nu \) falling on the patch of area \( dA \) on the piston mirror in time \( dt \) is

\[
I(\nu)d\nu dA \frac{dt}{c} = I(\nu) \cos \theta dA d\nu
\]  

(16)

If \( I(\nu')d\nu' \) is the intensity of the reflected radiation whose frequency lies in \( \nu' \) to \( \nu' + d\nu' \), the reflected energy from patch in time \( dt \) is

\[
I'(\nu')d\nu' dA \frac{dt}{c} = I'(\nu') \cos \theta dA d\nu'
\]  

(17)

then loss of the radiation energy will be

\[
-(I(\nu)d\nu - I'(\nu')d\nu') \cos \theta dA dt.
\]  

(18)

This loss appears in the form of work done on the piston by radiation. The pressure exerted by radiation on the piston is

\[
dP = 2\rho(\nu) \cos^2 \theta \ d\nu.
\]  

(19)

So, the work done will be

\[
2\rho(\nu) \cos^2 \theta \ d\nu \ dA \ u dt.
\]  

(20)

Equating this with the equation (18) and using \( I(\nu) = \rho(\nu)c \) we get

\[
I'(\nu')d\nu' = I(\nu) \left(1 + \frac{2u \cos \theta}{c}\right) d\nu
\]  

(21)

Again using the \( I(\nu) = \rho(\nu)c \) and integrating both side, we get

\[
\rho' = \rho \cdot \left(1 + \frac{2u \cos \theta}{c}\right)
\]  

(22)

where \( \rho' \) and \( \rho \) denote the integrated energy density. So, after reflection, energy density is increased by a factor \( \left(1 + \frac{2u \cos \theta}{c}\right) \).
To calculate the total energy change in time $dt$ of radiation whose frequency is in the range $\nu$ to $\nu + d\nu$, we need to take an angular average of the difference in final and initial state energy, which means

$$\Delta E = \int_{0}^{2\pi} \int_{0}^{\pi/2} c dt \ dA \ \cos\theta \ \frac{(\sin\theta d\theta d\phi)}{4\pi} (\rho'(\nu')d\nu' - \rho(\nu)d\nu)$$

(23)

All the radiation whose frequency is in the range $\nu$ to $\nu + d\nu$, after reflection that will be in range $\nu'$ to $\nu' + d\nu'$, where $\nu'$ is given by equation (15). So the term $\rho'(\nu')d\nu'$ is replaced by

$$\left(1 + \frac{2ucos\theta}{c}\right) \rho\left(\nu - \frac{2ucos\theta}{c}\nu\right) \left(1 - \frac{2ucos\theta}{c}\nu\right) d\nu.$$  

(24)

Writing Taylor expansion,

$$\rho\left(\nu - \frac{2ucos\theta}{c}\nu\right) = \rho(\nu) - \frac{2ucos\theta}{c}\nu \frac{\partial \rho}{\partial \nu} - \cdots$$

(25)

and neglecting terms of $\frac{u^2}{c^2}$ and higher in equation (24), and putting into equation (23), we get

$$\Delta E = \frac{cdt \ dA}{2} \int_{0}^{\pi/2} d\phi \sin\theta \cos\theta \left(-2 \frac{ucos\theta}{c}\nu \frac{\partial \rho}{\partial \nu}\right)$$

(26)

Now using the $dV = u \ dt \ dA$ and $\int_{0}^{\pi/2} d\theta \sin\theta \cos^2\theta = \frac{1}{3}$, we get

$$\Delta E = -dV \frac{\nu}{3} \frac{\partial \rho(\nu)}{\partial \nu} d\nu$$

(27)

The negative sign shows that there is a net reduction in energy which appears as work done by the piston. The reduction of the energy for the frequency range $\nu$ to $\nu + d\nu$ can be written as $-d(\rho(\nu)V)d\nu$. So we have

$$d(\rho(\nu)V)d\nu = dV \frac{\nu}{3} \frac{\partial \rho(\nu)}{\partial \nu} d\nu$$

(28)

since the process of moving the piston out by a very small amount $vdt$ is adiabatic, we can take $\rho(\nu)$ as the only function of volume since the energy density reduces with an increase in volume.

$$\rho(\nu) + V \frac{\partial \rho(\nu)}{\partial V} = \frac{\nu}{3} \frac{\partial \rho(\nu)}{\partial \nu}$$

(29)
It is easy to check that function $\rho(\nu) = \nu^3 \phi(\nu^3 V)$ is a solution of this equation.

Now, using the equation (8) for adiabatic process ($ds = 0$) we get

$$0 = \frac{V}{T} \frac{\partial \rho}{\partial T} dT + \frac{4}{3} \frac{\rho}{T} dV$$

(30)

Now, using the Stefan-Boltzmann law, we get

$$0 = \frac{V}{T} (4aT^3) dT + \frac{4}{3} \frac{aT^4}{V} dV$$

(31)

So,

$$\frac{dT}{T} = -\frac{1}{3} \frac{dV}{V}$$

(32)

which gives the solution $VT^3$ = constant.

So, we have

$$\rho(\nu) = \nu^3 \phi \left( \frac{\nu^3}{T} \right) \equiv \nu^3 f \left( \frac{\nu}{T} \right)$$

(33)

Using $\rho(\nu) d\nu = \rho(\lambda) d\lambda$ and $\lambda = c/\nu$, we get

$$\rho(\lambda, T) = \frac{c^4}{\lambda^5} f \left( \frac{c}{T\lambda} \right)$$

(34)

Thus Wien’s scaling law simplifies Kirchhoff’s universal function. This suggests that a plot of $\lambda^5 \rho(\lambda, T)$ vs $\lambda T$ gives a single curve regardless of the wavelength of radiation and temperature of the black-body. In other words, knowing a spectrum for a single temperature, a spectrum can be found for any other temperature. Later, other equivalent forms were given by Lord Rayleigh, and Max Ferdinand Thiesen [28].

Let $\rho(\lambda)$ is maximum at $\lambda_m$. Then derivative of $\rho(\lambda)$ with respect to $\lambda$ is zero at $\rho(\lambda)$, So

$$5f \left( \frac{c}{T\lambda_m} \right) + \frac{c}{T\lambda_m} f' \left( \frac{c}{T\lambda_m} \right) = 0$$

(35)

Let $x = \frac{c}{T\lambda_m}$ and rearranging terms we get

$$\frac{df(x)}{f(x)} = -5 \frac{dx}{x}$$

(36)
The solution to this differential equation is

\[ f(x) x^5 = \text{constant} \equiv k \]  

(37)

\[ f\left(\frac{c}{T\lambda_m}\right) = k \frac{(\lambda_m T)^5}{c^5} \]  

Then, \( \rho(\lambda_m) = \frac{kT^5}{c} \)  

(38)

This is not physical solution, so only solution is \( x = \frac{c}{T\lambda_m} \) is constant, i.e.

\[ \lambda_m T = \text{Const.} \]  

(39)

This is known as Wien’s displacement law or generally known as the Wien’s law. Any function which satisfies Wien’s scaling law will obey Wien’s displacement law. The first experimental verification of Wien’s law was given by German physicists Otto Lummer, and Ernst Pringsheim in 1895 [21][18].

In 1896, Wien published a second paper on black-body radiation [38]. He described a black-body spectrum using the Maxwell-Boltzmann distribution of atoms and assumed that the wavelength of radiation is the only function of its velocity (square of the velocity). He wrote energy density as

\[ \rho(\lambda, T) = F(\lambda)e^{-\frac{a}{\lambda T}} \]  

(40)

Now, using the Stefan-Boltzmann law, we have

\[ \int_0^{\infty} F(\lambda)e^{-\frac{a}{\lambda T}} d\lambda \propto T^4 \]  

(41)

Wien defined \( y := \frac{a}{\lambda T} \) and wrote expansion of function \( F(\lambda) \) as

\[ F(\lambda) = F\left(\frac{a}{yT}\right) = c_0 + c_1 \frac{yT}{a} + c_2 \frac{y^2 T^2}{a^2} + \cdots + c_n \frac{y^n T^n}{a^n} + \ldots \]

\[ c_{-1} \frac{a}{yT} + c_{-2} \frac{a^2}{y^2 T^2} + \cdots + c_{-n} \frac{a^n}{y^n T^n} + \ldots \]  

(42)
Putting into integral, we get
\[
\int_0^\infty F(\lambda) e^{-\frac{a}{\lambda T}} d\lambda = \frac{a}{T} \int_0^\infty F\left(\frac{a}{yT}\right) e^{-\frac{y}{y^2}} dy
\]
\[
= \sum_n c_n \frac{T^{n-1}}{y^{n-1}} \int_0^\infty e^{-y^2} y^{n-2} dy
\]
\[
= \sum_n c_n \frac{T^{n-1}}{y^{n-1}} \Gamma(n - 1) \tag{43}
\]
This should be proportional to the $T^4$, so we must have $c_5$ is non-zero, and all others must be zero, which gives $F(\lambda) = \frac{C}{\lambda^5}$ where $C$ is constant. So energy density has a form
\[
\rho(\lambda, T) = \frac{C}{\lambda^5} e^{-\frac{a}{\lambda T}}. \tag{44}
\]
This is known as Wien’s distribution law or Wien’s approximation. This agrees with the equation (34) and gives a quite close distribution to experimental results. In the same month, German physicist Friedrich Paschen found that his experimental data is best fitted to function [24]
\[
\rho(\lambda, T) = \frac{C_P}{\lambda^{5.56}} e^{-\frac{a}{\lambda T}}. \tag{45}
\]
Which is close to Wien’s empirical distribution. Wien did not give a rigorous physical argument for the empirical exponential factor. Many other attempts were made to find Kirchhoff’s universal function, but this was the first closest function that was found. In 1899, Lummer and Pringsheim checked that the Wien’s distribution agrees for small wavelengths, but at a larger wavelength and higher temperature, there is a systematic difference with experimental data.

5. Enter Lord Rayleigh

In June 1900, English mathematician and physicist John William Strutt, known as 3rd Baron Rayleigh, gave another form of universal function in a paper “Remarks upon the law of complete radiation”. Rayleigh was the founder of the theory of sound, published many papers, and wrote two books on it. Using the analogy
of the theory of sound, Rayleigh classically derived that the emission spectrum is proportional to the $T \lambda^{-4}$ \cite{29}. He assumed that radiation inside the cubical box formed a standing wave. His idea was to calculate the energy density according to

$$ \rho(\nu, T) = [N(\nu)d\nu]e(\nu, T) $$

(46)

where $N(\nu)d\nu$ is the number of standing waves per unit volume having a frequency in a range of $\nu$ to $\nu + d\nu$ and $e(\nu, T)$ is the mean energy of the wave. If $k$ is the wavenumber (reciprocal of $\lambda$), then the number of points for which $k$ lies between $k + dk$ is proportional to the $k^2dk$. According to the Maxwell-Boltzmann law, energy is equally divided among all modes. And since the energy of each mode is proportional to temperature $T$, the energy of radiation whose wavenumber is in the range $k$ and $k + dk$ will be proportional to the

$$ Tk^2dk. $$

(47)

Or the energy of radiation whose wavelength is the range $\lambda$ and $\lambda + d\lambda$ will be

$$ \rho(\lambda, T)d\lambda \propto T\lambda^{-4}d\lambda $$

(48)

In Rayleigh’s first paper on radiation, he did not calculate the proportionality constant. The correct form of Rayleigh’s law came five years later by Rayleigh himself and James Jeans. In between, Max Planck published two remarkable papers on radiation theory and introduced the idea of discretization of energy. In September 1900, Lummer and Pringsheim experimentally checked that
Rayleigh’s law agrees only for large wavelength. Lummer and Pringsheim put exponential factor $\exp[-c_2/(\lambda T)^3]$ and from the best-fitted data, they gave an empirical distribution function \[16\] [17]

$$\rho(\lambda, T) = c_1T\lambda^{-4} \exp[-c_2/(\lambda T)^{1.3}] \quad (49)$$

6. Enter Max Planck

**Figure 8.** Max Planck (1858-1947)  [Photo: Wikipedia Commons]

The German physicist Max Karl Ludwing Planck was trying to justify Wien’s distribution function using Heinrich Hertz’s idea of electric dipole radiation. From Kirchhoff’s law, Planck understood that the black-body radiation is independent of the material of the cavity, so he presumed that the cavity walls are made of a collection of damped oscillators of electric dipoles (Hertzian resonators).

Planck considered the steady state case where the energy emitted by the accelerating charges via electromagnetic radiation should be equal to the energy absorbed by the oscillators. So oscillations are damped-driven. In 1899, he proved that the average energy of oscillator $U(\nu, T)$ is related to the energy density of radiation
\[ \rho(\nu, T) \] in the following way\(^2\)

\[ \rho(\nu, T) = \frac{8\pi \nu^2}{c^3} U(\nu, T) \] (50)

In October 1900, Planck published the paper “On the theory of the energy distribution law of the normal spectrum”, where he guessed the function of the second derivative of entropy with respect to average energy and gave the first appearance of today known as Planck’s distribution function. Planck made first simplest guess that

\[ \frac{d^2S}{dU^2} = -\frac{\alpha}{U} \] (51)

Integrating the above equation with respect to \(U\), we get

\[ \frac{dS}{dU} = -\alpha \ln U + C \] (52)

At constant volume, the fundamental thermodynamic equation \(dU = TdS - PdV\) becomes

\[ \frac{dS}{dU} = \frac{1}{T} \] (53)

Now, using the equations (52) and (53), we get

\[ U = A \exp \left(-\frac{1}{\alpha T}\right) \] (54)

where \(A\) and \(\alpha\) depend upon the frequency of radiation. We get Wien’s distribution function by putting this into equation (50).

Now, if we say energy is linearly proportional to the temperature, then using equation (53), we get

\[ \frac{dS}{dU} = \frac{\alpha'}{U} \] (55)

So,

\[ \frac{d^2S}{dU^2} = -\frac{\alpha'}{U^2} \] (56)

\(^2\)For further details and proof, readers can refer to Max Jammer book *The Conceptual Development of Quantum Mechanics, Appendix A, pp. 470 and D. Ter. Haar book [36] Chapter 1, pp. 8"
Planck then combined both equations (51) and (56) and made an intelligent guess that

$$\frac{d^2S}{dU^2} = -\frac{\alpha}{U(U + \beta)}$$  \hspace{1cm} (57)

Integrating the above equation with respect to $U$, we get

$$\frac{dS}{dV} = \frac{\alpha}{\beta} \ln \left(1 + \frac{\beta}{U}\right) + \text{Constant}$$ \hspace{1cm} (58)

Using equation (53) and rearranging terms, we get

$$U = \frac{a}{\exp(b/T) - 1}$$ \hspace{1cm} (59)

Putting into equation (50) and comparing with Wien’s scaling law (equation (33)), we get the energy distribution function \cite{26, 27, 14, 6}

$$\rho(\nu, T) = \frac{c_1\nu^3}{\exp(c_2\nu/T) - 1}.$$ \hspace{1cm} (60)

After a week, physicists Heinrich Rubens and Friedrich Kurlbaum published experimental verification of Planck’s energy distribution for temperatures 85 K to 1773 K, and it matched with all ranges of wavelengths \cite{6}. In December 1900, Planck published a second paper with a detailed explanation of the energy distribution where he introduced an idea of energy quantization and a universal constant known as Planck’s constant \cite{36, 14}.

In the second paper, Planck thought that the total entropy of Hertzian resonators should be maximum at equilibrium. So it can be calculated from Boltzmann combinatorial method (microcanonical ensemble in statistical mechanics) \cite{25}. Let $U$ be the average energy and $S$ be the entropy of a single oscillator. Then total energy of $N$ oscillators is $U_N = NU$, and total entropy is $S_N = NS$. Let $W$ be the total number of micro-states, then total entropy will be $S_N = k_B \ln W$. Now, Planck thought that to calculate $W$ of $N$ oscillators having total energy $U_N$, we should take $U_N$ to be composed of discrete values of an element $\epsilon$ rather than a continuous one. So one can assume

$$U_N = n \cdot \epsilon$$ \hspace{1cm} (61)
where \( n \) is a large integer. Now, to calculate \( W \), Planck calculated the number of ways to distribute \( n \) bundles of energy \( \epsilon \) among \( N \) oscillators. Then,

\[
W = \frac{(N + n - 1)!}{(N - 1)! n!} \approx \frac{(N + n)!}{N! n!} \tag{62}
\]

Using the Stirling approximation for large number \( N! \approx N^N \), we get

\[
W \approx \frac{(N + n)^{N+n}}{N^N n^n} \tag{63}
\]

Then, the total entropy will be

\[
S_N = k_B \left[ (N + n) \ln(N + n) - N \ln N - n \ln n \right] \tag{64}
\]

Now, using the \( U_N = n\epsilon = NU \) and \( S_N = NS \) we get

\[
S = k_B \left[ \left( 1 + \frac{U}{\epsilon} \right) \ln \left( 1 + \frac{U}{\epsilon} \right) - \frac{U}{\epsilon} \ln \frac{U}{\epsilon} \right] \tag{65}
\]

Using the \( \rho = \frac{8\pi^2}{3c^3}U \) and Wien’s scaling law, we get

\[
U = \nu f \left( \frac{T}{\nu} \right) \tag{66}
\]

Rearranging terms, we can write a different form

\[
T = \nu g \left( \frac{U}{\nu} \right) \tag{67}
\]

using the \( \frac{1}{\nu} = \frac{dS}{dU} \), we get

\[
\frac{\partial S}{\partial U} = \frac{1}{\nu} g \left( \frac{U}{\nu} \right) \tag{68}
\]

Integrating, we get

\[
S = g \left( \frac{U}{\nu} \right) \tag{69}
\]

Comparing equation (65) and (69), we must have \( \epsilon \propto \nu \). The proportionality constant “\( h \)” is known as Planck’s universal constant. So putting \( \epsilon = \hbar \nu \) into equation (65) we get entropy

\[
S = k_B \left[ \left( 1 + \frac{U}{\hbar \nu} \right) \ln \left( 1 + \frac{U}{\hbar \nu} \right) - \frac{U}{\hbar \nu} \ln \frac{U}{\hbar \nu} \right] \tag{70}
\]
Again, using the $\frac{1}{T} = \frac{4\xi}{dT}$, we get

$$\frac{1}{T} = \frac{k_B}{h \nu} \ln \left( 1 + \frac{h \nu}{U} \right)$$

(71)

So,

$$U = \frac{h \nu}{\exp(h \nu/k_B T) - 1}$$

(72)

So we get the energy distribution

$$\rho(\nu, T) = \frac{8\pi \nu^2}{c^3} \frac{h \nu}{\exp(h \nu/k_B T) - 1}$$

(73)

This is known as Planck’s distribution function. The universality of the Boltzmann constant was also given by Planck. Using experimental results of F. Kurlbaum on total radiation energy and Lummer and Pringsheim on Wien’s displacement law Planck also calculated the value of the $k_B$ and $h$ [25]. The word ”quanta” was first used by Planck for energy elements. This strange non-classical theory of discretization of energy was uncomfortable to many scientists, including Planck himself. Albert Einstein’s concern was that equation (50) which Planck derived using the classical theory of absorption and emission of the energy by the oscillator, and in equation (61), Planck defined the mean energy of the oscillator in a non-classical way. Scientists started taking Planck’s idea seriously when Einstein explained Heinrich Hertz’s experiment photoelectric effect by assuming that the energy of the light comes in quanta (light quanta) in 1905-06.

Einstein’s concern about Planck’s theory got resolved when Peter Debye [9, 2] gave an alternative approach to Planck’s theory using the idea from Rayleigh-Jeans theory. Debye considered electromagnetic radiation in a cubical cavity similar to the proper vibration of the crystal. So the number of vibrations in the interval $\nu$ to $\nu + dv$ is the same as in the calculation of Rayleigh and Jeans. The energy of the proper vibration of frequency $\nu$ can be from possible values $n h \nu$, where $n$ is an integer. Using the Boltzmann probability distribution $P(n) = A \exp(-nh \nu/k_B T)$, the mean energy of the
proper vibration will be

\[
e(\nu, T) = \frac{\sum_n n h \nu P(n)}{\sum_n P(n)}
\]

(74)

\[
e(\nu, T) = \frac{\nu x}{1 + x + x^2 + \ldots}, \quad \text{where} \quad x = e^{-h\nu/k_B T}
\]

(75)

\[
e(\nu, T) = \frac{h\nu}{1 - x^2}
\]

(76)

\[
e(\nu, T) = \frac{h\nu}{\exp(h\nu/k_B T) - 1}
\]

(77)

Using the equation (46) and (81) gives the energy density

\[
\rho(\nu, T) = \frac{8\pi \nu^2 c^3}{\exp(h\nu/k_B T) - 1}
\]

(78)

7. Enter James Jeans

Even though Planck’s distribution agreed with experimental data for many years, physicists could not believe the idea of quantization of energy. Rayleigh was still working on his idea of radiation in a cubical enclosure. In April 1905, he calculated a numerical constant in his previously proposed radiation energy distribution [30]. Rayleigh considered the standing wave \( f(x, y, z, t) = A \sin(k_x x + k_y y + k_z z) \cos(\omega t) \) in cubical enclosure of length \( l \). In order to satisfy the boundary condition \( f = 0 \) at \( x, y, \) or \( z = 0 \) or
we must have \( k_x = \frac{n_x \pi}{l}, \quad k_y = \frac{n_y \pi}{l} \) and \( k_z = \frac{n_z \pi}{l} \) where \( n_x, n_y, n_z \in \mathbb{N} \). Since \( k^2 = k_x^2 + k_y^2 + k_z^2 \), we have

\[
    n_x^2 + n_y^2 + n_z^2 = \frac{k^2 l^2}{\pi^2}
\]

which is a sphere of radius \( \frac{kl}{\pi} \).

Now, to calculate the total number of modes, we need to take eighth part of the volume of a sphere since \( n_x, n_y, \) and \( n_z \) can only take positive values. Rayleigh considered the positive as well as negative values. This mistake was pointed out by English physicist sir James Hopwood Jeans in May 1905 [10].

With the Jeans correction factor, the total number of modes from 0 to \( k \) is

\[
    \frac{1}{8} \frac{4}{3} \pi \left( \frac{kl}{\pi} \right)^3
\]

then the number of modes per unit volume in frequency range \( \nu \) to \( \nu + d\nu \) is

\[
    N(\nu) d\nu = \frac{4\pi \nu^2}{c^3} d\nu
\]

where we have used \( k = \frac{2\pi \nu}{c} \) and factor ‘2’ is multiplied since there are two polarization states of vibration. In terms of wavelength, the number of modes in the range \( \lambda \) to \( \lambda + d\lambda \) is

\[
    N(\lambda) d\lambda = \frac{8\pi}{\lambda^4} d\lambda
\]

Now, since each mode is vibrating with average energy \( k_B T \), from equation (46), we have the energy density

\[
    \rho(\lambda, T) = \frac{8\pi}{\lambda^4} k_B T
\]

This is known as ‘Rayleigh-Jeans law’. It agrees with experiments for large wavelength, but as we go towards a small wavelength (in the UV region of the black-body spectrum), energy density becomes very large. In 1911, Paul Ehrenfest called this the ‘Ultraviolet catastrophe’. But the problem was already solved.
by Planck. It took many years even for experts working in the
field to appreciate that.

8. Conclusion

Textbooks usually do not explain Max Planck’s original derivation
of the black-body spectrum and how exactly he introduced
the quantization of energy. Planck’s first assumption that cavity
walls can be considered as a collection of oscillators of electric
dipoles came from Kirchhoff’s law of universality. Instead of di-
rectly working on the energy distribution function like Rayleigh
and Jeans, Planck used the microcanonical ensemble in statistical
mechanics and tried to calculate the entropy of the oscillators.
Planck showed that entropy is a function of the ratio of the aver-
age energy of oscillators and the frequency of radiation, and by
comparing that with entropy calculated from the Boltzmann com-
binatorial method, he concluded that there should be some uni-
versal constant (Planck’s constant $h$) such that energy of quanta
is $hv$. Most of the textbook explains Wien’s displacement law
and Stefan-Boltzmann law as a consequence of Planck’s distrbution
function, but historically Planck used Wien’s scaling law
to get the function of entropy of oscillator (equation (69)). To
derive the scaling law, Wien used the Stefan-Boltzmann $T^4$ law.
So not only for a historical purpose but also to understand the
black-body radiation rigorously, one needs to understand the orig-
inal calculation and arguments of Kirchhoff, Stefan, Boltzmann,
Wien, Rayleigh, Planck and Jeans in an actual timeline.

The logical way to explain the black-body radiation is to under-
stand the Rayleigh-Jeans law first and appreciate that the black-
body spectrum can not be explained using classical physics, and
we need some different new formalism. Then by introducing the
energy quantization and calculating the mean energy of Planck’s
cavity oscillators using the canonical ensemble in statistical me-
chanics, we can explain Planck’s distribution function as a modi-
Many physicists feel that Wien’s and Planck’s original derivations of their law are pretty long and not worth discussing, but we should at least appreciate the fact that without Kirchhoff’s universal law, Stefan-Boltzmann $T^4$ law and Wien’s scaling law, Planck may never have got the idea of energy quantization.

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**Appendix**

**A. Derivation of the radiation pressure**

Using the classical theory of electromagnetic radiation, we know that the momentum density is given by the ratio of the energy density and speed of light.

Let us say radiation is propagating, as shown in the figure. Consider a tube of length $c\Delta t$ with a cross-section area $L \times L$, making an angle $\theta$ with the normal to the walls. The energy contained in the tube is approximately

$$\rho_T c\Delta t L^2$$

(84)

Then the energy per unit area of the wall is given by

$$\rho_T c\Delta t \frac{L^2}{L \times \frac{L}{\cos\theta}} = \rho_T c\Delta t \cos\theta$$

(85)
Thus, the momentum per unit area along the incident line is
\[ \rho_T \Delta t \cos \theta \]  
(86)

But, the \( \cos \theta \) component of this momentum will contribute to the radiation pressure. And since the wall is perfectly reflecting, the net momentum transfer will be
\[ 2 \rho_T \Delta t \cos^2 \theta \]  
(87)

By dividing \( \Delta t \), we get the force per unit area, i.e. pressure on the wall is
\[ 2 \rho_T \cos^2 \theta \]  
(88)

The total pressure exerted can be calculated by taking the angular average. So,
\[ P = \frac{1}{4\pi} 2 \rho_T \int_0^{\pi/2} \cos^2 \theta \sin \theta d\theta \int_0^{2\pi} d\phi \]  
(89)

The \( \theta \) integration gives the factor \( \frac{1}{3} \), So the radiation pressure is
\[ P = \frac{\rho_T}{3} \]  
(90)

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