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I think chapters 1-3 are in great shape. And with a little more work/polishing,
Ch 4 will be great, too!
Nice job!

Modeling Passive Daytime Radiative Cooling Devices (PDRCs) using COMSOL Multiphysics™

In particular, please focus on addressing the two comments circled in red
in sections 4.1 & 4.2. Also on all of the blue comments in section 4.3.

by
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A thesis submitted in partial fulfillment
of the requirements for the
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in Physics

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Abstract

This thesis delves into the modeling and simulation of Passive Daytime Radiative Cooling (PDRC) devices using COMSOL Multiphysics™, a cutting-edge computational tool. At its core, PDRC technology harnesses the principle of radiative cooling to emit heat directly into outer space, offering a sustainable solution to the pressing challenges of global warming and the energy crisis. By optimizing the optical properties and layering strategies of these devices, the thesis seeks to enhance their cooling efficiency and operational performance, providing a blueprint for future innovations in cooling technologies. Through rigorous computational analysis and simulation, it validates theoretical models and investigates the potential of various materials and configurations to improve PDRC performance, aiming to contribute significantly to the development of energy-efficient cooling solutions.

Executive Summary

Your executive summary will give a detailed summary of your thesis, hitting the high points and perhaps including a figure or two. This should have all of the important take-home messages; though details will of course be left for the thesis itself, here you should give enough detail for a reader to have a good idea of the content of the full document. Importantly, this summary should be able to stand alone, separate from the rest of the document, so although you will be emphasizing the key results of your work, you will probably also want to include a sentence or two of introduction and context for the work you have done.

Acknowledgments

(Note to Prof. Hudgings: I intend to refine these acknowledgments to make them more personal and succinct by the time the thesis reaches the second reader).

In this section, I express my heartfelt gratitude to those who have supported me throughout this journey. First and foremost, a huge thanks to Professor Janice Hudgings, whose incredible guidance, patience, and unwavering optimism have been a beacon of light for me, especially during challenging times with COMSOL. Her enthusiasm and belief in my project have been invaluable.

A special acknowledgment goes to Professor James Higdon, whose generosity in providing his COMSOL license and desktop was instrumental in initiating my thesis work. His willingness to assist during technical difficulties and his provision of essential resources for mastering COMSOL have been fundamental to my progress.

My friends, who have been a constant source of support and encouragement, deserve a world of thanks...

The Hudgings lab members, for their insights and camaraderie, have contributed significantly to my journey...

My suitemates, for their understanding and encouragement, have made this process smoother...

And, of course, my parents, whose love and belief in me have been my foundation...

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Chapter 1

Introduction

This thesis centers on the modeling of Passive Daytime Radiative Cooling Devices (PDRCs) utilizing the COMSOL Multiphysics™ software. While all objects with a temperature above absolute zero emit blackbody radiation, PDRCs are distinct in their ability to efficiently radiate heat in the mid-infrared range where the Earth's atmosphere is most transparent. This allows PDRCs to effectively transfer heat directly to the cold sink of outer space during daylight hours, without the need for ~~input~~ ^{FIXED} energy. Therefore, PDRCs hold the promise of addressing two significant challenges: the energy crisis and global warming.

1.1 Cooling is Critical

Over the years, cooling has become more critical to humans due to global warming, rapid population growth and industrial development [4]. Various methods exist for cooling buildings, ranging from traditional practices, such as shading and solar orientation, to the use of electric fans. The most advanced approach is air conditioning (AC), encompassing systems that enhance indoor thermal comfort and air quality. While mechanical cooling techniques date back to the 19th century, widespread adoption of air conditioning began in the 1950s, driven by improved performance, affordability, and economic prosperity, primarily in the United States [5].

Modern AC systems vary widely in size and cost, catering to individual rooms or entire buildings, with electricity being the predominant power source. Today, the largest concentration of cooling systems is found in urban areas, both in industrialized nations and emerging economies, reflecting the higher population density and greater demand for climate control in these regions [5].

Particularly in the realm of residential air conditioners (ACs), China emerges as the foremost market with a staggering sale of 41 million units. Following China, ~~the USA~~ Japan and the European Union represent the subsequent largest markets for residential ACs. However, there is a significant uptick in sales within various emerging economies, notably in Asia (see ~~figure~~ ^{FIXED} figure 1.1).

Global sales of ACs have exhibited consistent growth in recent years. Over the period

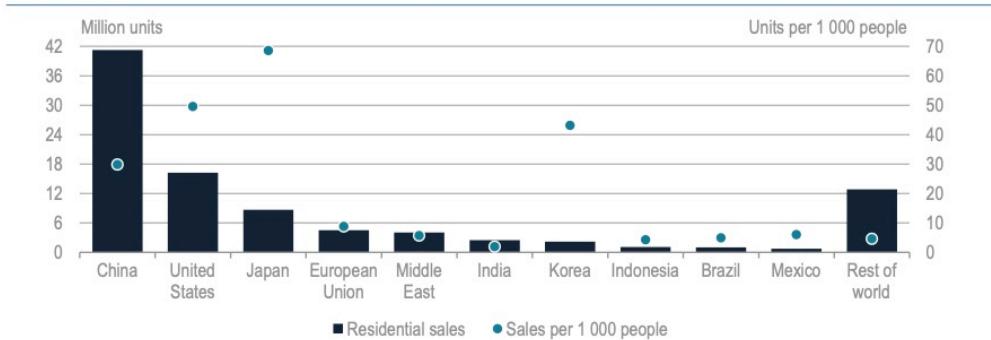
Figure 1.5 • Sales of residential ACs by country/region, 2016

Figure 1.1: Sales of Residential ACs by Country or Region, 2016. Source: [5].

from 1990 to 2016, annual AC sales experienced a nearly fourfold increase, reaching 135 million units. In 2016, China emerged as the leading market in terms of AC capacity sales, totaling nearly 390 gigawatts (53 million units) [5].

The growing demand for cooling is significantly influencing power systems, primarily due to the reliance on electricity-driven fans or air conditioners to meet cooling requirements. The escalating demand for air conditioning, in particular, not only elevates overall electricity consumption but also contributes to higher peak electricity loads. Additionally, the emission of greenhouse gases (GHGs) from ACs occurs through refrigerant leakage or improper disposal. ~~It's~~ ^{FIXED} It's noteworthy that these refrigerants are potent GHGs with adverse implications for climate change [5].

Improving the efficiency of air conditioning systems (ACs) is pivotal in mitigating peak electricity demand, thereby resulting in decreased emissions and associated financial implications. Endeavors focused on enhancing cooling efficiency necessitate a thorough assessment of the comparative costs linked to diverse cooling technologies.

1.2 Radiative Cooling

Objects with temperatures above absolute zero emit blackbody radiation, with a spectrum that depends on their temperature as governed by Planck's law. While the emitted radiation covers a range of wavelengths, it is not uniformly distributed across all wavelengths. Radiative passive cooling occurs when objects emit more radiation, particularly in the infrared spectrum, than the combined radiation they absorb, which includes both blackbody and solar radiation. Thus radiative passive cooling is an electricity-free method for cooling terrestrial entities [6].

The heat emitted by these objects, which exceeds the heat they absorb, is transferred to outer space via thermal radiation, leveraging the substantial temperature difference between Earth (approximately 300 K) and outer space (approximately 3 K) ~~See~~ figure 1.2 for net heat loss to space). This process efficiently exchanges heat with the infinite cold reservoir of deep space, achieving cooling without any energy consumption [4].

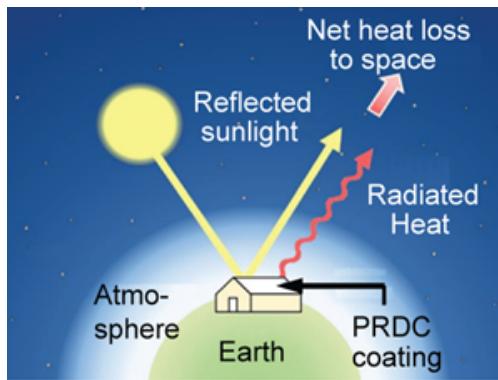


Figure 1.2: Schematic for Radiative Cooling. Source: [6]

Passive radiative cooling can be realized even during the daytime, necessitating precise tuning of optical properties across a broad spectrum of wavelengths, from ultraviolet to mid-infrared (see Figure 1.3). As a result, achieving effective passive daytime radiative cooling imposes stringent requirements on materials and structures to mitigate solar heating [6]:

1. Minimal absorptivity (α) approaching 0% (equivalent to nearly 100% reflectance, R) in the solar spectrum, ranging from $0.3\text{--}2.5\ \mu\text{m}$. This characteristic ensures that the surface absorbs minimal solar energy during daylight, thereby reducing the heat gained by the PRDC.
2. High thermal radiation in the atmospheric transparency window, with an emittance (ε) close to 1 within the long-wavelength infrared (LWIR) transmission window of the atmosphere ($\lambda = 8\text{--}13\ \mu\text{m}$). This range is significant due to the atmosphere's partial transparency and minimal infrared absorption by gas molecules in this spectrum.
3. An emittance (ε) close to 0 in other mid-infrared wavelengths, such as $5\text{--}8\ \mu\text{m}$ and those greater than $13\ \mu\text{m}$. This characteristic is crucial due to the atmosphere's opacity in these spectral ranges, *and helps to minimize atmospheric BB radiation* FIXED *heating of the device due to* FIXED

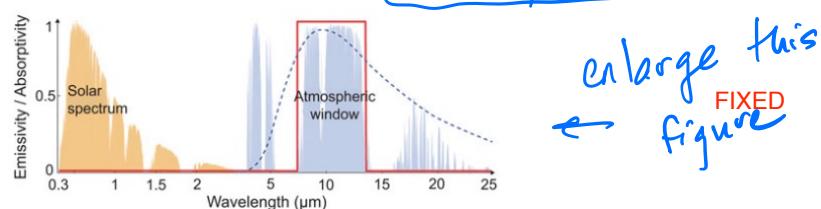


Figure 1.3: Ideal Optical Properties of a Radiative Cooling Surface. Source: [6]

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1.3 Literature Review

At the nexus of physics and technological advancement, Passive Daytime Radiative Cooling Devices (PDRCs) stand as a symbol of optimism amidst the challenges of global warming and the energy crisis. PDRCs are notable not just for their operational efficiency but also for their capacity to transform our approach to energy consumption and environmental conservation. While radiative cooling for nighttime use was established decades ago, only recently has there been significant progress in achieving cooling directly under sunlight, suggesting a promising upward trajectory for the technology's cooling capabilities.

1.3.1 Historical Overview of Radiative Cooling.

The concept of radiative cooling, hypothesized by French Engineer Félix Trombe in 1965, initially focused on nighttime cooling [7]. Significant strides have recently been made in achieving daytime cooling under direct sunlight, marking a pivotal advancement in the field. Trombe's collaboration at Montlouis Laboratory led to a cooling method that significantly reduced temperatures ~~without insulation~~, leveraging a system that could cool by 14 to 30 degrees Celsius relative to the surroundings. This system involved a north-facing, slightly tilted facade with infrared-transparent panels and a selective radiator, optimizing both night and daytime cooling as seen in figure 1.4 [8].

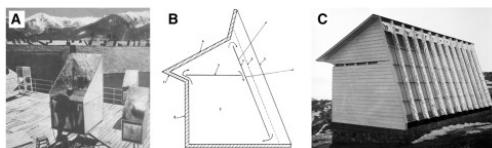


Figure 1.4: Experimental Cooling Container by Trombe and his team in Mont-Louis, France. (A) depicts the experimental cooling setup developed by Trombe's team in Mont-Louis. (B) presents a cross-sectional view of Trombe's patented cold house design. (C) offers a northeast perspective of the cold house, showcasing its radiative cooling facade, a creation of Felix Trombe and his collaborators in 1963. Source: [8]

Building on Trombe's groundbreaking efforts, in 1975, Catalanotti and his team developed an experimental setup for daytime radiative cooling. This setup combined metals and TEDLAR (polyvinyl-fluoride plastic) to protect the cooling device from direct sunlight, although it was more effective for nighttime use [9]. This TEDLAR film, offering somewhat selective and high infrared (IR) absorption within the 9 to 13 μm wavelength range, also absorbs IR significantly beyond 20 μm , impacting its selectivity and, consequently, the efficiency of radiative cooling. Despite this, they achieved a temperature decrease of 10 degrees Celsius below ambient under diffused sunlight. The primary drawback of using PVF polymer for daytime cooling is its substantial absorption in the solar spectrum [10].

Further research explored the potential of thin films on aluminum substrates, with specific material compositions, such as coated thin solid film on an aluminum substrate with $SiO_6N_{0.2}$

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enough from the sources
to avoid any risk
of plagiarism.*

and double layers of SiO_2 and $\text{SiO}_{0.25}\text{N}_{1.52}$, aimed at achieving selective emission. Despite these innovative approaches, effectively cooling during daylight remained a challenge [9].

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1.3.2 Challenges with Material Selection.

Ongoing advancements have aimed to enhance the cooling capabilities of radiative emitters, achieving notable success in nocturnal cooling. Yet, the reliance on both naturally sourced and synthetic polymers has imposed constraints. The absence of materials that combine high solar reflectivity with potent IR emission has hindered effective cooling under direct sunlight. Investigations into selective IR emitters, including polymer films, white pigmented paints, and SiO films, have encountered limitations due to their inadequate emissivity and the wide range of their IR absorption, which absorbs too much atmospheric radiation, thus failing to significantly lower temperatures below ambient conditions [10].

1.3.3 Exploration of Polymer Films and Pigmented Paints.

Additional polymer films such as polyvinylchloride (PVC) and poly(4-methylpentene) (TPX) were evaluated for their potential in radiative cooling. However, their performance, particularly in terms of IR emissivity and selectivity within the crucial $8\text{-}13 \mu\text{m}$ range, fell short when compared to PVF [10].

Furthermore, investigations into pigmented paints, inorganic compounds, and gases capable of IR emission for nighttime cooling have shown promise. Specifically, white paint containing titanium dioxide (TiO_2), applied to aluminum plates, demonstrated notable cooling effects under clear sky conditions at night, despite less effectiveness under direct midday sun. This suggests pigmented paints might offer advantages for nocturnal cooling due to their application versatility on different substrates [10].

The application ~~of~~ *of* Silicon monoxide (SiO) films and other inorganic materials in radiative cooling technologies leverages their capacity for targeted infrared (IR) emission. These substances are recognized for their potential to efficiently radiate heat from surfaces, aiding in cooling processes. Nevertheless, their effectiveness is somewhat diminished by a narrow emission bandwidth, limiting the spectrum of IR radiation they can emit. This constraint may decrease the volume of heat dispersed into the environment, thereby affecting the device's cooling efficacy [10].

1.3.4 Innovations in Solar Reflectors and Convection Shields.

Investigations into solar reflectors and IR-transparent materials for daytime cooling have proven effective in both reflecting solar radiation and facilitating IR emission. This capability is essential for daytime cooling to both prevent solar heat accumulation and promote surface heat release. However, maintaining this delicate balance during peak sunlight, when solar irradiance is most intense, presents a considerable challenge. Achieving the optimal mix of

solar reflection to avoid overheating while ensuring adequate IR emission for cooling under direct sunlight is particularly complex [10].

Beyond solar reflectors, convection shields significantly enhance a radiative cooler's effectiveness. In daytime operations, when a cooler's net radiative output exceeds its solar absorption, it can significantly lower temperatures beneath the ambient level, assuming convective heat gains are minimized. However, employing convection shields might not always be beneficial, as natural convection can help dissipate heat from the device. Yet, without direct sunlight, convection shields can markedly boost cooling efficiency to its maximum potential [10].

1.3.5 Photonic Devices and Microstructure Advances.

The advancement in photonic-based radiative coolers represents a significant leap towards achieving efficient cooling directly under sunlight, allowing temperatures to fall below ambient levels. This innovative approach differs from traditional methods that rely on the natural optical characteristics of materials, as it harnesses engineered photonic properties in tandem with those natural characteristics to enhance cooling performance [10].

Raman and colleagues conducted the inaugural experiment showcasing radiative cooling under direct sunlight with planar photonic structures (see figure 1.5), achieving a notable 4.9°C drop below ambient temperature. Their device, composed of seven layers alternating between hafnium oxide (HfO_2) and silicon dioxide (SiO_2) of varied thicknesses, utilized the upper thick layers for IR emission across the $8\text{--}13 \mu\text{m}$ range. The lower, thinner layers on an aluminum-coated silicon wafer acted as a chirped 1D photonic crystal, reflecting up to 97% of solar radiation [1].

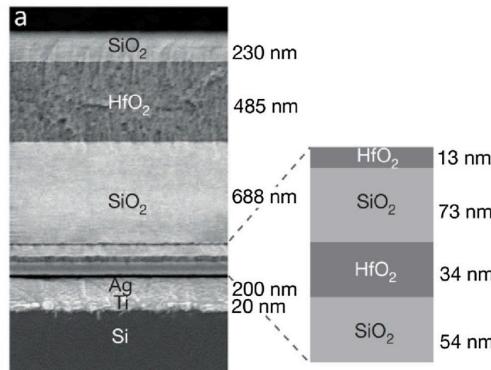


Figure 1.5: Radiative cooling device consisting of seven layers of HfO_2 and SiO_2 , whose thicknesses are defined by extensive numerical optimization (see Methods), on top of 200 nm of Ag, a 20-nm-thick Ti adhesion layer, and a 750-mm-thick, 200-mm-diameter Si wafer substrate. Source: [1]

1.3.6 Future Directions and Microfabrication Techniques.

Metallic microstructures, like plasmonic structures and metallic photonic crystals, offer the potential for highly selective infrared (IR) emission. Achieving this selective emission across the full 8–13 μm wavelength spectrum remains a challenge. Anisotropic metamaterials have shown promise, capable of strong and precise IR emission within this range. This technology could potentially enable cooling significantly below ambient temperatures by more than 10 degrees Celsius, illustrating the advanced capabilities and potential of these engineered materials for effective radiative cooling [10].

Leveraging photonic microstructures for efficient cooling shows great promise but faces hurdles in practical application, particularly for daytime use where such radiators need to be paired with IR transparent solar reflectors to minimize solar absorption. This pairing, however, compromises the precise IR emission. Nonetheless, advancements in microfabrication techniques, like UV-nanoimprint lithography, offer improved prospects for scaling up production, potentially overcoming some of the challenges in deploying these technologies widely [10].

1.4 Previous Project Work and Project Goals

Research on PDRCs has been an ongoing project in the Hudgings lab. This thesis aims to build upon the work of Paul McKinley ~~Fixed~~ (class of 2022) and Genevieve DiBari ~~Fixed~~ (class of 2023).

McKinley laid the groundwork by developing a rooftop testbed and the fabrication process for PDRCs (a process I plan to replicate after completing the modeling of various PDRC iterations on COMSOL). DiBari contributed by modeling PDRCs in Python and enhancing the initial outdoor testing setup.

For my thesis, the goal is to model different PDRC structures using COMSOL, exploring materials that could enhance the optical properties of an ideal PDRC device. This includes stacking materials of varying thicknesses and refractive indices on current PDRC models in the Hudgings lab to ~~spectrally tailor reflectance~~ increase reflectivity (R). If time permits, promising PDRC models (with high R and/or emissivity in the atmospheric window) will be fabricated after the modeling phase.

→ Maybe change this goal to developing a COMSOL model of the PDRC devices used in the Hudgings lab, quantitatively comparing your COMSOL predictions of $R(\lambda)$, $\varepsilon(\lambda)$ with experimental measurements.

Excellent job on this chapter! Would you please just double check the lit review sections and make sure they are substantially reworded from the source articles?

Chapter 2

The Physics Behind PDRCs

Be careful with reflectivity vs. reflectance, etc

A Passive Daytime Radiative Cooling (PDRC) device operates by absorbing a lower amount of radiation than it emits, thereby facilitating electricity-free cooling, even in daylight conditions. Consequently, one of the pivotal attributes of a PDRC device is the imperative for an absorptivity (α) as close to 0% or, conversely, a reflectivity (R) of 100% within the solar spectrum (ranging from 0.3 to 2.5 micrometers). This specification ensures that the device's surface remains entirely unaffected by solar heating during daylight hours. Additionally, it is critical that the device exhibits an emissivity or absorptivity close to 100% over long wavelengths, particularly within the atmospheric window (ranging from 8 to 13 micrometers), to maximize thermal radiation emission back into space. In this wavelength range, the device's reflectivity should be nearly 0%, which further enhances its cooling efficiency by minimizing the absorption of Earth-emitted radiation, thereby optimizing the device for passive cooling under direct sunlight. See figure 2.1 for these ideal properties. Fixed: deleted

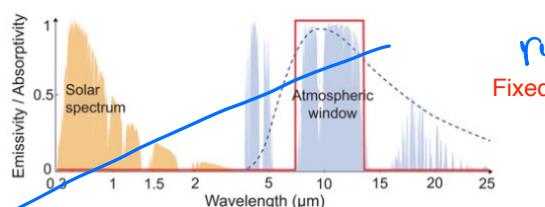


Figure 2.1: Ideal Optical Properties of a Radiative Cooling Surface. Source: [6]

To enhance the effectiveness of PDRC, it becomes essential to optimize this reflectivity (R) spectrum. One approach for achieving this goal is to perceive light as an electromagnetic wave, and from this perspective, ultimately derive R through the renowned *Fresnel equations*. Fixed

The Fresnel equations are mathematical expressions that delineate the proportion of incident electromagnetic field Fixed that is either transmitted or reflected at the interface of two materials with differing refractive indices. This concept aligns precisely with our objectives, as we plan to stack planar surfaces featuring distinct reflective properties and refractive indices. This chapter serves as an exploration of the theoretical framework underpinning the derivation of R via the Fresnel Equations, delving into associated phenomena such as total

internal reflection. Additionally, we explore the practical application of these principles to PDRC devices.

2.1 Fresnel Equations

The derivation below follows that given in [Fixed \(cite\)](#). *between 2 materials*

Consider a light ray incident at point P upon a planar interface, leading to the generation of both reflected and refracted rays. It is important to recognize that the refractive index before the interface (n_1) and after the interface (n_2) are properties intrinsic to the materials themselves, determining how light propagates through them. Specifically, n_1 is the refractive index of the material through which the incident and reflected rays travel, while n_2 is the refractive index of the material through which the refracted ray travels. The plane of incidence, lying within the x-z plane, is defined by both the surface normal and the direction of the incident ray. See [Fixed](#) Figure 2.2.

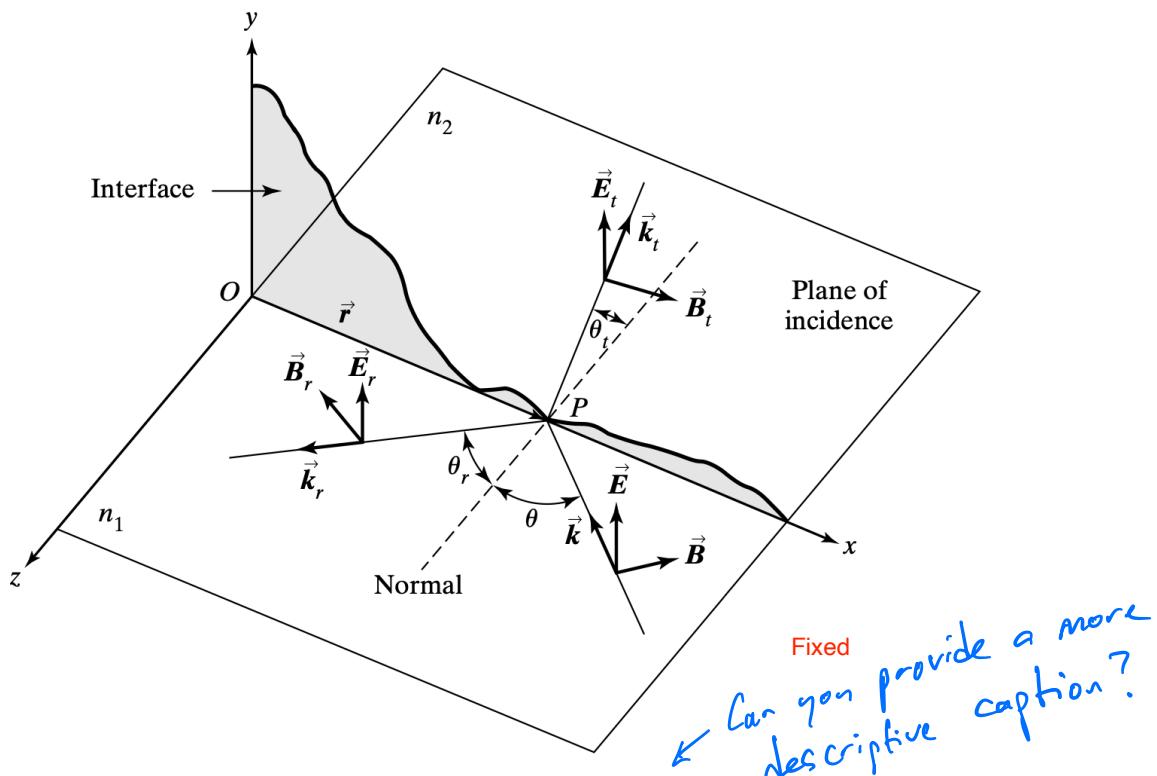


Figure 2.2: The transverse electric (TE) set-up. Source: [3]

In the context of each ray, the direction of wave propagation (\vec{k}), can be established by the vector cross product of the electric field (\vec{E}), and magnetic field (\vec{B}) vectors, expressed as $\vec{E} \times \vec{B}$. This relationship can be conveniently determined using the right-hand rule, offering a valuable method for directional assessment.

Let us consider that the incident light comprises of plane harmonic waves:

$$\vec{E} = \vec{E}_0 e^{i(\vec{k} \cdot \vec{r} - \omega t)} \quad (2.1)$$

where

- \vec{E} represents the electric field vector of the wave at any point.
- \vec{E}_0 is the amplitude of \vec{E} which represents the maximum strength or value of \vec{E} .
- \vec{k} is the wave (direction) vector and it indicates the direction of wave propagation and has a magnitude equal to the wave number $k = \frac{2\pi}{\lambda}$.
- \vec{r} represents the position vector in space where the field is being examined.
- ω is the angular frequency of the wave.
- t represents the time.
- i , the imaginary number, indicates the wave has both real and imaginary components.

In our analytical approach, we will specifically examine a linearly polarized light wave, where the electric field vector \vec{E} is oriented perpendicular to the plane of incidence. According to the right-hand rule, this configuration places the magnetic field vector \vec{B} within the plane of incidence. This particular polarization is known as the *transverse electric* (TE) mode

The reflected and transmitted waves can then be expressed as plane harmonic waves:

$$\vec{E}_r = \vec{E}_{0r} e^{i(\vec{k}_r \cdot \vec{r} - \omega_r t)} \quad (2.2)$$

$$\vec{E}_t = \vec{E}_{0t} e^{i(\vec{k}_t \cdot \vec{r} - \omega_t t)} \quad (2.3)$$

where

- \vec{E}_r represents the electric field vector of the reflected wave at any point.
- \vec{E}_t represents the electric field vector of the transmitted wave at any point.
- \vec{k}_r is the wave (direction) vector of the reflected wave.
- \vec{k}_t is the wave (direction) vector of the transmitted wave.

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published paper for an example

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2.1.1 Boundary Conditions for TE Waves

At the interface where all three waves ~~emerge~~ ^{Fixed} simultaneously, a crucial boundary condition must be established to govern the relationship between their respective wave amplitudes. This boundary condition stipulates that the waves both incident upon and emerging from the plane of incidence should exhibit continuity and differentiability. This requirement is contingent upon the assumption that the interface is isotropic meaning that the physical properties of the material do not vary with different directions of measurement at the interface.

$$\vec{E}_r = E_{0r}^{\text{Fixed}} e^{i(\vec{k}_r \cdot \vec{r} - \omega_r t)} \quad (2.4)$$

In the context of our TE mode configuration (see figure 2.2), we can express the waves for the incident, reflected, and transmitted electric field components waves as follows:

$$\vec{E} = E \hat{y} e^{i(\vec{k} \cdot \vec{r} - \omega t)} \quad \vec{E}_r = E_r \hat{y} e^{i(\vec{k}_r \cdot \vec{r} - \omega_r t)} \quad \vec{E}_t = E_t \hat{y} e^{i(\vec{k}_t \cdot \vec{r} - \omega_t t)}$$

where E , E_r , and E_t denote the complex field amplitudes corresponding to the incident, reflected, and transmitted waves, respectively. These waves adhere to the boundary conditions, ensuring the continuity of electric field components parallel to the interface, as in:

$$E + E_r = E_t \text{ Fixed} \quad (2.5)$$

Using basic trigonometry and Maxwell's equations, we can derive the corresponding magnetic fields for the incident, reflected, and transmitted waves. The expressions for these fields, assuming a TE mode, are:

$$\begin{aligned} \vec{B} &= (B \cos(\theta \hat{x}) - B \sin(\theta \hat{z})) e^{i(\vec{k} \cdot \vec{r} - \omega t)} \\ \vec{B}_r &= (-B_r \cos(\theta_r \hat{x}) - B_r \sin(\theta_r \hat{z})) e^{i(\vec{k}_r \cdot \vec{r} - \omega_r t)} \\ \vec{B}_t &= (B_t \cos(\theta_t \hat{x}) - B_t \sin(\theta_t \hat{z})) e^{i(\vec{k}_t \cdot \vec{r} - \omega_t t)} \end{aligned}$$

where

- θ represents the angle of incidence (the angle between the incident wave's direction of propagation and the normal to the interface). See figure 2.2.
- θ_r represents the angle of reflection (the angle between the reflected wave's direction of propagation and the normal to the interface).
- θ_t represents the angle of transmission (the angle between the transmitted wave's direction of propagation and the normal to the interface).

As the magnetic field vector lies transversely to the plane of incidence, adherence to the boundary conditions necessitates the connection of parallel components of the magnetic field, as defined by:

$$B \cos(\theta) - B_r \cos(\theta) = B_t \cos(\theta_t) \quad (2.6)$$

I'm ^{Fixed} unsure what this means

parallel to what?

Here, it's important to note that $\theta = \theta_r$ according to the Law of Reflection. Equations 2.5 and 2.6 stand as two pivotal equations arising from the boundary conditions for TE waves. These equations, instrumental in determining R , serve as a critical foundation. Nevertheless, before we delve into the calculation of R through these boundary conditions, it is imperative to demonstrate the applicability of the same procedure to the transverse magnetic case.

2.1.2 Boundary Conditions for TM Waves

Another polarization mode for electromagnetic waves is known as the *transverse magnetic* (TM) polarization. In this mode, the magnetic field vector is oriented perpendicular to the plane of incidence, while the electric field vector lies transverse to the plane of incidence. Within the framework of our TM mode configuration, we can formulate the wave equations governing the electric field components of the incident, reflected, and transmitted waves as follows (see figure 2.3):

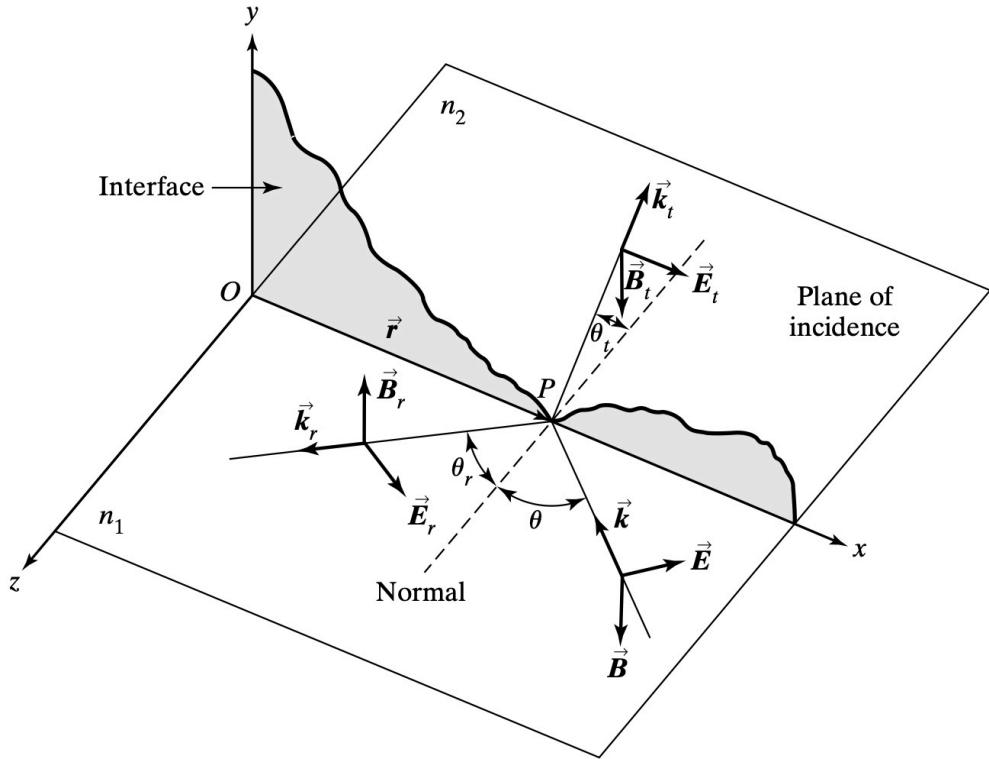


Figure 2.3: The transverse electric (TM) set-up. Source: [3]

$$\vec{E} = (E \cos(\theta \hat{x}) - E \sin(\theta \hat{z})) e^{i(\vec{k} \cdot \vec{r} - \omega t)}$$

$$\vec{E}_r = (E_r \cos(\theta_r \hat{x}) + E_r \sin(\theta_r \hat{z})) e^{i(\vec{k}_r \cdot \vec{r} - \omega t)}$$

$$\vec{E}_t = (E_t \cos(\theta_t \hat{x}) - E_t \sin(\theta_t \hat{z})) e^{i(\vec{k}_t \cdot \vec{r} - \omega t)}$$

Consequently, the magnetic field components can be written as (see figure 2.3):

$$\begin{aligned}\vec{B} &= -B\hat{y}e^{i(\vec{k} \cdot \vec{r} - \omega t)} \\ \vec{B}_r &= B_r\hat{y}e^{i(\vec{k} \cdot \vec{r} - \omega t)} \\ \vec{B}_t &= -B_t\hat{y}e^{i(\vec{k}_t \cdot \vec{r} - \omega t)}\end{aligned}$$

These equations adhere to the boundary conditions, guaranteeing the continuity of electric field components parallel to the interface, as demonstrated by:

$$-B + B_r = -B_t \quad (2.7)$$

$$E\cos(\theta) + E_r\cos(\theta) = E_t\cos(\theta_t) \quad (2.8)$$

2.1.3 Reflection and Transmission Coefficients

To determine the reflectance R and, consequently, the transmittance T , it is imperative to define the reflection (r) and transmission coefficients (t), as follows:

$$r \equiv \frac{E_r}{E} \quad (2.9)$$

and

$$t \equiv \frac{E_t}{E}. \quad (2.10)$$

However, it is essential to calculate these coefficients for both the Transverse Electric (TE) and Transverse Magnetic (TM) polarization cases. Recall that the electric (E) and magnetic (B) field amplitudes are interconnected through the following relation:

$$E = \nu B = \left(\frac{c}{n}\right) B \quad (2.11)$$

where:

- 
- E represents the amplitude of the electric field.
 - ν represents the phase velocity of the wave within the medium, $\nu = \frac{c}{n}$.
 - n is the refractive index of the medium.
 - c is the speed of light.

We can employ 2.11 to replace every occurrence of B in the boundary conditions with its corresponding E . Initiating with the TE case, remember the two vital boundary condition equations for the TE mode:

$$TE : \begin{cases} E + E_r = E_t \\ n_1 E \cos(\theta) - n_1 E_r \cos(\theta) = n_2 E_t \cos(\theta_t) \end{cases} \quad (2.12)$$

We can proceed to solve the system of equations above for the TE case to determine r_{TE} (by eliminating all instances of E_t). Introducing the concept of the *relative refractive index*, denoted as n and defined as $n \equiv \frac{n_2}{n_1}$ where n_1 is the refractive index of the initial medium through which the incident wave travels, and n_2 is the refractive index of the second medium into which the wave is transmitted.

We can derive:

$$r_{TE} = \frac{E_r}{E} = \frac{\cos(\theta) - n \cos(\theta_t)}{\cos(\theta) + n \cos(\theta_t)}. \quad (2.13)$$

By the Law of Refraction, $\sin(\theta) = n \sin(\theta_t)$, so we can eliminate θ_t by noting that:

$$n \cos(\theta_t) = n \sqrt{\cos^2(\theta_t)} = n \sqrt{1 - \sin^2(\theta_t)} = \sqrt{n^2 - \sin^2(\theta)} \quad (2.14)$$

Finally, our reflection coefficient is:

$$r_{TE} = \frac{E_r}{E} = \frac{\cos(\theta) - \sqrt{n^2 - \sin^2(\theta)}}{\cos(\theta) + \sqrt{n^2 - \sin^2(\theta)}} \quad (2.15)$$

Likewise, employing our boundary conditions for the TM case along with 2.11, we arrive at the reevaluated boundary conditions for the TM mode, which are as follows:

$$TM : \begin{cases} -n_1 E + n_1 E_r = -n_2 E_t \\ E \cos(\theta) + E_r \cos(\theta) = E_t \cos(\theta_t) \end{cases} \quad (2.16)$$

With this revised form of the boundary condition, we can compute r_{TM} in a manner similar to our determination of r_{TE} , resulting in:

$$r_{TM} = \frac{E_r}{E} = \frac{-n^2 \cos(\theta) + \sqrt{n^2 - \sin^2(\theta)}}{n^2 \cos(\theta) + \sqrt{n^2 - \sin^2(\theta)}} \quad (2.17)$$

Similarly, if we follow the same steps we did for evaluating r_{TM} and r_{TE} , we can subsequently figure out t_{TM} and t_{TE} (we now eliminate E_r instead of E_t). We find:

$$t_{TE} = \frac{E_t}{E} = \frac{2 \cos(\theta)}{\cos(\theta) + \sqrt{n^2 - \sin^2(\theta)}} \quad (2.18)$$

$$t_{TM} = \frac{E_t}{E} = \frac{2 n \cos(\theta)}{n^2 \cos(\theta) + \sqrt{n^2 - \sin^2(\theta)}} \quad (2.19)$$

not a
sentence
Fixed

Observing, for instance, the interconnection between t_{TE} and r_{TE} , it is evident that they share the same denominator, suggesting that one can be expressed in terms of the other. To establish the relationship between the two equations, one can subtract r_{TE} from t_{TE} , yielding 1. With some additional manipulation, the corresponding equation linking t_{TM} and r_{TM} can be derived. Ultimately, r can be expressed in terms of t as follows (see figure 2.3):

$$t_{TE} = 1 + r_{TE}$$

$$nt_{TM} = 1 - r_{TM}$$

The set of equations r_{TE} (2.15), r_{TM} (2.17), t_{TE} (2.18), and t_{TM} (2.19) constitutes the **Fresnel Equations**. These equations provide reflection and transmission coefficients that establish the connection between incident and reflected ~~energy~~ ^{Fixed} field amplitudes in any linear, isotropic, and homogeneous medium. The primary objective of the Fresnel equations is to facilitate the determination of reflectance R and transmittance T , which will be discussed in the subsequent subsection.

2.1.4 Reflectance and Transmittance

For the sake of energy conservation, it is imperative that, at a specific boundary, the power incident upon the boundary equals the combined power of both the reflected and transmitted energy at that boundary.

~~Fixed~~ ^{Waves}

$$P_i = P_r + P_t \quad (2.20)$$

We define reflectance R and transmittance T as:

$$R = \frac{P_r}{P_i} \quad (2.21)$$

$$T = \frac{P_t}{P_i} \quad (2.22)$$

Hence, R represents the proportion of reflected power relative to incident power, and transmittance T signifies the proportion of transmitted power in relation to incident power. This implies that 2.20 conforms to the well-known unity equation, under the assumption of zero absorption:

$$R + T = 1 \quad (2.23)$$

Power can be defined as the product of the irradiance (I) of the electromagnetic wave and the cross-sectional area (A) through which the wave passes, so 2.20 can be written as:

$$I_i A_i = I_r A_r + I_t A_t \quad (2.24)$$

To ascertain A_i , A_r , and A_t , it is essential to calculate the cross-sectional area covered by the incident, reflected, and transmitted electromagnetic waves, respectively. By employing trigonometric relationships, as illustrated in ^{2.4}, we can calculate the cross-sectional areas

~~Fixed~~ ^{Figure}

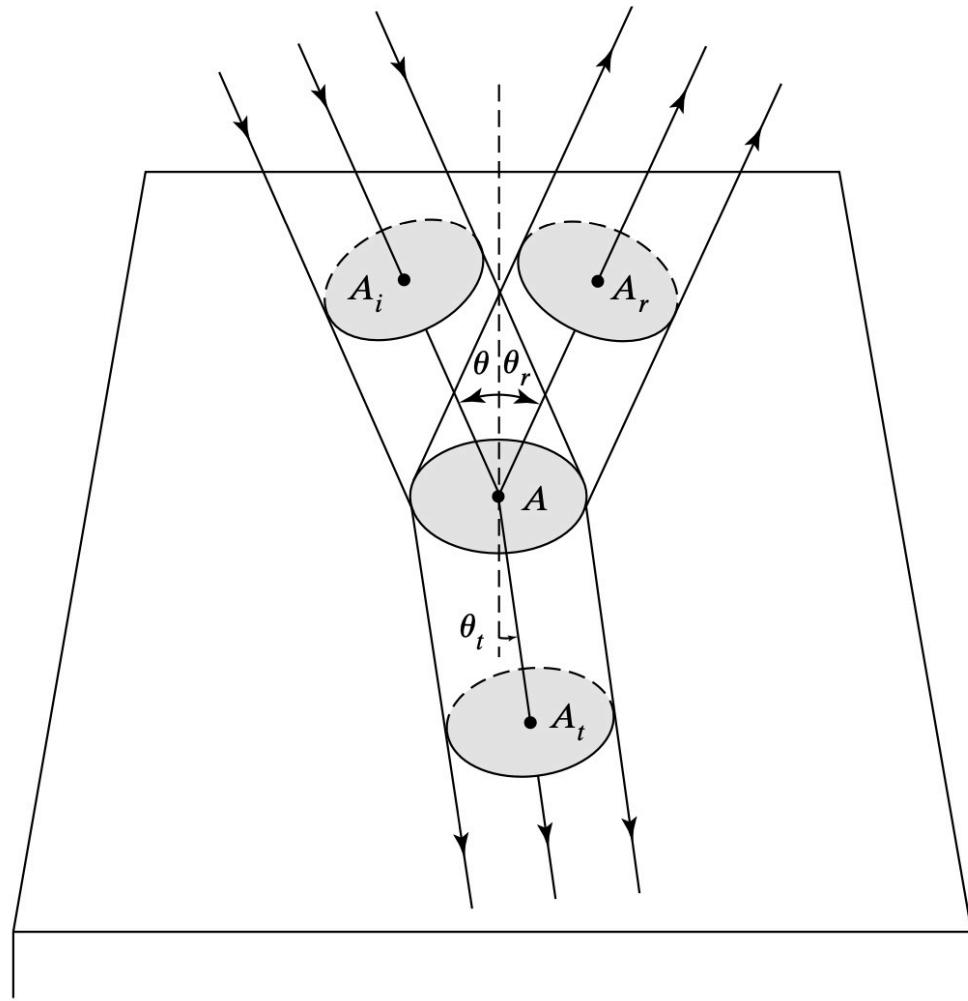


Figure 2.4: Cross sections of the incident, reflected, and transmitted beams. Source: [3]

encompassed by the incident, reflected, and transmitted waves and subsequently substitute them into the equation 2.24 to be:

$$I_i A \cos(\theta) = I_r A \cos(\theta) + I_t A \cos(\theta_t) \quad (2.25)$$

where A represents the cross-sectional area through which the wave propagates.

Using the relation between irradiance and electrical field amplitude,

$$I = E_0^2 \left(\frac{\varepsilon \nu}{2} \right) \quad (2.26)$$

where:

- I represents the irradiance, which is the power per unit area received by a surface
- E_0 is the amplitude of the electric field, indicating the maximum strength of the field.

Fixed

- ε denotes the permittivity of the medium.
- ν is the phase velocity of the electromagnetic wave in the medium.

and that $\nu_i = \nu_r$ and $\varepsilon_i = \varepsilon_r$ since they correspond to the same medium, we can rephrase the power balance equation to be:

$$E_{0i}^2 = E_{0r}^2 + E_{0t}^2 \left(\frac{\nu_t \varepsilon_t}{\nu_i \varepsilon_i} \right) \left(\frac{\cos(\theta_t)}{\cos(\theta)} \right) \quad (2.27)$$

where

- E_{0i} , E_{0r} , and E_{0t} represent the electric field amplitudes of the incident, reflected, and transmitted waves, respectively.
- ν_i and ν_r denote the phase velocities in the incident and reflected paths, which are equal since these paths are within the same medium.
- ε_i and ε_r are the permittivities of the medium for the incident and reflected waves, equal because they pertain to the same medium.
- ν_t and ε_t refer to the phase velocity and permittivity in the transmission path, respectively.
- The angles θ and θ_t are the angles of incidence and transmission, respectively.

The ratio $\frac{\nu_t \varepsilon_t}{\nu_i \varepsilon_i}$ is a complicated way of expressing the relative refractive index. Considering that $\mu_i = \mu_t = \mu_0$ for nonmagnetic materials, where μ_i and μ_t are the magnetic permeabilities of the incident and transmitted materials, and μ_0 is the magnetic permeability of free space. The relationship, $\nu^2 = \frac{1}{\mu \varepsilon}$ further elucidates the link between phase velocity, magnetic permeability, and permittivity. Hence, we derive:

$$\frac{\nu_t \varepsilon_t}{\nu_i \varepsilon_i} = \frac{\nu_i^2 \mu_i \nu_t}{\nu_t^2 \mu_t \nu_i} = \frac{\nu_i}{\nu_t} = n \quad (2.28)$$

This transformation streamlines the comprehension of how waves interact with various media by introducing the concept of the relative refractive index, n . This index represents the ratio of the phase velocities of the wave as it moves from the incident to the transmitted medium, thus quantifying the change in phase speed resulting from the passing of the wave between these two media.

Thus we can include n in the power balance equations:

$$E_{0i}^2 = E_{0r}^2 + n E_{0t}^2 \left(\frac{\cos(\theta_t)}{\cos(\theta)} \right) \quad (2.29)$$

Dividing through by E_{0i}^2 , we get:

$$1 = r^2 + n t^2 \left(\frac{\cos(\theta_t)}{\cos(\theta)} \right) \quad (2.30)$$

Reflectance (R) and transmittance (T) quantify the fractions of incident electromagnetic power that are reflected and transmitted across a boundary, respectively. These concepts are grounded in the principle of energy conservation, which, in the context of electromagnetic waves at an interface, is expressed as:

$$R = \frac{P_r}{P_i} = \frac{I_r}{I_i} = \left(\frac{E_{0r}}{E_0 i} \right)^2 = r^2 \quad (2.31)$$

where in the last step, I have utilized equation 2.26. The term r^2 is introduced as a shorthand notation for this ratio, encapsulating the reflectance in terms of the electric field amplitude ratios. ~~Fixed~~

Looking at equation 2.23, we gather that:

$$T = nt^2 \left(\frac{\cos(\theta_t)}{\cos(\theta)} \right) \quad (2.32)$$

Fixed It's worth noting that T is not merely t^2 , as we must consider the altered speed of the electromagnetic wave when it enters a medium with a different refractive index. This change in speed impacts the rate of energy propagation and, consequently, the power of the beam.

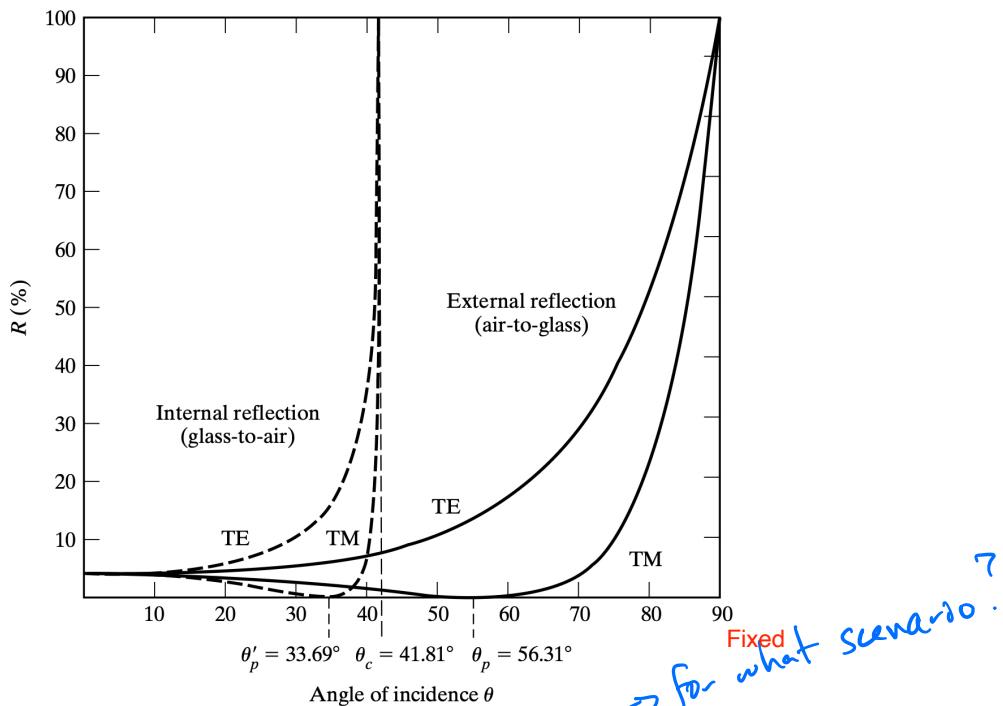


Figure 2.5: Reflectance vs Angle of Incidence. Source: [3]

Figure 2.5 illustrates how the reflectance R of light varies with the angle of incidence θ for two different polarizations of light we've looked at, the transverse electric (TE) and transverse

magnetic (TM). The graph is split into two regions that represent different physical scenarios of light interaction with a boundary: internal reflection (glass-to-air) and external reflection (air-to-glass).

The critical angle, denoted as θ_c , represents the specific angle of incidence at which a refracted light ray grazes the interface between two media. At this angle, the ray runs parallel to the boundary, without penetrating into the second medium or retreating into the first. When the angle of incidence surpasses θ_c , a scenario of total internal reflection unfolds: all the incident light is entirely reflected into the originating denser medium, and no light is refracted into the less dense medium, such as air. This results in the graph indicating a reflectance of 100% for both TE and TM polarization states in situations of glass-to-air transition. This effect is exclusive to internal reflection; conversely, when light travels from a less dense to a denser medium, it is almost always refracted into the denser medium, a behavior consistent across all incidence angles.

Brewster's Angles, marked by vertical dashed lines at 56.31° for external reflection (air-to-glass) and 33.69° for internal reflection (glass-to-air), correspond to the specific angles of incidence where the reflectance reaches a minimum for TM-polarized light in external reflection and internal reflection respectively. At these angles, the reflected intensity is significantly reduced for the respective polarization states.

n is a function of the wavelength (λ) of light. Consequently, both reflectance (R) and transmittance (T) are wavelength-dependent. This wavelength dependence is particularly relevant to the scope of this work as it influences the behavior of light across different spectra. The refractive index's variation with wavelength means that the critical and Brewster's angles, and therefore the efficiency of reflection and transmission, will vary with the wavelength of the incident light, which has significant implications for applications that rely on precise control of light propagation, such as in optical filters and coatings.

2.2 Multilayer films

In the development of Passive Daytime Radiative Cooling Devices (PDRCs), we intend to employ multiple stacks comprising diverse materials with distinct refractive indices, arranged in layers to enhance the overall reflection coefficient and consequently increase reflectance. It is crucial to comprehend the electromagnetic wave interaction and optimal layer stacking physics to maximize reflectance. We again need a rigorous consideration of boundary conditions as dictated by Maxwell's equations.

The derivation below follows that given in ^{Fixed} [cite].

2.2.1 Transfer Matrix

At the heart of analyzing how electromagnetic waves interact with thin film structures lies a powerful tool, the transfer matrix. We shall spend most of this section elucidating the process by which we derive a general form of the transfer matrix, M , for arbitrary thin films. This 2×2 matrix is instrumental in establishing a quantitative link between the electric and magnetic fields at one interface. By introducing the transfer matrix, we enable the

prediction of wave behavior across various interfaces encountered within a multilayer stack. The transfer matrix not only does this for ~~double~~^{two} ~~Fixed~~ layers (single interfaces) but it also offers a scalable approach to tackling the dynamics of multiple layers.

Consider the set-up ~~above~~^{Fixed} where the electric field amplitude is perpendicular to the plane of incidence (a transverse electric (TE) wave).

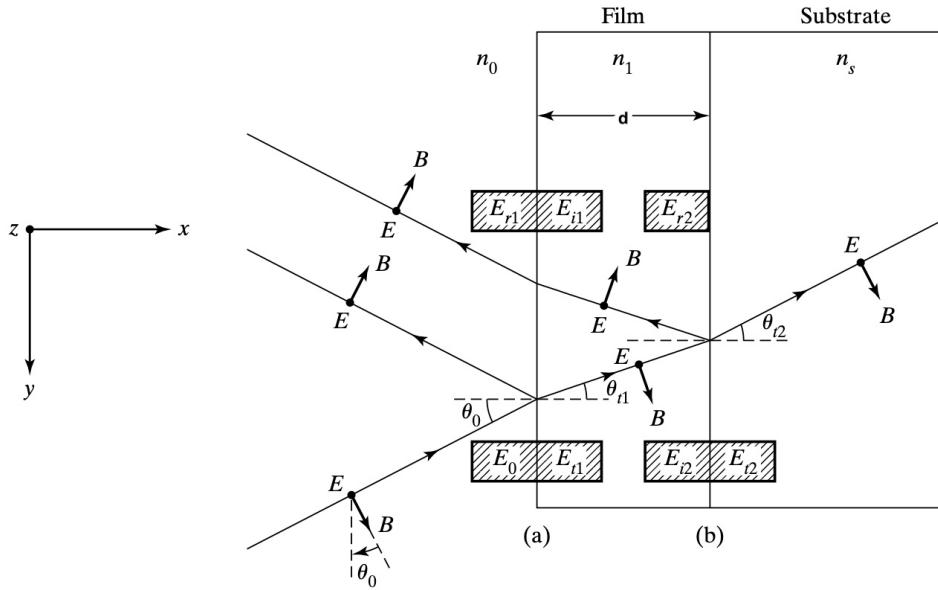


Figure 2.6: Multilayer Film Set-Up. Source: [3]

A TE wave propagates from air, reaching the air-film boundary. A fraction of the wave is transmitted, while another portion is reflected. The transmitted portion encounters the film-substrate interface, where part is further transmitted, and another part is reflected. This reflected component then becomes incident on the film-air interface, leading to transmission and reflection, repeating the process. This scenario can be extrapolated to involve multiple films. It is important to note that the film and substrate possess distinct refractive indices, denoted as n_1 and n_s respectively.

To capture the complex interaction between reflected and transmitted beams, we employ a systematic notation, as seen in the provided setup image. For example, E_{r1} denotes the aggregate of multiply reflected beams at interface (a), while E_{i2} signifies the sum of multiple incident beams at interface (b), directed toward the substrate.

Continuing with our established assumptions, the film is presumed to be homogeneous, (has the same properties at every point) and isotropic (the physical properties do not differ regardless of the direction or orientation in which it is examined). Furthermore, we hold that the film's thickness is comparable to the wavelength of the incident light because it allows for the film to be a medium where constructive and destructive interference patterns emerge, significantly affecting how light is transmitted and reflected by the film. Such interference

phenomena are fundamental in the function of anti-reflective coatings and other optical devices, where precise control of light behavior is required.

Good!
This discussion
might be
useful in
the Fresnel
derivation
above when
you first
introduce
the boundary
conditions.

The boundary conditions derived from Maxwell's equations stipulate that the tangential components of the electric and magnetic fields in plane waves must exhibit continuity across the interface, remaining equal on both sides. It is essential to recognize that while the electric field component is tangential to the interfaces everywhere, the orthogonal nature of the magnetic field vector to the electric field vector necessitates the determination of corresponding tangential magnetic field vectors. When we refer to a field component as being "tangential," we imply that it lies parallel to the plane of the interface, skimming along it without crossing. This is similar to drawing a line that touches a circle at just one point, running along its edge without entering its interior.

This endeavor to identify and equate tangential components of both electric and magnetic fields across an interface lays the foundation for deriving expressions that describe the electromagnetic field behavior across different media. It allows us to formulate relationships that link the characteristics of the fields on one side of the interface to those on the other side, essential for predicting how electromagnetic waves propagate through varying materials:

$$E_a = E_0 + E_{r1} = E_{t1} + E_{i1} \quad (2.33)$$

$$E_b = E_{i2} + E_{r2} = E_{t2} \quad (2.34)$$

To determine the tangential magnetic field vectors, trigonometry can be applied to establish the following boundary conditions:

$$B_a = B_0 \cos(\theta_0) - B_{r1} \cos(\theta_0) = B_{t1} \cos(\theta_{t1}) - B_{i1} \cos(\theta_{t1}) \quad (2.35)$$

$$B_b = B_{i2} \cos(\theta_{t1}) - B_{r2} \cos(\theta_{t1}) = B_{t2} \cos(\theta_{t2}) \quad (2.36)$$

← We can build on equation 2.11 by expressing B in terms of E as:

$$B = n\sqrt{\epsilon_0 \mu_0} E \quad (2.37)$$

and since $c = \frac{1}{\sqrt{\epsilon_0 \mu_0}}$, we can rewrite 2.35 and 2.36 employing 2.37:

$$B_a = \gamma_0 (E_0 - E_{r1}) = \gamma_1 (E_{t1} - E_{i1}) \quad (2.38)$$

$$B_b = \gamma_1 (E_{i2} - E_{r2}) = \gamma_s E_{t2} \quad (2.39)$$

where

$$\gamma_0 \equiv n_0 \sqrt{\epsilon_0 \mu_0} \cos(\theta_0), \quad (2.40)$$

$$\gamma_1 \equiv n_1 \sqrt{\epsilon_0 \mu_0} \cos(\theta_{t1}) \quad (2.41)$$

$$\gamma_s \equiv n_s \sqrt{\epsilon_0 \mu_0} \cos(\theta_{t2}) \quad (2.42)$$

E_{i2} differs from E_{t1} solely due to a phase difference δ arising from a single traversal of the film. The optical path length linked with one traversal is denoted as $\Delta_1 = n_1 d \cos(\theta_{t1})$. The optical path length represents the distance that light "perceives" it has covered in a medium and can be quantified in terms of the number of cycles the light beam has undergone within that specific medium. It is inherently proportional to the refractive index and the geometric length of the medium through which light is propagating.

We can express the phase difference that develops due to one traversal of the film as:

$$\delta = k_0 \Delta_1 = \left(\frac{2\pi}{\lambda_0} \right) n_1 d \cos(\theta_{t1}) \quad (2.43)$$

Therefore we can express the pair electric field sums E_{i2} & E_{t1} and E_{i1} & E_{r2} as follows:

$$\begin{aligned} E_{i2} &= E_{t1} e^{-i\delta} \\ E_{i1} &= E_{r2} e^{-i\delta} \end{aligned}$$

and consequently rephrase 2.34 and 2.39 as:

$$E_b = E_{t1} e^{-i\delta} + E_{i1} e^{i\delta} = E_{t2} \quad (2.44)$$

$$B_b = \gamma_1 (E_{t1} e^{-i\delta} - E_{i1} e^{i\delta}) = \gamma_s E_{t2} \quad (2.45)$$

Looking at the middle terms, we can solve for E_{t1} and E_{i1} in terms of E_b and B_b .

$$E_{t1} = \left(\frac{\gamma_1 E_b + B_b}{2\gamma_1} \right) e^{i\delta} \quad (2.46)$$

$$E_{i1} = \left(\frac{\gamma_1 E_b - B_b}{2\gamma_1} \right) e^{-i\delta} \quad (2.47)$$

Utilizing the Euler identities $2i\sin(\delta) \equiv e^{i\delta} - e^{-i\delta}$ and $2\cos(\delta) \equiv e^{i\delta} + e^{-i\delta}$, we can use the equations 2.46 and 2.47 and substitute them into 2.33 and 2.39 to have both:

$$E_a = E_b \cos(\delta) + B_b \left(\frac{i\sin(\delta)}{\gamma_1} \right) \quad (2.48)$$

$$B_a = E_b (i\gamma_1 \sin(\delta)) + B_b \cos(\delta) \quad (2.49)$$

which we can rewrite in matrix notation as:

$$\begin{bmatrix} E_a \\ B_a \end{bmatrix} = \begin{bmatrix} \cos(\delta) & \frac{i\sin(\delta)}{\gamma_1} \\ i\gamma_1 \sin(\delta) & \cos(\delta) \end{bmatrix} \begin{bmatrix} E_b \\ B_b \end{bmatrix}$$

This notation introduces the insightful 2×2 matrix that establishes a connection between the electric and magnetic fields at one interface and those at the subsequent interface. Termed the *transfer matrix*, we will refer to this significant matrix in a generic manner as:

$$M = \begin{bmatrix} m_{11} & m_{12} \\ m_{21} & m_{22} \end{bmatrix}$$

~~Fixed~~ Expanding this framework, we can extend the consideration to the stacking of multiple films, leading to the emergence of multiple interfaces. In a more general context, for an arbitrary number N of layers:

$$\begin{bmatrix} E_a \\ B_a \end{bmatrix} = M_1 M_2 M_3 \dots M_N \begin{bmatrix} E_N \\ B_N \end{bmatrix}$$

with the multiplication of individual transfer matrices representing the entire multilayer stack in the order in which the beam encounters them.

Returning to the comprehensive form of the transfer matrix presented above, we can substitute E_a and B_a with the middle segments of equations 2.33 and 2.38. Similarly, E_b and B_b can be replaced with the rightmost segments of equations 2.44 and 2.45, resulting in:

$$\begin{bmatrix} E_0 + E_{r1} \\ \gamma_0(E_0 - E_{r1}) \end{bmatrix} = \begin{bmatrix} \cos(\delta) & \frac{i \sin(\delta)}{\gamma_1} \\ i \gamma_1 \sin(\delta) & \cos(\delta) \end{bmatrix} \begin{bmatrix} E_{t2} \\ \gamma_s E_{t2} \end{bmatrix} = \begin{bmatrix} m_{11} & m_{12} \\ m_{21} & m_{22} \end{bmatrix} \begin{bmatrix} E_{t2} \\ \gamma_s E_{t2} \end{bmatrix}$$

Recall $r \equiv \frac{E_{r1}}{E_0}$ and $t \equiv \frac{E_{t2}}{E_0}$ (the reflection and transmission coefficients, respectively). By matrix multiplication, we can simplify the resulting equation to obtain:

$$1 + r = m_{11}t + m_{12}\gamma_s t \quad (2.50)$$

$$\gamma_0(1 - r) = m_{21}t + m_{22}\gamma_s t \quad (2.51)$$

Solving for r by solving the system of linear equations above, we get

$$r = \frac{\gamma_0 m_{11} + \gamma_0 \gamma_s m_{12} - m_{21} - \gamma_s m_{22}}{\gamma_0 m_{11} + \gamma_0 \gamma_s m_{12} + m_{21} + \gamma_s m_{22}} \quad (2.52)$$

2.2.2 Antireflecting Films

To gain deeper insights into optimizing reflectance, ~~let's~~ explore the characteristics conducive to anti-reflection using the structure shown in figure 2.7. Employing a double layer of quarter-wave-thickness films allows for the attainment of minimal reflectance at a specific wavelength. We will examine the simplest scenario for normal incidence, where all angles of incidence, reflection, and refraction are zero.

At normal incidence ($\theta = 0$) and for a film of quarter-wave thickness where $d = \frac{\lambda}{4} = \frac{\lambda_0}{4n_1}$, the phase difference, denoted as δ , becomes:

$$\delta = \left(\frac{2\pi}{\lambda_0} \right) n_1 d \cos(\theta_{t1}) = \frac{\pi}{2} \quad (2.53)$$

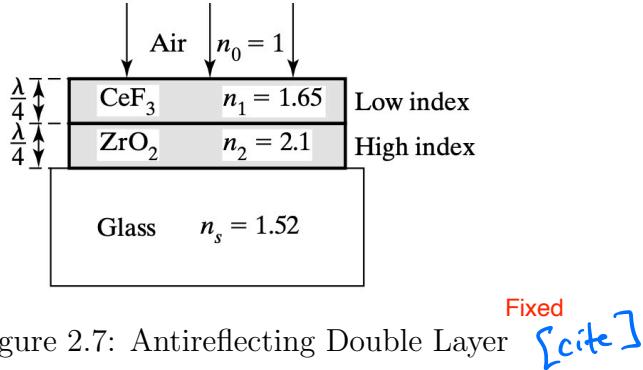


Figure 2.7: Antireflecting Double Layer [cite]

so that $\cos(\delta) = 0$ and $\sin(\delta) = 1$. With the phase difference established, the corresponding transfer matrix is expressed as:

Should this be a subscript?

$$M = \begin{bmatrix} \cos(\delta) & \frac{i \sin(\delta)}{\gamma_1} \\ i \gamma_1 \sin(\delta) & \cos(\delta) \end{bmatrix} = \begin{bmatrix} 0 & \frac{i}{\gamma_1} \\ i \gamma_1 & 0 \end{bmatrix}$$

The term γ_1 is related to the refractive index n_1 of the first film layer from the top, Cerium Fluoride (CeF₃), as depicted in figure 2.7. According to our earlier definitions (equation 2.41), $\gamma_1 \equiv n_1 \sqrt{\epsilon_0 \mu_0} \cos(\theta_{t1})$ where n_1 is the refractive index of the CeF₃ layer and θ_{t1} is the angle of transmission within this first layer. γ_1 is related to the wave impedance within the film relative to the impedance of free space, modified by the cosine of the angle of transmission.

Given two quarter-wave thickness films, the individual transfer matrices are expressed as:

$$M_1 = \begin{bmatrix} 0 & \frac{i}{\gamma_1} \\ i \gamma_1 & 0 \end{bmatrix}$$

and

$$M_2 = \begin{bmatrix} 0 & \frac{i}{\gamma_2} \\ i \gamma_2 & 0 \end{bmatrix}$$

The overall transfer matrix for the system is then the product of M_1 and M_2 :

$$M = M_1 \cdot M_2 = \begin{bmatrix} 0 & \frac{i}{\gamma_1} \\ i \gamma_1 & 0 \end{bmatrix} \begin{bmatrix} 0 & \frac{i}{\gamma_2} \\ i \gamma_2 & 0 \end{bmatrix}$$

Calculating the product of these matrices, we obtain the overall transfer matrix for this double layer:

$$M = \begin{bmatrix} -\frac{\gamma_2}{\gamma_1} & 0 \\ 0 & -\frac{\gamma_1}{\gamma_2} \end{bmatrix}$$

Equation 2.52, the reflection coefficient, then simplifies to:

$$r = \frac{\gamma_2^2 \gamma_0 - \gamma_s \gamma_1^2}{\gamma_2^2 \gamma_0 + \gamma_s \gamma_1^2} \quad (2.54)$$

The reflectance, R , is simply $|r|^2$ and using equations 2.40 through 2.42, we can calculate R to be

$$R = \left(\frac{n_0 n_2^2 - n_s n_1^2}{n_0 n_2^2 + n_s n_1^2} \right)^2 \quad (2.55)$$

Zero reflectance is then expected when the numerator is zero

$$\begin{aligned} n_0 n_2^2 &= n_s n_1^2 \\ \frac{n_2^2}{n_1^2} &= \frac{n_s}{n_0} \end{aligned}$$

so that

$$\frac{n_2}{n_1} = \sqrt{\frac{n_s}{n_0}} \quad (2.56)$$

For a glass substrate with a refractive index of $n_s = 1.52$ and incidence from air with $n_0 = 1$, the optimal ratio of the refractive indices for the two films should be $\frac{n_2}{n_1} = \sqrt{1.52} \approx 1.23$. However, achieving near-zero reflectance is challenging over the broader region of the visible spectrum because the refractive index of each material in the layers vary with wavelength.

In the visible spectrum, which ranges approximately from 400 to 700 nm, each wavelength interacts differently with the optical materials based on their refractive index at that specific wavelength. The optimal thickness and refractive index that reduce reflectance for one wavelength will not necessarily be optimal for another due to this dispersion. Furthermore, the phase change upon reflection at the boundary of a medium also varies with wavelength, which affects the constructive and destructive interference patterns needed for anti-reflective properties.

Practically, this requires a careful design of the coating's material properties and thicknesses to achieve broader-wavelength anti-reflection. This design must account for the wavelength-dependent change in refractive indices and the cumulative effect of multiple reflections and transmissions at different interfaces within the multilayer stack, which becomes increasingly complex as the wavelength range widens, hence the difficulty of zero reflectance across the broader region over the visible spectrum. This dependence of refractive index on λ introduces a level of complexity that is difficult to treat analytically and is one of the motivations for using *consol* in this work.

2.2.3 High-Reflectance Layers.

In the previous section, we established that to optimize for anti-reflectance, we stack layers in the order of air-low index-high index-substrate. Conversely, to optimize for high-reflectance, we follow the opposite order: air-high index-low index-substrate. A set of double layers designed to enhance reflectance is referred to as a *high-reflectance stack* or *dielectric mirror*.

The transfer matrix for the two films in the order of high index-low index is

$$M_{HL} = \begin{bmatrix} 0 & \frac{i}{\gamma_H} \\ i\gamma_H & 0 \end{bmatrix} \begin{bmatrix} 0 & \frac{i}{\gamma_L} \\ i\gamma_L & 0 \end{bmatrix} = \begin{bmatrix} -\frac{\gamma_L}{\gamma_H} & 0 \\ 0 & -\frac{\gamma_H}{\gamma_L} \end{bmatrix}$$

I wonder if this paragraph would make a good conclusion to Ch 2, setting up the transition to Ch 3?

For N similar double layers, we obtain $M = (M_{HL})^N$. In matrix form,

$$M = \begin{bmatrix} 0 & \frac{i}{\gamma_L} \\ i\gamma_L & 0 \end{bmatrix}^N = \begin{bmatrix} \left(-\frac{\gamma_L}{\gamma_H}\right)^N & 0 \\ 0 & \left(-\frac{\gamma_H}{\gamma_L}\right)^N \end{bmatrix}$$

Considering normal incidence and using equations 2.40 through 2.42,

$$\frac{\gamma_L}{\gamma_H} = \frac{n_L}{n_H}. \quad (2.57)$$

Equation 2.52, the reflection coefficient, then simplifies to:

$$r = \frac{n_0 \left(\frac{-n_L}{n_H}\right)^N - n_s \left(\frac{-n_H}{n_L}\right)^N}{n_0 \left(\frac{-n_L}{n_H}\right)^N + n_s \left(\frac{-n_H}{n_L}\right)^N} \quad (2.58)$$

To get R , we calculate $|r|^2$ to get

$$R = \left[\frac{\left(\frac{n_0}{n_s}\right) \left(\frac{n_L}{n_H}\right)^{2N} - 1}{\left(\frac{n_0}{n_s}\right) \left(\frac{n_L}{n_H}\right)^{2N} + 1} \right]^2 \quad (2.59)$$

Maximum (100%) reflectance is either achieved when:

~~Fixed~~ When N (the number of layer pairs) approaches infinity.

~~Fixed~~ When $\frac{n_L}{n_H}$ approaches zero. For example, when this ratio is 0.5 and there are 3 layers, a reflectance of 95.97 is achieved.

Since the optimal reflectance is obtained with the smallest ratio of $\frac{n_L}{n_H}$, high-reflectance stacks can be constructed using alternating layers of MgF_2 ($n_L = 1.38$) and ZnS ($n_H = 2.35$) or TiO_2 ($n_H = 2.40$). For example, employing alternating double layers of MgF_2 and ZnS can achieve a reflectance of 99.95% with a layer count (N) of 8.

In this chapter, we have meticulously explored the theoretical underpinnings of PDRCs, delving into the Fresnel Equations, the boundary conditions for TE and TM polarization states, and the complex behavior of multilayer films. Our investigation highlighted the delicate balance between film thickness, the coherence of light, and the manipulation of refractive indices necessary to realize passive cooling without external energy. Moreover, the transfer matrix is a formidable mathematical apparatus, enabling us to analyze the traversal of light through successive film layers. The knowledge acquired is instrumental for advancing PDRC technologies that promise to redefine sustainable energy practices. These strategies leverage essential physical laws to confront the urgent environmental and energy demands of our times.

Excellent job on this chapter.
Note that I have not checked your equations for typos, though - you should carefully double-check them against the book.

Chapter 3

Computational Methods

Cite your source here. Is this sufficiently rephrased so as not to be plagiarized?

Fixed COMSOL Multiphysics™ is a comprehensive software suite for finite element analysis, solving, and simulation across a wide array of physics and engineering disciplines, particularly focusing on coupled phenomena and multi-physics interactions. It enables the creation and simulation of physics-based models and applications in an intuitive, interactive workspace.

COMSOL software supports a broad spectrum of applications, from electromagnetics and structural mechanics to acoustics, fluid dynamics, heat transfer, and chemical engineering. COMSOL features an extensive array of modules, including the Wave Optics module for optical applications simulations like wave propagation in fiber optics and photonics, the Semiconductor module for the analysis of semiconductor devices such as diodes and transistors, and the AC/DC module for examining electric and magnetic fields in static and low-frequency scenarios.

For my thesis, which involves analyzing ray paths in optical systems, I will be utilizing the Ray Optics module.

3.1 The COMSOL Modelling Workflow

Briefly the modelling workflow consists of the following steps, if you start a model completely from scratch:

1. **Initialization of the Model Environment:** Preparing the foundational settings for the simulation.
2. **Geometry Construction:** Designing the model's physical layout.
3. **Material Property Specification:** Assigning specific materials to the model components.
4. **Physics Boundary Conditions Definition:** Establishing the physical constraints and conditions.
5. **Mesh Generation:** Creating the computational grid over the model.

Fixed *Illustrated in Figure 3.1,*

6. **Simulation Execution:** Performing the actual computational analysis.
7. **Results Post-Processing:** Analyzing and visualizing the simulation outcomes.

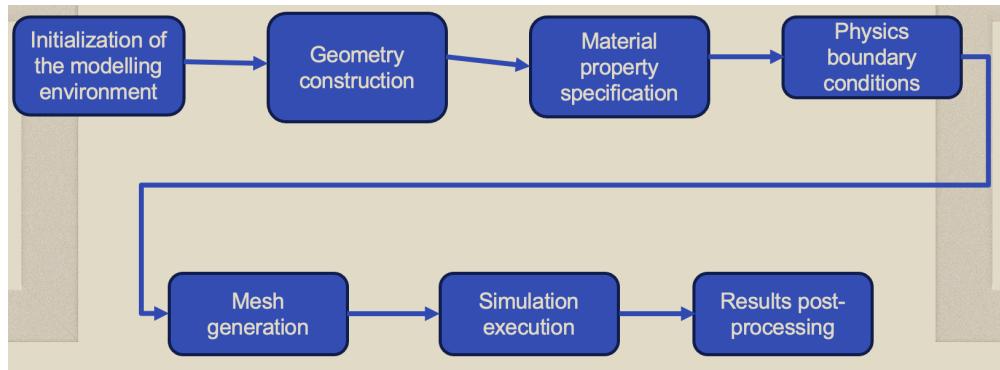


Figure 3.1: The modeling workflow.

In this context, I will demonstrate this workflow first through a simple project, illustrating COMSOL's robust capabilities and the straightforwardness of applying this workflow across various projects. It's important to recognize that this workflow remains consistent across different physics simulations, which will guide the development of subsequent models for my thesis.

following the sample tutorial given in ~ [cite] Fixed

This demonstration will focus on simulating Joule heating within a busbar, a process where electric current passing through metal results in heating due to electrical resistance.

The busbar in question will incorporate three titanium bolts, with electrical current flowing from a single bolt on one end to a pair of bolts at the opposite end. The objective is to observe the resulting temperature distribution as the busbar undergoes heating.

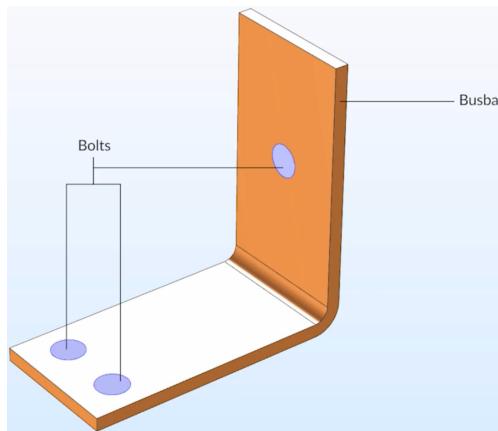
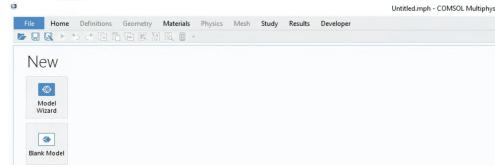


Figure 3.2: The busbar model. Source: [2]

Again, please double check throughout Ch 3 that your wording/instructions are sufficiently distinct from the tutorial you followed.

3.1.1 Setting Up the Model Environment



Can you make all these figures bigger?
They are hard to see -

Figure 3.3: Set-up Screen. Source: [2]

Upon launching COMSOL, you are presented with an option to initiate a new project: selecting a Blank Model for starting anew or opting for the Model Wizard for a guided setup. The Model Wizard is advisable for its structured selection process integral to configuring your model. Conversely, choosing a Blank Model grants immediate access to the COMSOL desktop interface, bypassing the preliminary selections required by the Model Wizard.

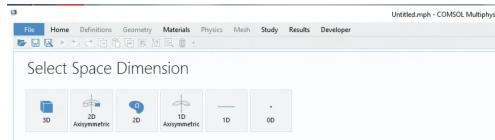


Figure 3.4: Select space dimension screen. Source: [2]

Since our busbar model requires a three-dimensional framework, we select the “3D” option.

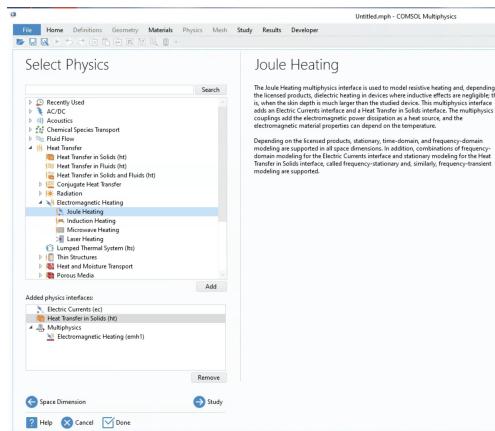


Figure 3.5: Select physics screen. Source: [2]

Subsequently, we arrive at a prompt to select the desired physics for the model. In our case, we opt for Joule heating. In another scenario, for instance, if one's project was on thin-film fluid flow, navigating to the “Fluid Flow” node and selecting “Thin-Film Flow”

would be the procedure. Adding “Joule Heating” automatically associates it with related physics categories like “Electric Currents (ec)” and “Heat Transfer in Solids (ht),” indicating that “Joule Heating” encompasses a composite interface with multiple physics elements integrated.

The selection of physics for exploration significantly hinges on the project’s specific requirements and the level of access provided by your COMSOL subscription. For example, acquiring the “Ray Optics” module necessary for my thesis entailed a waiting period due to initial subscription limitations. Therefore, early verification of the required physics for your project is recommended.

To proceed, we select “Study”.

modules
Fixed

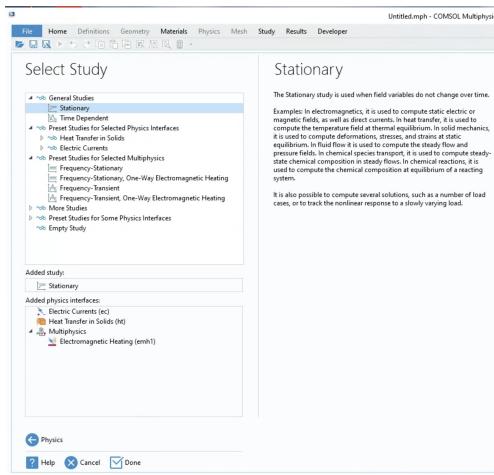


Figure 3.6: Select study screen. Source: [2]

Upon accessing this interface, we are presented with an assortment of study options, determined by our earlier selections regarding the model’s spatial dimensions and physics. For our busbar analysis, the “Stationary” study suffices, given its static nature. After selecting “Done”, we transition to the COMSOL desktop environment.

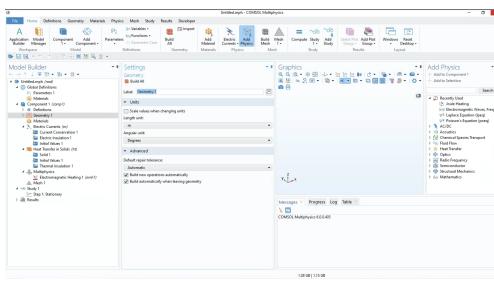


Figure 3.7: The COMSOL desktop. Source: [2]

The COMSOL desktop serves as the central hub for constructing and engaging with your model from the ground up. It is designed with an intuitive interface, featuring buttons and

ribbons aligned with the modeling process steps. Within the Model Builder window, you'll find the modeling hierarchy, where you have the capability to define your model's dimensions and select specific physics equations for adherence. Additionally, the layout of the COMSOL desktop windows can be customized to suit your preferences.

3.1.2 Building the Geometry

When developing the geometry of your model, multiple pathways are available. You may opt to utilize the drawing tools or geometric shapes provided by COMSOL or import a geometry from an external source.

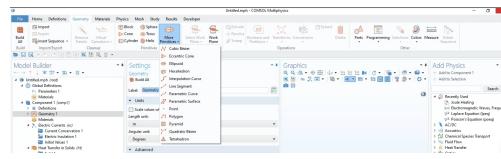


Figure 3.8: Add geometry button selection. Source: [2]

For this instance, given that the busbar geometry already exists within COMSOL's comprehensive library of pre-configured geometries, accessing this resource simplifies the process. This is achieved by navigating to the File menu and selecting Application Libraries.

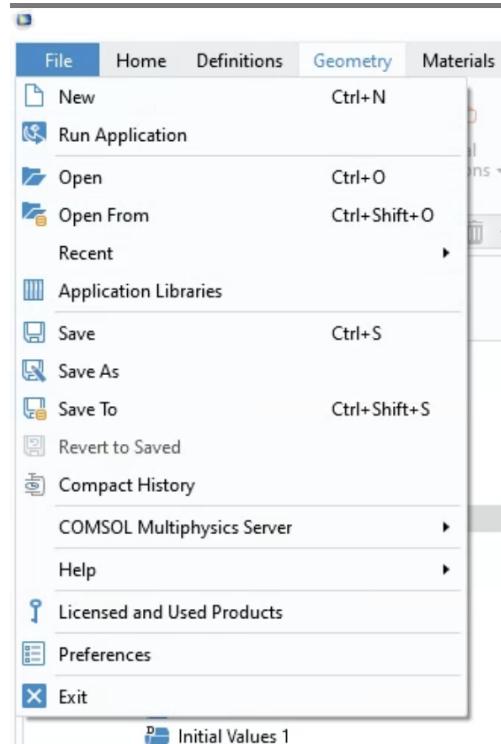


Figure 3.9: Application libraries selection. Source: [2]

Within the Application Libraries, proceed to the COMSOL Multiphysics section, locate the 'busbar_geom' file, open it, and proceed to save your project accordingly.

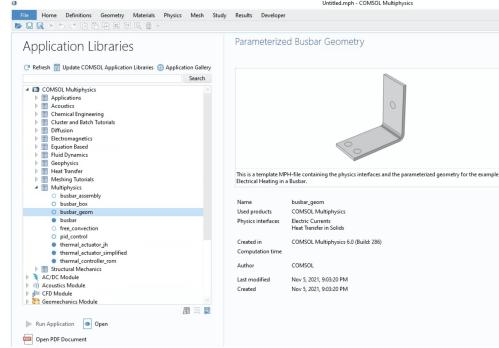


Figure 3.10: Application libraries. Source: [2]

The geometry will now appear in the Graphics window, allowing for zoom and rotational exploration to closely inspect the model. The alteration in the Geometry node within the Model Builder window reflects the sequence of actions executed to construct the busbar geometry. On closer inspection, it becomes evident that specific parameters were employed in crafting this geometry. These parameters are accessible within the "Parameters 1" node, which contains a table detailing the adjustable parameters utilized in the geometry's creation.

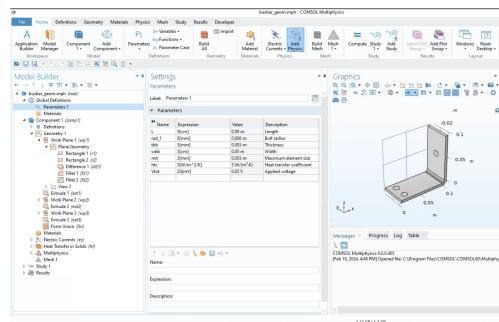


Figure 3.11: COMSOL desktop after adding the busbar geometry. Source: [2]

3.1.3 Specifying the Material Properties

Our busbar comprises various components each requiring distinct materials, necessitating batch assignment of materials to selected parts rather than individual allocation. To commence, access the Definitions tab and opt for the Explicit selection.

Following this, in the Graphics window, we identify and select the bolts, ensuring both their front and back sides are chosen due to their differing material composition from the busbar's main body.

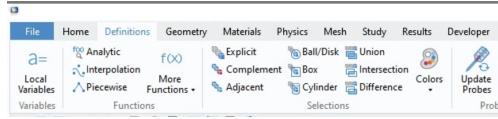


Figure 3.12: Explicit choice from the Definitions tab. Source: [2]

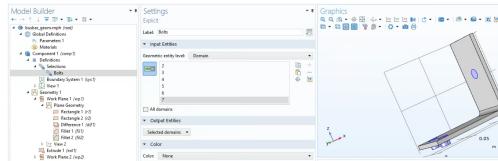


Figure 3.13: Adding bolt selections. Source: [2]

Proceeding to material assignment, we move to the Materials ribbon and select the Add Materials feature. This action opens the “Add Material” window, offering a selection of materials to apply to the chosen components.

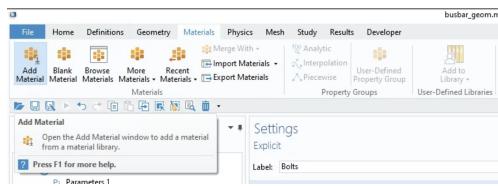


Figure 3.14: Add Materials option from Materials tab. Source: [2]

Select the Built-In option and opt for Titanium (for the bolts) and Copper as the materials.

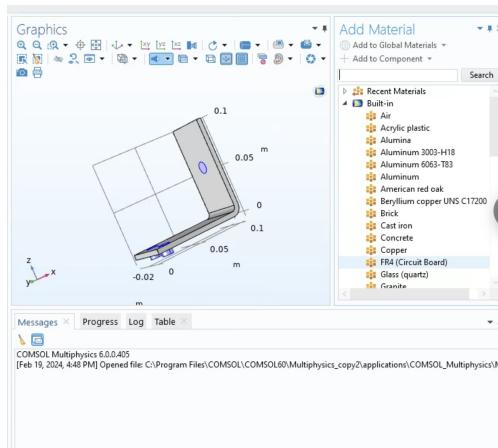


Figure 3.15: Add Materials window. Source: [2]

The earlier defined selection for Bolts proves useful now as we assign Titanium to each bolt by selecting the Bolts option, as illustrated below.

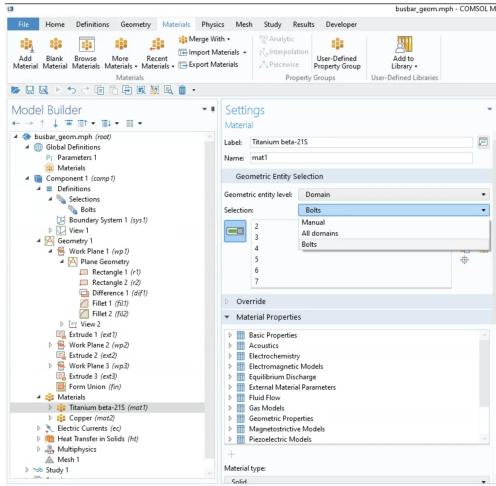


Figure 3.16: Assigning some bolts to be made of Titanium. Source: [2]

Every time a material is added, and consequently ~~Fixed~~ a material node is introduced in the Model Builder window, a table appears in the Settings window under Material Contents. This table details the physical properties of the respective materials, including Electrical Conductivity, Heat Capacity, and more.

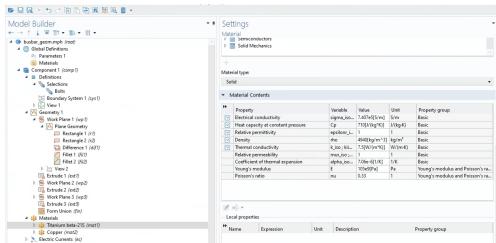


Figure 3.17: Material Properties. Source: [2]

3.1.4 Defining the Physics Boundary Conditions

~~Fixed~~ It's time to apply mathematical formulas across various sections of our model to replicate the intended physics. This involves highlighting specific segments of the geometry and imposing relevant formulas and physical parameters that accurately represent those segments.

~~Fixed~~ We'll begin by setting up the physics for the electric current interface, aiming to model the flow of electricity from a singular bolt across to the twin bolts at the opposite end of the busbar. The specific physics chosen during the initial model setup might already have certain equations predefined within their physics nodes. For example, within the "Electric Currents (ec)" node, the settings window reveals equations pertinent to this physics. A key equation presented is the fundamental one linking the electric field with electric potential, expressed as $\mathbf{E} = -\nabla V$.

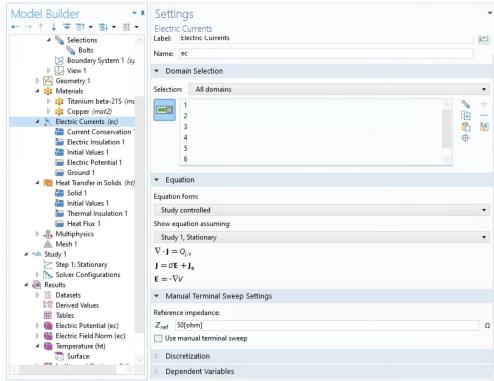


Figure 3.18: The corresponding guiding equation. Source: [2]

COMSOL Multiphysics operates transparently, allowing users to fully grasp the software's actions and methodologies. *ha!*

Upon integrating physics into our model, several default nodes were automatically established within each physics category. We add specific boundary conditions and constraints only if they deviate from these defaults. For example, the “Electric Currents (ec)” physics automatically applies the “Electric Insulation 1” condition to all boundaries, which may not suit our model’s requirements.

For our purpose, we aim to impose a voltage on the bottom surface of the single bolt. This is achieved by navigating to the Physics tab, selecting Boundaries, and then choosing Electric Potential. This action adds an “Electric Potential 1” node. The next step is to identify and select the geometry segment this potential will affect. To specify the electric potential value, we input 20 mV, denoted as 20[mV], with the brackets indicating unit specification.

To simplify the process, remember that the “Parameters 1” node contains a predefined variable, V_{tot} , representing the applied voltage. Utilizing this variable instead of inputting 20[mV] directly facilitates the setup of parametric studies in the future.

Finally, to complete our setup, we establish a ground boundary condition on the bottom surfaces of the two bolts at the busbar’s opposite end. This is accomplished by selecting Ground from the Physics tab’s boundaries options, then choosing the relevant geometry segments.

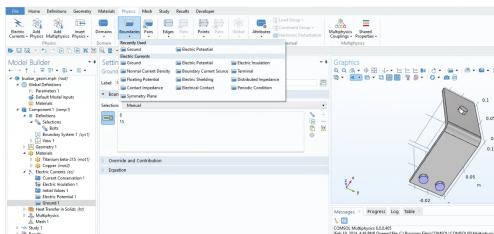


Figure 3.19: Electric Currents boundary conditions. Source: [2]

Given the selection of Joule Heating as a coupled physics in our model, it's essential

Fixed

to define the thermal boundary conditions as well. The default “Thermal Insulation 1” condition, which applies universally across our model, does not suit our specific needs. To address this, proceed to the Physics tab, access Boundaries, and opt for Heat Flux. Within the settings, under Boundary Selection, choose “All Boundaries” but exclude the previously selected bolt surfaces for the Electric Currents boundaries—namely, boundaries 8, 15, and 43—by deselecting these numbers using the “Remove from Selection” option.

For the Heat Flux settings, opt for the “Convective heat flux” under the Flux Type category, entering a value of $5 \text{ W}/(\text{m}^2 \cdot \text{K})$. Similar to our approach with the electric potential, revert to the Parameters¹ 1 node and input htc , which denotes the value of $5 \text{ W}/(\text{m}^2 \cdot \text{K})$.

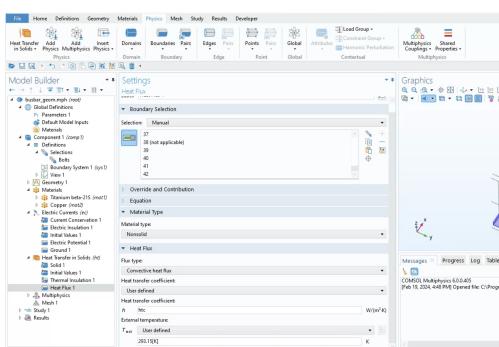


Figure 3.20: Heat Flux boundary conditions. Source: [2]

In our busbar model, the two physics components we’ve incorporated, namely Electric Currents and Heat Transfer in solids, are universally applicable throughout the model.

3.1.5 Build the Mesh

. Navigate to the Mesh ribbon and initiate mesh configuration by selecting “Mesh 1.” Within the Settings, under Mesh Setting, accessing the Sequence Type menu presents two alternatives: the default Physics-controlled mesh, which automatically tailors the mesh to the model’s physics requirements, and the user-controlled mesh, granting manual oversight over mesh granularity and enabling the use of diverse element types.

COMSOL Multiphysics accommodates a variety of 2D and 3D element shapes, including pyramids, triangles, and prisms. The software offers nine predefined element size settings, ranging from extremely fine to extremely coarse. To construct the mesh, simply press the “Build All” button.

3.1.6 Run the Study

. By selecting and expanding the “Study 1” node within the Settings, there’s an option to “Generate default plots”, which automatically creates visual representations tailored to the physics defined in the model. Given our focus on electric currents and heat transfer,

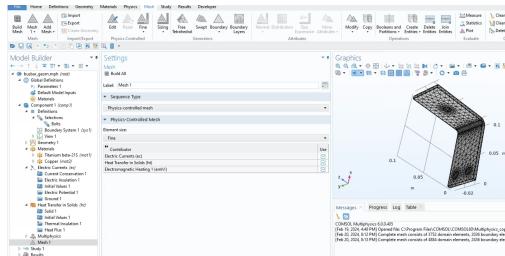


Figure 3.21: Fine mesh type. Source: [2]

this function will produce plots detailing both the electric potential and the temperature distribution within the busbar. *Can you show these?*

Under the “Step 1: Stationary” sub-node, found in the “Physics and Variables” dropdown, the integration of the multi-physics problem within the “Solve For” section becomes visible. Proceed by clicking the “Compute” button to perform the simulation. Once computation is complete, the process advances to the next phase, which involves the post-processing of the generated results.

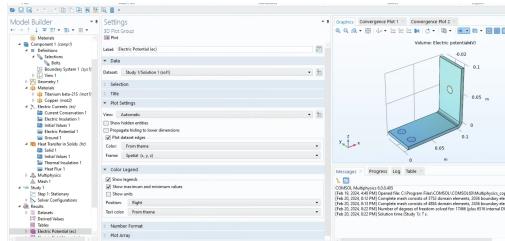


Figure 3.22: Computed results. Source: [2]

3.1.7 Post-Processing the Results

It's not obvious to ^{fixed} what is happening in this section. What is the goal here?

Upon finalizing the model construction process, the next step involves encapsulating it into a simulation application, achievable via the Application Builder. This tool enables the transformation of the busbar model into an interactive application, providing the flexibility to manipulate specific parameter values.

The Application Builder is readily accessible from the home toolbar.

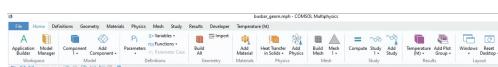


Figure 3.23: Application builder button. Source: [2]

The quick and easy way to create an app is to use the New Form Wizard.

From our model, we select parameters whose values can be adjusted, serving as input variables for our simulation application. We opt for the busbar’s length, width, and applied



Figure 3.24: New Form Wizard button Source: [2]

voltage as these inputs. Following this, we move to the graphics tab to specify which visuals will be shown in the app. In the subsequent Buttons tab, it's crucial to add the “Compute Study” button to initiate simulations.

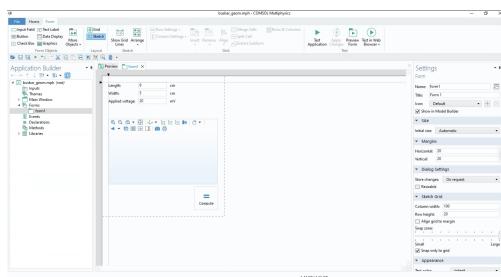


Figure 3.25: Form Editor. Source: [2]

Upon clicking “Done”, the Form Editor is launched, allowing us to refine the app’s user interface. This includes customizing its look, structuring the display through various forms, and implementing methods to extend its functionality significantly.

If we click the Test Application button, our simulation app appears.

We can change some of the values of the parameters such as increasing the length and the width and re-run the app by clicking “Compute”.

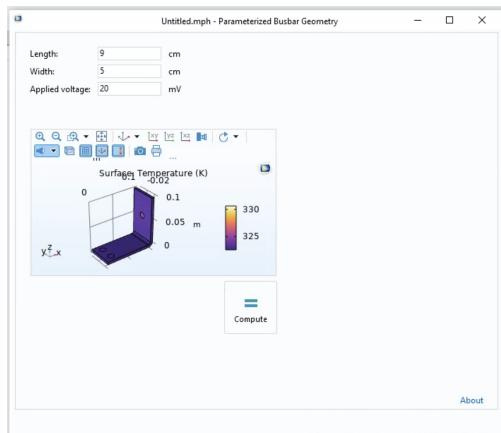


Figure 3.26: Test Application button. Source: [2]

Add a concluding paragraph. What should the reader take away from this chapter?

Chapter 4

Computational Results

Embarking on the findings of this chapter, we transition from the established COMSOL modeling workflow to the critical stage of validating theoretical concepts through the lens of computational analysis. The essence of this chapter is twofold: first, to harness the computational prowess of COMSOL in substantiating key optical phenomena such as anti-reflectivity thus corroborate our simulation outcomes with theoretical models; and second, to attempt to model the simplest Passive Daytime Radiative Cooling devices (PDRCs) at Hudgings Lab by showing how their reflectance varies with wavelength.

Our investigation begins with an empirical analysis of anti-reflective behaviors, examining both simple and composite multilayer arrangements. These simulations are meticulously compared to the theoretical predictions presented in *Introduction to Optics* by Frank L. Pedrotti et al., ~~ensuring our findings adhere to the rigorous standards of optical precision.~~

We delve into the intricacies of high reflectance by simulating stacks of alternating high and low refractive index layers, shedding light on the nuanced ways layer configurations impact reflectance. These insights are bench-marked against scholarly literature, reinforcing the accuracy of our models.

Moreover, the thesis explores the layered design of PDRCs. By systematically layering glass, silver, and Polydimethylsiloxane (PDMS), we dissect the optical properties crucial to the development of passive cooling systems.

This chapter stands at the intersection of physics and simulation, merging the realms of theory and experimentation. The collection of results and graphs here encapsulate more than just data; they are catalysts for continuous inquiry and deeper understanding of optical physics as it related to PDRCs.

4.1 COMSOL: Modeling Anti-reflectivity.

Recalling the formula for the reflectance of a two-layer anti-reflecting film, given by equation 2.55, and assuming light strikes normally, we have:

I'm finding 4.1 & 4.2 a bit hard to follow. For now, can you do the two things circled in RED below? We can work on organization together when I return.

Maybe include in your intro a reminder that while we can analytically derive $R(\theta)$ & $R(\lambda)$ for materials with constant n , as detailed in Ch 2, this becomes very difficult when refractive index varies with λ . Hence, the need for COMSOL!

*Work on grouping into paragraphs
and on equation punctuation
(see comments in Ch 2)*

$$R = \left(\frac{n_0 n_2^2 - n_s n_1^2}{n_0 n_2^2 + n_s n_1^2} \right)^2 \quad (4.1)$$

Here, we anticipate zero reflectance when the expression in the numerator becomes zero, leading us to equation 2.56, which stipulates the condition for achieving no reflectance:

$$\frac{n_2}{n_1} = \sqrt{\frac{n_s}{n_0}} \quad (4.2)$$

This relationship delineates the necessary ratio between the refractive indices of the layers that leads to a reflectance of zero.

When working with a glass substrate of refractive index $n_s = 1.5$ and air of refractive index $n_0 = 1$, the ideal refractive index ratio, $\frac{n_2}{n_1}$, is calculated to be $\sqrt{1.5} \approx 1.225$.

In COMSOL, our goal is to identify and define (in our parameters table) two materials for the double-layer structure that closely match this ratio to maximize anti-reflective properties. It's crucial to note that within the visible spectrum, ranging from 400 to 700 nm, each wavelength uniquely interacts with materials based on their specific refractive index at that wavelength. This variability, known as dispersion, means that optimal thickness and refractive indices for minimizing reflectance at one wavelength may not be as effective at another. While it's challenging to select materials that precisely align their refractive indices to achieve exactly zero reflectance, our aim is to get as close to 0% reflectance as practically possible.

*the refractive index of most materials varies with wavelength.
over as wide a λ range*

The limitation of a quarter-wavelength layer is its selective reflectivity; it effectively eliminates reflection at a specific wavelength but substantially reflects at other wavelengths in proximity. Additionally, its performance varies significantly with the angle at which light strikes the surface. A viable solution to overcome this is the application of multilayer coatings. These coatings, compared to their single-layer counterparts, tend to lower the reflection coefficient over a broader spectrum and accommodate a greater diversity of tangible materials [3].

In the following section, we will compare the reflectivity of two distinct multilayer coatings across an extensive spectral range. The analysis will cover a dual-layer, quarter-quarter wavelength coating, and a triple-layer, quarter-half-quarter wavelength coating. It will be demonstrated that the triple-layer coating achieves a more uniform low reflectance throughout the majority of the visible light spectrum.

4.1.1 Anti-reflectance Layers: Setup

The pursuit of anti-reflective coatings leads us to the utilization of thin film dielectrics within COMSOL. In line with the criteria for zero reflectance, the first dielectric film, starting from the base, is cerium trifluoride (CeF_3) with a refractive index of 1.63, while the second film is magnesium fluoride (MgF_2) with a refractive index of 1.38. The refractive index ratio of approximately 1.181 aligns closely with the optimal ratio of 1.225. As established in chapter

*Ce = Cesium?
 CeF_3 ? MgF_2*

Please add a figure showing each of these 2 test structures & thicknesses.

2, effective anti-reflectivity for a dual-layer arrangement requires each film to be a quarter-wavelength in thickness, which corresponds to the wavelength at which minimal reflectance is desired.

In our enhanced three-layer structure, designed to broaden anti-reflectivity across a larger wavelength spectrum, the middle layer is a half-wavelength film of zirconium dioxide (ZrO_2), chosen for its refractive index of 2.1. This multi-layer configuration allows for increased anti-reflective performance compared to the more wavelength-specific two-layer model.

This screenshot from the COMSOL desktop highlights the configuration for modeling anti-reflective coatings. Pay particular attention to the 'Film Properties' area within the 'Thin Dielectric Film' settings where the quarter-wavelength calculation is outlined.

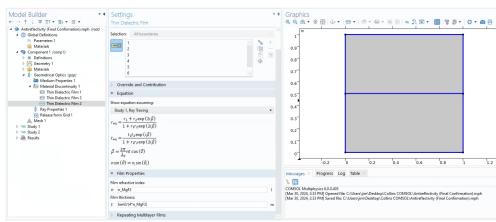


Figure 4.1: COMSOL desktop setup showcasing anti-reflectivity modeling.

Here, the findings from the anti-reflectivity modeling using CeF_3 and MgF_2 for the quarter-quarter wavelength combination, as well as CeF_3 , MgF_2 , and ZrO_2 for the quarter-half-quarter wavelength setup, are presented.

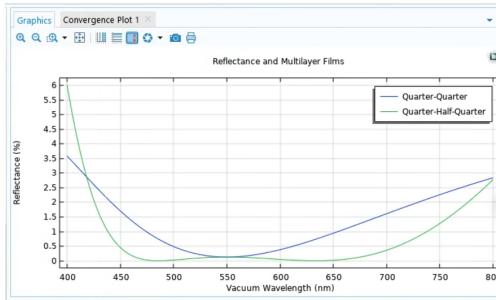


Figure 4.2: Illustration of the reflectance behavior for two anti-reflective coating configurations across the visible light spectrum. The blue line indicates the performance of a quarter-quarter wavelength coating, whereas the green line shows the effect of a quarter-half-quarter wavelength coating.

In the quarter-quarter wavelength design, we employ two layers, each precisely a quarter of the desired wavelength in thickness. While the aim is to achieve minimal reflectance at a particular wavelength, the actual reflectance seldom reaches absolute zero due to the inherent limitations of material characteristics.

On the other hand, the quarter-half-quarter arrangement introduces an additional layer, half a wavelength thick, nestled between two quarter-wavelength films. This configuration is tailored to minimize reflectance across an extended spectrum of wavelengths.

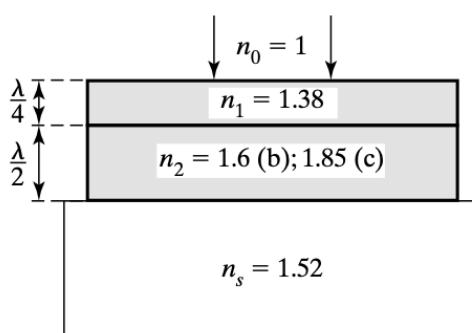
Such multi-layered coatings surpass the effectiveness of single-layer films for anti-reflective purposes by accommodating a wider wavelength range. This versatility is particularly beneficial in applications where reducing reflection is paramount, such as in the manufacturing of optical lenses.

4.1.2 Verification of Theoretical Results from Optics Literature

This section aims to validate the principles outlined in *Introduction to Optics* by Frank L. Pedrotti et al. through the simulation of multilayer film behavior, ensuring our computational results align with established optical theories. *I think you're validating your simulations, not the existing theory*

Beginning with a straightforward approach, we examine the reflectance across various wavelengths for three distinct multilayer film configurations:

1. Films with quarter-quarter wavelength thickness.
2. Films with quarter-half wavelength thickness.
3. Films with quarter-half wavelength thickness, incorporating a different material for the half-wavelength layer.



Please show a figure like this for each test case used by Pedrotti or in your CONSOL work. Label the material used in each layer.

Figure 4.3: Antireflective double-layer arrangement. Source: [3]

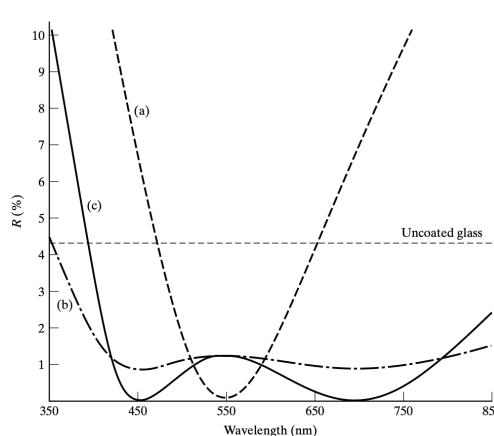
We first examine films with quarter-quarter wavelength thickness. The reference from Pedrotti et al. specifies the lower thin dielectric layer as ZrO_2 and places CeF_3 directly above. The choice of materials is flexible, provided they adhere to the optimal ratio criterion for antireflectivity as defined in equation 4.2. This results in a refractive index ratio of $\frac{n_2}{n_1} = \frac{2.1}{1.65} \approx 1.273$, aligning closely with the ideal ratio of approximately 1.225 for normal incidence, given a substrate refractive index (n_s) of 1.5.

Expanding the spectrum of low reflectance within the visible region becomes achievable by diverging from the constraint of equal quarter-wavelength ($\lambda/4$) coatings. By integrating a layer of quarter-wavelength thickness as the second layer (considered from the bottom upwards), we attain more extensive zones of diminished reflectance. Consequently, in the scenario of item 2, we employ magnesium fluoride (MgF_2), characterized by a refractive

index of 1.38, as the quarter-wavelength thick material. The intermediate half-wavelength layer utilizes aluminum oxide, with a refractive index of 1.60. For item 3, thorium dioxide, featuring a refractive index of 1.85, serves as the material for the half-wavelength thick layer, as noted in [3].

For the specific wavelength of 550 nm, where the thicknesses of the quarter-wavelength ($\lambda/4$) and half-wavelength ($\lambda/2$) layers are calculated, the half-wavelength layer does not influence reflectance. In this case, the double-layer system acts akin to a solitary quarter-wavelength layer, resulting in a reflectance of 1.3%. At wavelengths close to 550 nm, the presence of the half-wavelength layer contributes to maintaining reflectance levels below those achieved by a two quarter-wavelength layers. Below is the plot illustrating reflectance against wavelength for scenarios 1 through 3:

The graph below illustrates anti-reflectivity trends as detailed in the Optics textbook, indicating that while the reflectance at a wavelength of 550 nm stands at approximately 1.3%, this value surpasses the performance of the quarter-quarter wavelength coating. Nevertheless, reflectance stays below this threshold across a wide wavelength span, extending from roughly 420 to 800 nm. This suggests that alternative configurations for dual-layer reflective films could be viable if the layers are not strictly constrained to quarter-wavelength multiples.



Can you duplicate
this exact figure
using COMSOL?
And then show the 2
side-by-side.

Figure 4.4: Anti-reflectivity Graphs as shown in the Optics Textbook. Source: [3]

Presented below are the modeled results for scenarios (a), (b), and (c) as detailed previously, with corresponding legends found in figure 4.4. Notice how each individual plot resembles each plot in figure 4.4.

These results derive from the principles outlined in Chapter 2. Initially, the overall transfer matrix elements are calculated by multiplying the transfer matrices of the individual layers together. Within these calculations, the phase difference, δ , varies with λ , aligning the film thickness with either $\lambda/4$ or $\lambda/2$ at the reference wavelength of 550 nm.

Subsequently, the reflection coefficient, as outlined in 2.52, is squared to produce reflectance as a wavelength function. COMSOL's Ray Optics module significantly simplifies this process by automating several steps, although careful consideration is needed for layer treatment, particularly regarding their classification as thin dielectric films.

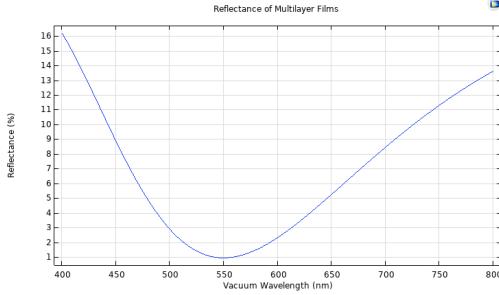


Figure 4.5: Modeling results of an anti-reflective coating with two layers, each a quarter-wavelength thick.

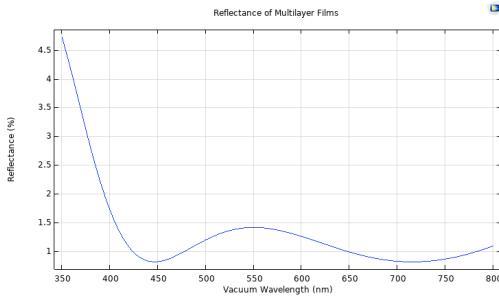


Figure 4.6: Reflectance outcomes for an anti-reflective coating with a bottom layer a quarter-wavelength thick and a top layer half-wavelength thick, using a material with a refractive index of 1.6.

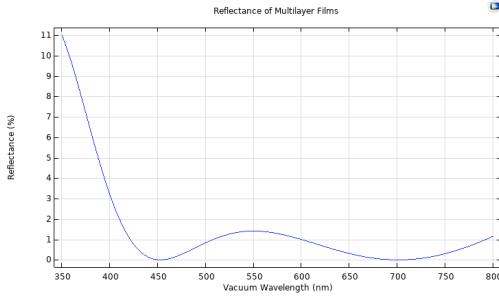


Figure 4.7: Reflectance results for an anti-reflective double layer with a bottom layer a quarter-wavelength thick and a top layer half-wavelength thick, using a material with a refractive index of 1.85.

4.2 COMSOL: Modeling High Reflectance.

To achieve anti-reflective properties, layers are arranged starting from the air and progressing through low-index to high-index materials before reaching the substrate. In contrast, for enhanced reflectivity, the sequence is reversed, starting from the air to high-index and then to low-index materials before ending at the substrate. This arrangement encourages multiple

reflections of the incident light within the layered structure.

A combination of layers arranged to maximize reflectivity is known as a *dielectric mirror*, *high-reflectance stack*, or *distributed Bragg reflector*. It's important to note that achieving high reflectance across the solar spectrum is a key factor for effective passive daytime radiative cooling, as previously discussed.

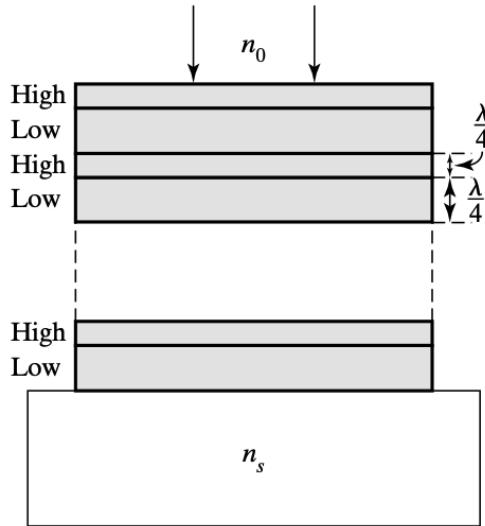


Figure 4.8: High-reflectance stack comprising double layers with alternating high and low refractive indices. Source: [3]

Again, you're verifying your GmsoL results, not the theory. ☺

4.2.1 Verification of Theoretical Results from Optics Literature.

The formula to ascertain high reflectance, R , at a chosen design wavelength (λ_0) for a specific layer arrangement is denoted by 2.59.

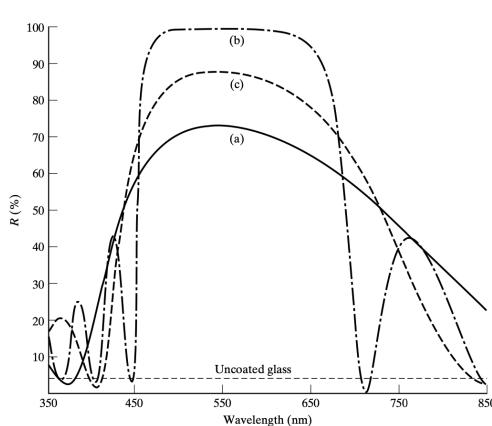
$$R = \left[\frac{\left(\frac{n_0}{n_s} \right) \left(\frac{n_L}{n_H} \right)^{2N} - 1}{\left(\frac{n_0}{n_s} \right) \left(\frac{n_L}{n_H} \right)^{2N} + 1} \right]^2 \quad (4.3)$$

Achieving maximal reflectance (100%) occurs under the following conditions:

1. As N , representing the count of layer pairs, becomes very large.
2. When the ratio $\frac{n_L}{n_H}$ approaches zero.

The optics textbook illustrates high reflectance with the following depiction:

In these configurations, layers are designed to be $\lambda/4$ thick at a design wavelength $\lambda_0 = 550$ nm. Here, the high-index material (Zinc Sulfide - ZnS) has $n_H = 2.35$, the low-index



Can you replicate exactly this using COMSOL and then show the two side-by-side?

Figure 4.9: Reflectance spectra for high-low index stacks: (a) for a two-layer double stack, (b) for a six-layer double stack, and (c) for a two-layer double stack enhanced with an additional high-index layer at the very top. Source: [3]

material (Magnesium Fluoride - MgF₂) has $n_L = 1.38$, and the incident medium (air) has $n_0 = 1.00$. The ratio of $\frac{n_L}{n_H}$ is thus approximately $\frac{1.38}{2.35} \approx 0.587$.

Our analysis will specifically address graph (c), which demonstrates how adding an extra high-index layer between the substrate and the final low-index layer enhances maximum reflectance in a two-layer double stack ($N = 2$). This addition improves reflectance efficiency compared to simpler two-layer double configurations.

The findings showcased below stem from applying the high reflectance principles discussed in Chapter 2.

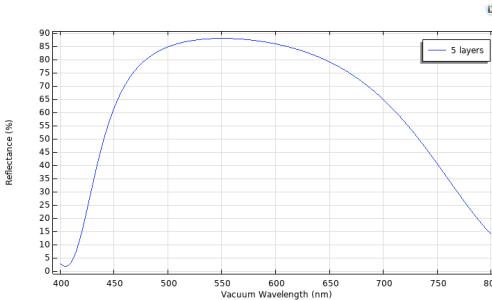


Figure 4.10: Spectral reflectance for a structure with two pairs of high-low refractive index layers, topped with an additional high-index layer.

Observe the alignment of peak reflectance with the textbook's findings, achieving 90

Employing a parametric sweep in COMSOL allows for the exploration of reflectance variation across wavelengths by systematically adjusting the count of double layer pairs (N) to values such as 2, 5, 10, and 20. This approach calculates solutions across a range of parameter sets. For all depicted graphs, an extra top layer with a high refractive index is incorporated to broaden the spectrum of maximum reflectance.

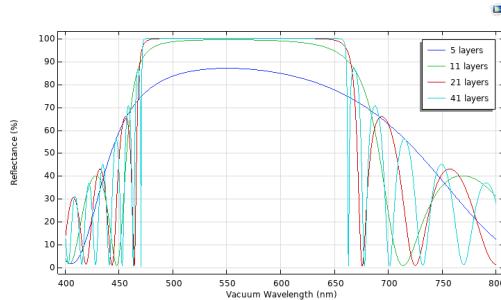


Figure 4.11: Reflectance comparison for configurations with 5, 11, 21, and 41 layers, each enhanced by an additional high-index surface layer.

Unlike conventional metallic mirrors, DBRs leverage a careful and simple engineered structure of alternating thin layers of materials with contrasting refractive indices. This design typically features an odd number of layers, anchored by high refractive index materials at both ends, optimizing their reflectivity [11].

At the core of a DBR's functionality is the concept of constructive interference. As optical waves encounter the boundaries between layers, each boundary induces a partial reflection. When the optical wave's wavelength is four times the layers' optical thickness, these reflections interfere constructively, transforming the layers into an efficient reflector. This principle gives rise to a *stopband*, a wavelength range where the reflector achieves heightened reflection. With an adequate number of layers, a DBR can achieve a high-quality reflection, making it useful in the operation of vertical cavity surface emitting lasers [11].

Expanding beyond the *stopband*, DBRs exhibit a characteristic where reflectance transitions into a pattern of oscillating maxima and minima.

By strategically adjusting these parameters, it is possible to shift the stopband's center, enhancing the filter's spectral transmittance across a broader range. This capability to finely control the light's passage through these structures not only underscores the versatility of DBRs but also paves the way for their integration into a wide array of optical devices, where precise manipulation of light is paramount [3].

Building on the principles of DBRs and their implications for enhanced reflectivity, a parallel can be drawn to PDRCs. PDRCs, which aim to achieve cooling without energy input by radiating heat into the cold outer space, can significantly benefit from the reflective properties and spectral selectivity offered by DBR structures.

PDRCs leverage similar concepts to reflect the solar spectrum while allowing the thermal infrared range to pass through effectively. By incorporating multilayer structures like those used in DBRs, PDRC devices can optimize their performance, achieving a delicate balance between high solar reflectance and high thermal emittance. This enables PDRC systems to reflect the majority of incoming solar radiation, minimizing heat absorption, while simultaneously maximizing the emission of infrared radiation for cooling.

I wonder if this might go well in Ch5 under future work?

4.3 COMSOL: Modelling PDRCs

At Hudgings Lab, one of the foundational PDRC designs comprises a straightforward sequence of layers. Beginning with silica glass as the foundation, a layer of silver is applied directly above, topped with a final layer of Polydimethylsiloxane (PDMS).

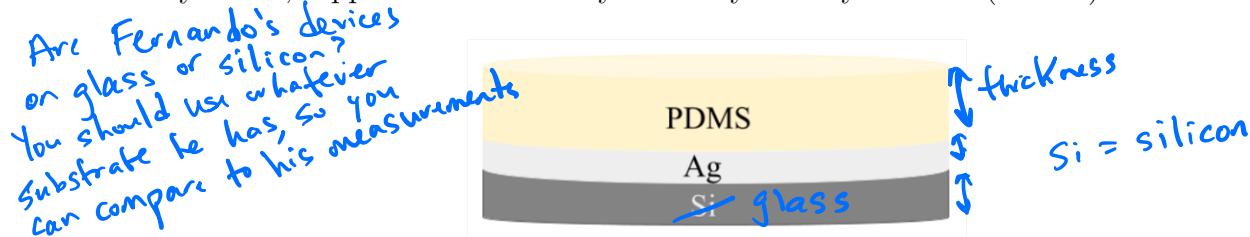


Figure 4.12: Illustration of a basic PDRC configuration at Hudgings Lab, featuring a silica glass base with subsequent layers of silver and PDMS.

To understand the optical behavior of this structure, I systematically approached the modeling process from the bottom layer upwards. The following results present a sequence of reflectance versus wavelength analyses for each layer configuration: initially for glass, followed by glass with an added silver layer, and concluding with the composite structure of glass, silver, and PDMS. This progression illustrates how each layer contributes to the overall reflectance spectrum.

4.3.1 Analysis of Glass Layer Only

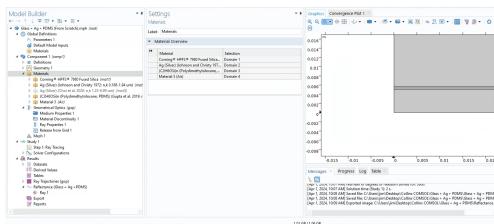


Figure 4.13: COMSOL desktop setup for PDRC analysis demonstrating the simulation framework.

For simulations that span across various wavelengths, it is crucial to utilize materials with wavelength-dependent refractive indices. In COMSOL Multiphysics, this requirement is met through the use of interpolation functions, which map refractive indices to specific wavelengths. This method allows for dynamic adjustment of a material's refractive index in response to changes in wavelength during simulations, providing a more accurate representation of the material's optical behavior.

Utilizing the interpolation table enables the plotting of an interpolation function, offering a graphical representation of how refractive indices fluctuate with wavelength. This capability enriches the COMSOL simulation by providing a more precise depiction of light-material

→ COMSOL tabulates $n(\lambda)$ for specific λ measured in the literature. Then it interpolates between these experimentally measured points.

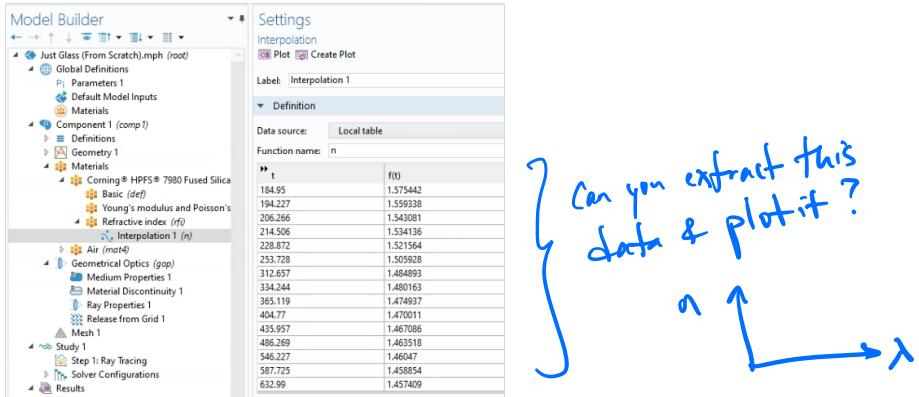


Figure 4.14: Interpolation table for the glass substrate, showing the wavelength-dependent refractive index.

interaction over diverse wavelengths, a critical aspect in optics where dispersive phenomena play an important role.

For this reason, I selected Corning™ HPFS™ 7980 Fused Silica for the substrate, taking advantage of its accompanying interpolation function. *It's vital to acknowledge that interpolation functions, which depict the wavelength-dependent refractive index variations of materials, can also be derived from experimental data found in scientific literature.*

When generating a reflectance versus wavelength graph using only the glass substrate (specifically, Corning™ HPFS™ 7980 Fused Silica in this case), the reflectance indeed shows variation across different wavelengths.

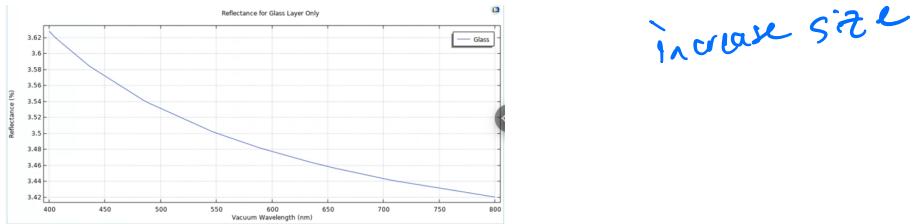


Figure 4.15: Reflectance variation with wavelength for Corning™ HPFS™ 7980 Fused Silica.

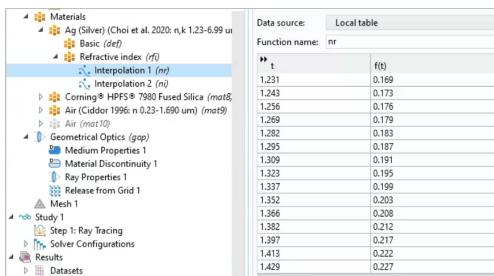
Within the displayed wavelength range of 400 – 800 nm, a sole substrate yields a reflectance below 4%, *which is expected for a largely transparent glass substrate.* This falls short of the desired high reflectivity within the solar spectrum for optimal PDRC performance. This observation underscores the need for integrating materials capable of meeting the dual objectives of high solar reflectance and substantial emissivity within the atmospheric window.

Focusing initially on enhancing solar reflectance, designs at Hudgings Lab incorporate silver on top the substrate due to its exceptional reflectivity in the solar spectrum. Having illustrated the outcomes with merely the glass substrate and underscored the importance of a highly reflective layer in the solar spectrum, I will venture into simulating the combined effects of glass and silver layers.

4.3.2 Analysis of Glass plus Silver Layers.

Next, I added a silver layer on top the glass substrate, recognizing the continued necessity for an interpolation function to accurately represent silver's refractive index across various wavelengths.

COMSOL offers a comprehensive collection of silver materials, each accompanied by interpolation tables that chart the refractive indices over preferred wavelength spans. The entry *Ag (Silver)* (*Choi et al. 2020: n,k 1.23-6.99 μm*) provides data on silver's refractive indices from 1.23 to 6.99 (μm), where n is the real component of the complex refractive index and k —the extinction coefficient—is the imaginary part, indicating the material's degree of light absorption.



The screenshot shows the COMSOL Materials library. The 'Ag (Silver)' entry is highlighted. To the right, a table titled 'Local table' shows the refractive index (n) and extinction coefficient (k) for various wavelengths (λ in nm). The table is as follows:

Data source:	Local table
Function name:	nr
» t	nr
1,231	0.169
1,243	0.173
1,256	0.176
1,269	0.179
1,282	0.183
1,295	0.187
1,309	0.191
1,323	0.195
1,337	0.199
1,352	0.203
1,366	0.208
1,382	0.212
1,397	0.217
1,413	0.222
1,429	0.227

Figure 4.16: Interpolation table for silver's refractive index.

COMSOL's materials library offers various definitions of silver to cater to different wavelength ranges. For example, the definition *Ag (Silver)* (*Johnson and Christy 1972: n,k 0.188-1.94 μm*) provides an interpolation table covering 0.188-1.94 (μm).

The figures below display the outcomes utilizing *Ag (Silver)* (*Johnson and Christy 1972: n,k 0.188-1.94 μm*) and using *Ag (Silver)* (*Choi et al. 2020: n,k 1.23-6.99 μm*), respectively.

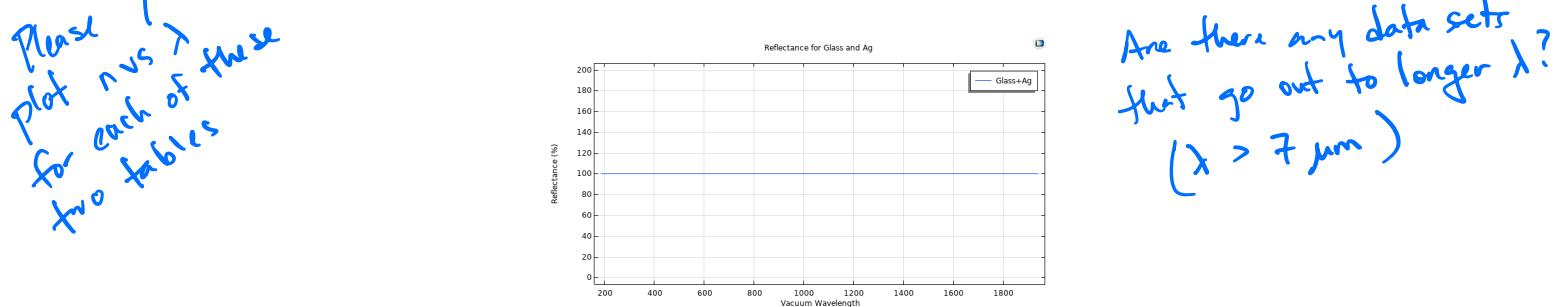


Figure 4.17: Reflectance analysis for glass with Ag (Johnson and Christy).

These graphs mostly reflect the expected outcomes. Given silver's high reflectivity across the visible wavelength spectrum, we observe near 100% reflectance within the visible range (400 - 700 nm).

Did you originally get ≈ 100 at small λ ?

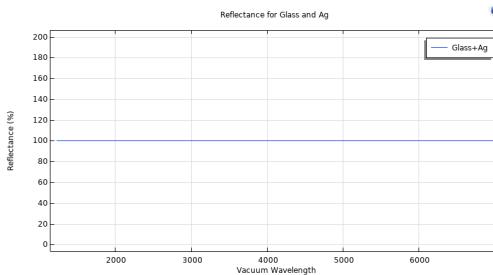
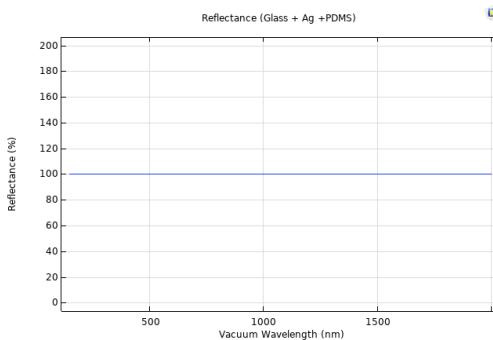


Figure 4.18: Reflectance analysis for glass with Ag (Choi et al.).

4.3.3 Analysis of Glass plus Silver plus PDMS Layers.

In the final step, a PDMS layer was applied on top of the silver layer. This step also utilized a PDMS material equipped with a pre-defined interpolation function for its refractive index, specifically $(C_2H_6OSi)_n$ (*Polydimethylsiloxane, PDMS*) (Gupta et al. 2019: n 0.30-1.69 μm).

For each simulation plot of R vs λ , please plot it with Fernando's experimental data plots.



Again, we need to go to much longer λ (out to 20 μm) for our devices

Figure 4.19: Reflectance analysis for a composite of glass, silver, and PDMS.

This configuration resulted in reflectance maintaining a peak at 100% across an extensive range of wavelengths.

We conclude our examination of optical phenomena using COMSOL. This chapter had two aims: first, to validate theoretical concepts like anti-reflectivity and high reflectance using COMSOL, and second, to initiate the simulation of basic PDRC configurations as designed at Hudgings Lab.

We began by evaluating anti-reflective strategies, ranging from simple to complex multilayer systems, against theoretical predictions from *Introduction to Optics* by Frank L. Pedrotti et al. This comparison affirmed the accuracy of our simulations. Additionally, we modeled PDRC devices by layering glass, silver, and Polydimethylsiloxane (PDMS), revealing key optical properties essential for passive cooling effectiveness.

This chapter demonstrates the synergy between theoretical physics and computational modeling, offering a solid foundation for future research to expand upon. It underscores the

importance of simulation in confirming theoretical models and opens avenues for improving PDRC device design.

Looking forward, the results from this chapter not only validate our understanding of optical principles but also suggest strategies for enhancing PDRC performance through material and design optimization.

Chapter 5

Conclusion

The initial chapters laid the groundwork by illustrating the critical role of cooling technologies in addressing the pressing issues of global warming and the energy crisis. With an emphasis on PDRCs, we embarked on a journey through the theoretical underpinnings and practical applications of these devices, which offer the potential for efficient, energy-free cooling by radiating heat directly to outer space during daylight hours.

Throughout this exploration, we leveraged the capabilities of COMSOL Multiphysics™ to simulate various PDRC configurations, aiming to enhance their performance by optimizing material properties and layering strategies. The results chapter validated our approach, demonstrating the effectiveness of PDRCs in achieving significant cooling effects under real-world conditions. *I'm not clear on how Ch4 demonstrates this.* ?.

As we culminate this research, this concluding chapter aims to distill the insights gained from our investigations, outlining the contributions of this work to the field of cooling technologies and the broader context of sustainable energy solutions.

5.1 Our work

COMSOL Multiphysics™ has proven itself as an indispensable tool for computational modeling, facilitating the simulation of intricate PDRC structures with relative ease. The modeling approach, exemplified in Chapter 3 through a detailed walkthrough with a busbar example, laid the foundation for all subsequent simulations. *You are corroborating/validating your COMSOL models against the established theory*

This thesis has not only corroborated established theoretical principles within the realms of anti-reflectivity and high reflectance but has also ventured into the modeling of Fresnel equations to observe the behavior of reflection coefficients across varying angles of incidence.

The core of our efforts was the computational modeling of PDRC designs from Hudgings Lab, layer by layer, revealing critical insights into their optical performance. Notably, the simulation results confirmed the imperative for PDRCs to exhibit high reflectivity within the solar spectrum, a requirement clearly illustrated by the limited reflectivity observed in the sole glass substrate model.

5.2 Future Directions

Moving forward, several avenues for further research have been identified:

1. Enhanced definition in the reflectance versus wavelength graphs for glass plus silver and glass plus silver plus PDMS models, possibly through the utilization of more detailed refractive index interpolation tables or functions.
2. Exploration of adding more materials on top of the PDMS layer to fulfill PDRC design criteria more effectively, experimenting with various thicknesses and refractive indices to exploit constructive interference across multiple interfaces more efficiently.
3. The translation of computational models into physical prototypes for empirical validation in lab settings, comparing simulated reflectance with real-world performance.
4. Development of reflectance versus angle of incidence models to aid in the characterization of PDRC behavior throughout the day, providing a benchmark for experimental testing on rooftop installations.

5.3 Final Remarks

This thesis not only validates the utility of PDRCs within the context of sustainable cooling technologies but also charts a course for their further development and application especially on the computational modeling front.

As we look to the future, the insights and methodologies presented here will serve as a valuable resource for those endeavoring to advance the efficiency and applicability of PDRCs and similar technologies in our collective pursuit of environmental sustainability and energy efficiency.

Appendix A

An appendix

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