Chemical Modification of Lignin into Advanced Materials

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Abstract

Fossil fuel resources are being used today for most of humankind’s energy and chemical/material needs. The inevitable demise of these resources has created significant interest in the field of biomass and particularly, lignin valorization. As the world’s second most abundant polymer, more than 98% of the annually produced lignin is under-utilized either as an on-site heat source or as landfill. Thus, finding practical approaches to modifying this inexpensive, sustainable resource into materials of high value can be the next leap in lessening the dependence on fossil fuel resources and thus, developing a sustainable future.

In this thesis, kraft lignin is modified into methacrylated lignin (ML), lignin grafted with methacrylate functionalities at its hydroxyl sites. ML was incorporated at loading percentages of 10-31 wt.% into UV-cured coatings, and the developed coatings were characterized according to their different properties.

The methacrylation process was optimized via response surface methodology using a central composite design to determine the effect of different reaction variables on the process’ recovered mass yield and to obtain the maximum possible recovered ML. The ML obtained from the optimized reaction conditions was then incorporated at 30 wt.% into a siloxane-based UV-cured coating as a proof of concept for the practicality of the optimization process.

ML was afterwards converted into a primary polyphosphine (lignophine) by a phosphane-ene reaction. The primary lignophine was then capped into tertiary alkylated and fluorinated lignophines, respectively using corresponding alkenes. The ability of the tertiary alkylated lignophine to coordinate to transition metals and sequester transition metal-containing catalysts was determined using silver triflate a ring closing metathesis (RCM) reaction using Grubbs I (GI) catalyst, respectively.

Finally, the tertiary alkylated lignophine was converted into a free-standing UV-cured polyphosphonium (lignophonium) network. The obtained films showed high cure percentages and water contact angles, along with a surface charge density value well over the necessary threshold for antimicrobial films and swelling degrees suitable for controlled drug release. The
films were probed for their controlled drug release ability using diclofenac as the loaded drug and examining the release in PBS and showed a promising release profile.

Keywords
Co-Authorship Statement

Chapter 3: UV-curable Coatings of Modified Lignin

Authors: Soheil Hajirahimkhan, Chunbao (Charles) Xu, Paul J. Ragogna

The experimental work, characterization and, result analyses were conducted by Soheil Hajirahimkhan under the supervision of Prof. Paul J. Ragogna and Prof. Chunbao (Charles) Xu. The manuscript was written by Soheil Hajirahimkhan and revised by Prof. Paul J. Ragogna and Prof. Chunbao (Charles) Xu and submitted to “Sustainable Chemistry and Engineering”.

Chapter 4: Methavrylation of Kraft Lignin for UV-curable Coatings: Process Optimization Using Response Surface Methodology

 Authors: Soheil Hajirahimkhan, Paul J. Ragogna, Chunbao (Charles) Xu

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Chapter 5: “Lignophines”: Lignin-based Tertiary Phosphines with Metal-Scavenging Ability

Authors: Soheil Hajirahimkhan, Devon E. Chapple, Johanna M. Blacquiere, Chunbao (Charles) Xu, Paul J. Ragogna

Reactor setup for the hydrophosphination reaction, tertiary lignophile synthesis, all product characterizations, and silver triflate coordination studies was conducted by Soheil Hajirahimkhan under the supervision of Prof. Paul J. Ragogna and Prof. Chunbao (Charles) Xu. The PH3 line was operated by Dr. Tyler J. Cuthbert, Dr. Cameron M. E. Graham, Tristan D. Harrison, Vanessa A. Béland, and Dr. Kaijie Ni at different times during the project under the supervision of Prof. Paul J. Ragogna. Metal sequestration studies were performed, and the results were analyzed by Devon E. Chapple under the supervision of Dr. Johanna M. Blacquiere. The manuscript was written by Soheil Hajirahimkhan and Devon E. Chapple and
Chapter 6: “Lignophonions”: Stimuli-Responsive Lignin-based Polyphosphonium Networks for Controlled Drug Delivery

Authors: Soheil Hajirahimkhan, Tristan D. Harrison, Chunbao (Charles) Xu, Paul J. Ragogna

All synthesis, characterization, and result analyses were conducted by Soheil Hajirahimkhan under the supervision of Prof. Paul J. Ragogna and Prof. Chunbao (Charles) Xu. Drug release studies and result analyses were carried out with the collaboration of Tristan D. Harrison. The manuscript was written by Soheil Hajirahimkhan and Tristan D. Harrison and revised by Prof. Paul J. Ragogna and Prof. Chunbao (Charles) Xu and will be submitted to “Chemical Communications”.

revised by Prof. Paul J. Ragogna, Prof. Johanna M. Blacquierre, and Prof. Chunbao (Charles) Xu and submitted to “Angewandte Chemie International Edition”.
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<th>Description</th>
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<td>°</td>
<td>Degrees</td>
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<tr>
<td>IMIM</td>
<td>1-methylimidazole</td>
</tr>
<tr>
<td>AA</td>
<td>Acetic anhydride</td>
</tr>
<tr>
<td>AAL</td>
<td>Acetic Acid Lignin</td>
</tr>
<tr>
<td>ABDA</td>
<td>4-acryloyloxybutyl dehydroabieticcarboxylate</td>
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<tr>
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<td>American Society for Testing and Materials</td>
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<tr>
<td>ATRP</td>
<td>Atom Transfer Radical Polymerization</td>
</tr>
<tr>
<td>BA</td>
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<tr>
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<tr>
<td>CER</td>
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<td>CL</td>
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<td>Abbreviation</td>
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<td>Dimethylsulfoxide</td>
</tr>
<tr>
<td>DSA</td>
<td>Drop Shape Analysis</td>
</tr>
<tr>
<td>EB-1360</td>
<td>EBERCYL® 1360</td>
</tr>
<tr>
<td>EG</td>
<td>Ethylene glycol</td>
</tr>
<tr>
<td>EGDGE</td>
<td>Ethylene glycol diglycidyl ether</td>
</tr>
<tr>
<td>EGDMA</td>
<td>Ethyleneglycol dimethacrylate</td>
</tr>
<tr>
<td>EHL</td>
<td>Enzymatically Hydrolyzed Lignin</td>
</tr>
<tr>
<td>EOM</td>
<td>Extent of Methacrylation</td>
</tr>
<tr>
<td>FPInnovations</td>
<td>Forestry Products Innovations</td>
</tr>
<tr>
<td>FT-IR</td>
<td>Fourier Transform Infrared</td>
</tr>
<tr>
<td>G</td>
<td>Guaiacyl</td>
</tr>
<tr>
<td>GI</td>
<td>Grubbs I</td>
</tr>
<tr>
<td>GLY</td>
<td>Glycerol</td>
</tr>
<tr>
<td>GLYDGE</td>
<td>Glycerol diglycidyl ether</td>
</tr>
<tr>
<td>H</td>
<td>$p$-hydroxyphenyl</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Definition</td>
</tr>
<tr>
<td>--------------</td>
<td>------------</td>
</tr>
<tr>
<td>HDMAP</td>
<td>2-hydroxy-2-methylpropiophenone</td>
</tr>
<tr>
<td><em>in vacuo</em></td>
<td>In a vacuum</td>
</tr>
<tr>
<td>KL</td>
<td>Kraft Lignin</td>
</tr>
<tr>
<td>KWF</td>
<td>Aminosilane-Treated Wood Flour</td>
</tr>
<tr>
<td>LGA</td>
<td>Lignin-based Hydrogel</td>
</tr>
<tr>
<td>LGBA</td>
<td>ABDA polymer grafted lignin</td>
</tr>
<tr>
<td>LGEA</td>
<td>AEDA polymer grafted lignin</td>
</tr>
<tr>
<td>LGEMA</td>
<td>MAEDA polymer grafted lignin</td>
</tr>
<tr>
<td>Lignin-g-DHA</td>
<td>DHA-grafted lignin</td>
</tr>
<tr>
<td>Lignin-g-PS</td>
<td>lignin-graft-polystyrene</td>
</tr>
<tr>
<td>LPF</td>
<td>Lignin-Phenol-Formaldehyde</td>
</tr>
<tr>
<td>LS</td>
<td>Sodium lignosulfonate</td>
</tr>
<tr>
<td>LWF</td>
<td>Lignin Amine-Treated Wood Flour</td>
</tr>
<tr>
<td>MA</td>
<td>Maleic anhydride</td>
</tr>
<tr>
<td>MAEDA</td>
<td>2-methacryloyloxyethyl dehydroabieticcarboxylate</td>
</tr>
<tr>
<td>MC</td>
<td>Methacyryloyl chloride</td>
</tr>
<tr>
<td>Me₆-TREN</td>
<td>tris[2-(dimethylamino)ethyl] amine</td>
</tr>
<tr>
<td>MeCN</td>
<td>Acetonitrile</td>
</tr>
<tr>
<td>MEHQ</td>
<td>Monomethyl ether hydroquinone</td>
</tr>
<tr>
<td>MethA</td>
<td>Methacrylic anhydride</td>
</tr>
<tr>
<td>MethAcid</td>
<td>Methacrylic acid</td>
</tr>
<tr>
<td>ML</td>
<td>Methacrylated Lignin</td>
</tr>
<tr>
<td>MR₅₀₀</td>
<td>Mass residue at 500 °C</td>
</tr>
<tr>
<td>MT</td>
<td>Million Tons</td>
</tr>
<tr>
<td>Mw</td>
<td>Molecular Weight</td>
</tr>
<tr>
<td>Acronym</td>
<td>Abbreviation</td>
</tr>
<tr>
<td>---------</td>
<td>--------------</td>
</tr>
<tr>
<td>NEt₃</td>
<td>Triethylamine</td>
</tr>
<tr>
<td>NL</td>
<td>Nitrolignin</td>
</tr>
<tr>
<td>NMR</td>
<td>Nuclear Magnetic Resonance</td>
</tr>
<tr>
<td>OIT&lt;sub&gt;temp&lt;/sub&gt;</td>
<td>Oxidation Induction Temperature</td>
</tr>
<tr>
<td>PA</td>
<td>Propionic anhydride</td>
</tr>
<tr>
<td>PAA</td>
<td>Phenolic Aldehyde Amine</td>
</tr>
<tr>
<td>PBD-(COOH)₂</td>
<td>Dicarboxy-terminated polybutadiene</td>
</tr>
<tr>
<td>PBS</td>
<td>Phosphate buffered saline</td>
</tr>
<tr>
<td>PEG</td>
<td>Polyethylene glycol</td>
</tr>
<tr>
<td>PEI</td>
<td>Polyethyleneimine</td>
</tr>
<tr>
<td>PF</td>
<td>Phenol Formaldehyde</td>
</tr>
<tr>
<td>PhD</td>
<td>Doctor of Philosophy</td>
</tr>
<tr>
<td>phr</td>
<td>Parts per Hundred parts of Rubber</td>
</tr>
<tr>
<td>PS</td>
<td>Polystyrene</td>
</tr>
<tr>
<td>PU</td>
<td>Polyurethane</td>
</tr>
<tr>
<td>PUNL</td>
<td>Nitrolignin-incorporated Polyurethane</td>
</tr>
<tr>
<td>PVC</td>
<td>Polyvinyl chloride</td>
</tr>
<tr>
<td>RCM</td>
<td>Ring Closing Metathesis</td>
</tr>
<tr>
<td>RSM</td>
<td>Response surface methodology</td>
</tr>
<tr>
<td>S</td>
<td>Syringyl</td>
</tr>
<tr>
<td>SAH</td>
<td>Succinic acid anhydride</td>
</tr>
<tr>
<td>SC</td>
<td>Sebacoyl chloride</td>
</tr>
<tr>
<td>SL</td>
<td>Soda Lignin</td>
</tr>
<tr>
<td>SPF</td>
<td>Sun Protection Factor</td>
</tr>
<tr>
<td>SPS</td>
<td>Solvent Purification System</td>
</tr>
<tr>
<td>Abbreviation</td>
<td>Description</td>
</tr>
<tr>
<td>--------------</td>
<td>-------------</td>
</tr>
<tr>
<td>TC</td>
<td>Terphthaloyl chloride</td>
</tr>
<tr>
<td>TCI</td>
<td>Tokyo Chemical Industry</td>
</tr>
<tr>
<td>$T_d$</td>
<td>thermal decomposition temperature</td>
</tr>
<tr>
<td>TDI</td>
<td>2,4-toluene diisocyanate</td>
</tr>
<tr>
<td>$T_{d\text{ onset}}$</td>
<td>Onset decomposition temperature</td>
</tr>
<tr>
<td>$T_g$</td>
<td>Glass transition temperature</td>
</tr>
<tr>
<td>TEGDA</td>
<td>Tetraethyleneglycoldiacrylate</td>
</tr>
<tr>
<td>TGA</td>
<td>Thermogravimetric Analysis</td>
</tr>
<tr>
<td>THF</td>
<td>Tetrahydrofuran</td>
</tr>
<tr>
<td>TRS</td>
<td>Totally Reduced Sulfur</td>
</tr>
<tr>
<td>UATR</td>
<td>Universal Attenuated Total Reflection</td>
</tr>
<tr>
<td>UV</td>
<td>Ultraviolet</td>
</tr>
<tr>
<td>UWF</td>
<td>Untreated Wood Flour</td>
</tr>
<tr>
<td>WCA</td>
<td>Water Contact Angle</td>
</tr>
<tr>
<td>wt%</td>
<td>Weight percent</td>
</tr>
</tbody>
</table>
Dedication

To Ava
Chapter 1

1 General Introduction

1.1 Objective

The overall objective of this PhD project was to synthesize and develop novel advanced functional materials from lignin, a byproduct of the pulp and paper milling industries and an industrial waste material. In this thesis, lignin was chemically modified using a methacrylation process to convert it into a more suitable material for post-modification processes. The methacrylation process was optimized using a response surface method, and the modified lignin was post-modified as part of three different classes of materials: UV-cured coatings, tertiary polyp phosphines, and free-standing polyphosphonium films. The synthesized lignin-based tertiary polyp phosphines reported in this thesis were probed for their metal-scapenging abilities, and the polyphosphonium films were probed as suitable materials for utilization as potential antimicrobial and controlled drug delivery vehicles.

1.2 Background

Almost all carbon-based products such as synthetic polymers currently used are derivatives of petroleum. This high demand for fossil fuel resources, as well as their high prices and their consequently inevitable depletion, along with their detrimental effects on climate change and the environment, have been the main driving points behind the pursuit of renewable, abundant, and comparably clean alternatives. Another critical issue these resources is that their products (e.g. synthetic polymers) are generally resistant to biodegradation, which has resulted in excessive increase of fossil fuel-based waste in the environment. Renewable resources on the other hand have various unique functions and are vastly available, which turn them into viable potential candidates for reducing the cost and improving the properties of existing products derived from non-renewable resources.

Biomass is organic matter available on a renewable basis and can be used to produce fuels, as well as chemicals that are currently being produced from fossil fuel sources. Biomass
also carries significant environmental advantages, such as being carbon neutral and abundant.\textsuperscript{9} More than 220 BT of biomass is produced worldwide each year\textsuperscript{10,11} from different sources such as grains, sugar and oilseed crops, agricultural residues, food processing wastes (liquid and solid), wood and wood chips, bark, mill residues, forestry residues and by-products (such as lignin), animal fats, manures, algae, and exotic crops (i.e. guayule, jojoba, and euphorbia).\textsuperscript{12,13} Each biomass resource, depending on its origin, has varying compositions of three main components: cellulose, hemicellulose, and lignin,\textsuperscript{14} while lignocellulosic biomass (woody biomass and crop residues) containing a significant amount of lignin is of particular interest in this thesis work.

Lignocellulosic biomass is the most readily available renewable source of carbon (next to CO\textsubscript{2}) and can be used to produce various chemicals.\textsuperscript{10,15,16} The term “lignocellulosic biomass”, otherwise known as lignocellulose, refers to biomass that comes from plant dry matter (i.e. woody biomass and crop residues)\textsuperscript{17} and like any other form of biomass, consists of three major components: cellulose (30 – 50%), hemicellulose (20 – 35%), and lignin (15 – 30%).\textsuperscript{18,19}

Cellulose (Figure 1-1A) is perhaps the most well-known constituent of lignocellulose. It is the world’s most abundant biopolymer. It has two main structural parts including a crystalline (organized) component and an amorphous (unorganized) section. Strains of cellulose are linked to each other to form cellulose fibrils (bundles), which are usually independent and bound weakly via hydrogen bonding.\textsuperscript{20,21}

Hemicellulose (Figure 1-1B) is a complex carbohydrate made of different polymers (i.e. pentoses, hexoses, and sugar acids). While cellulose is made of long unbranched fibrils of glucose, the dominant component of each hemicellulose sample can vary depending on its source. For example, the hemicellulose from hardwoods and agricultural plants (e.g. grass and straw) consists mostly of xylan, whereas hemicellulose from softwood is mainly made of glucomannan.\textsuperscript{18,22} One function of hemicellulose is to act as a connection between the cellulose fibres and lignin that increase the rigidity of the cellulose-hemicellulose-lignin network in the plant cell walls.\textsuperscript{20}
Last but not least, lignin is the world’s second most abundant biopolymer. It is a three-dimensional heteropolymer that occurs naturally through an in vivo enzyme-mediated dehydrogenation polymerization process, otherwise known as “lignification.” Following the lignification process and during the cell differentiation of plants, lignin is deposited in the cell wall. Lignin is considered responsible for giving structural integrity to plant cells, as well as contributing to the total stiffness of the plants. Due to its hydrophobicity, lignin plays a crucial role in transporting water and other nutrients inside the plant while protecting it from insect- or pathogen-invasion.
As a byproduct of the pulp and paper milling industry, lignin is today considered an industrial waste, meaning that of the sheer volume of lignin being produced each year, only a small portion (less than 2%) has defined industrial usage and the rest, is either burned as a source of heat or buried in landfill. Facile approaches to utilizing this natural polymer are of utmost importance and can play an important role in the future of sustainable development.

Many reports are available on utilizing lignin as a filler or an additive (without chemical modification) in various systems such as rubber latexes, starch films, and sunscreens. However, the main challenge in such incorporations is that valorization processes that don’t chemically modify lignin before utilizing it in a system, generally can incorporate limited amounts of the material in their respective systems and the reports on this class of incorporation generally utilize a low percentage of lignin in the final material. On the other hand, various studies have focused on adding value to lignin by chemically modifying the macromolecule to make it a better candidate for incorporation in different systems. Several reactions have been used in this research area such as phenolation, esterification, and oxopropylation, all of which aim at replacing an existing component in a specific system with modified lignin and examining the effect of the replacement. Moreover, lignin has also been utilized in advanced functional materials, where many studies have focused on converting the macromolecule into an advanced system with unique and specific properties such as carbon fibres and nanofibers, nanocarriers, and hydrogels.

Despite the headway made in upconverting this natural polymer, the material is still woefully underutilized, which highlight the great need for more research in the area of lignin valorization and the development of facile processes capable of converting lignin into new and highly valuable advanced functional systems such as UV-curable coatings, bio-based metal-scavengers, and antimicrobial and controlled drug delivery vehicles, which are targeted in this PhD project.
1.3 Research Objectives

The overall objective of this PhD project was to develop novel, facile processes for the conversion of lignin into high-value materials. This overall objective was achieved by accomplishing the following tasks:

**Task 1:** Developing a facile methacrylation process for lignin.

Methacrylation was chosen as the desired chemical modification process for the purpose of introducing vinyl groups onto the chemical structure of lignin. Such modification has the ability to convert lignin into a material suitable for post-modification into various useful systems.

**Task 2:** Optimizing the developed methacrylation process.

Response surface methodology and central composite design were used to accomplish this task, which gave the ability to obtain the maximum possible yield of the desired product (methacrylated lignin) under the optimal process parameters, particularly the reaction time, temperature, and the catalyst/lignin molar ratio.

**Task 3:** Incorporating the methacrylated lignin into UV-curable coatings.

Introduction of the vinyl groups onto the chemical structure of lignin gave the ability to incorporate the macromolecule as a crosslinked moiety in UV-curable coatings at high percentages (31 wt%).

**Task 4:** Synthesizing lignin-based polyphosphines.

This task was achieved by performing a hydrophosphination reaction on lignin using phosphine gas, reported for the first time. Utilizing -C-P linkages within the chemical structure of lignin became the first step in synthesizing a new class of advanced functional materials with metal-scavenging abilities from lignin.

**Task 5:** Synthesizing lignin-based polyphosphonium salts.
This task was achieved by further modifying the synthesized lignin-based polyphosphines. The synthesized polyphosphonium salts were crosslinked to obtain UV-cured free-standing films, which showed promise in being used as antimicrobial surfaces and controlled drug delivery vehicles.

1.4 Scope of Thesis

This thesis focuses on developing strategies for converting lignin, a pulp and paper industry waste, into value-added materials.

Chapter 1 provides a general introduction to bioresources, particularly lignin and the research motivation and objectives, as well as the scope of the thesis.

Chapter 2 presents a detailed introduction of lignin, establishing the significant need for more research in lignin valorization, as well as a comprehensive literature review of the most common lignin valorization procedures and research themes in general, identifying knowledge gaps in this research field.

Chapter 3 describes the synthesis of methacrylated lignin as an approach to introducing UV-curability into the structure of lignin, followed by the comprehensive characterization of the final products. This chapter then reports a modified method developed for the quantification of the hydroxyl functionalities within lignin using a halophosphine. The methacrylated lignin preparation procedure, characterization, and analysis data for lignin-incorporated UV-cured coatings containing 10-31 wt.% methacrylated lignin are reported.

Chapter 4 Reports a process optimization study for the synthesis of methacrylated lignin as reported in chapter 3, using response surface methodology, employing a central composite design. Three variables were examined and optimized via the CCD: reaction time, reaction temperature, and catalyst/lignin molar ratio, and their corresponding effects on the recovered mass yield were investigated. The mathematical model derived from the employed CCD was proved accurate in predicting the optimal reaction conditions and consequently, optimizing the recovered mass yield of the lignin methacrylation reaction. The methacrylated lignin synthesized under the optimal reaction conditions was
characterized and proved to be of great promise in a UV-curable coating system at 30 wt.% loading.

**Chapter 5** describes the synthesis of primary and tertiary “lignophines”, lignin-based polyphosphines. Methacrylated lignin was reacted with phosphine gas to prepare a phosphorus-rich bio-based polymer containing PH/PH$_2$ functional groups, which were converted to tertiary phosphine units via the phosphane-ene reaction. The synthesized tertiary alkylated lignophine was then probed for its ability to coordinate to transition metals using silver triflate and subsequently, analyzed for its effectiveness in metal sequestration, where it was shown to effectively sequester the Grubbs I catalyst from a ring closing metathesis reaction system under normal laboratory conditions.

**Chapter 6** reports the synthesis and characterization of “lignophonions”, lignin-based polyphosphonium networks, from lignophines as synthesized in chapter 5. The lignophonium was then incorporated into siloxane-crosslinked UV-cured free-standing films at 15 wt.% and analyzed with regards to its surface-accessible charge density, water contact angle, cure percentage, and swelling degree and showed great potential as antimicrobial and controlled drug delivery vehicles.

**Chapter 7** presents the main conclusions obtained from the research presented in this thesis and suggests future research avenues.

### 1.5 References


(12) Rasmussen, H. Carbohydrate Degradation Mechanisms and Compounds from Pretreated Biomass, Technical University of Denmark, Department of Chemical and Biochemical Engineering, 2016.


Chapter 2

2 Literature Review

In this chapter, lignin is introduced regarding its origin, biosynthesis, chemical structure, isolation, and annual production and demand, highlighting the need for its valorization. Subsequently, current lignin valorization techniques are fundamentally categorized into techniques without and with chemical modification of lignin and are each discussed in depth. In accordance to the interest of this thesis, valorization techniques employing a chemical modification of lignin are categorized into fragmentation/depolymerization, addition of new chemical active sites, and hydroxyl group modification, and several notable reports for each category are introduced and discussed in detail. Finally, lignin-based materials as the most promising lignin valorization avenue are introduced and knowledge gaps in the field of lignin valorization are discussed.

2.1 Lignin

Lignin is a complex, water-insoluble, amorphous, racemic, and aromatic heteropolymer that has higher carbon and a lower oxygen content compared to the holocellulose (cellulose and hemicellulose) fraction of lignocellulosic biomass and thus, can be an attractive candidate as the feedstock for the production of fuels and chemicals. Lignin is primarily derived from three hydroxycinnamyl alcohol monomers (monolignols): \( p \)-coumaryl (2-1), sinapyl (2-2), and coniferyl (2-3) alcohol, which only differ in their degree of methoxylation (Figure 2-1).
Figure 2-1 Three standard monolignols of p-coumaryl alcohol (2-1), coniferyl (2-2), and sinapyl alcohol (2-3) as the building blocks of lignin.

2.1.1 Biosynthesis of Lignin

As the monolignols are incorporated into lignin, they are turned into corresponding phenylpropanoid units (Figure 2-2) of p-hydroxyphenyl (H, from p-coumaryl alcohol), guaiacyl (G, from coniferyl alcohol), and syringyl (S, from sinapyl alcohol).

Figure 2-2 Representative general chemical structures for p-hydroxyphenyl (H), guaiacyl (G), and syringyl (S) groups.

Lignin is deposited in plants cell walls during the cell differentiation process. As a filler material in the cell walls, lignin is vital to plant life as it is responsible for the structural stiffness of the cells and subsequently, the stiffness of the plants.\textsuperscript{4,5}

Depending on the taxon, the cell type, and the individual cell wall layers, the volume, and composition of lignins in different plants can differ, as it can be profoundly affected by environmental and developmental cues.\textsuperscript{6} For example, dicotyledonous angiosperm (hardwood) lignins are composed mainly of G and S units and trace amounts of H units,
while gymnosperm (softwood) lignins mostly consist of units and low amounts of units. On the other hand, monocot lignins (lignins from grasses) are composed of comparable amounts of units and have generally higher levels of units than the dicots. These differences in monomer contents lead to subsequent differences in the bonds that are formed during the formation of lignin, which make the task of determining the exact chemical structure of lignin an extremely difficult and maybe even impossible one.

2.1.2 Chemical Structure of Lignin

The in vivo enzyme-mediated dehydrogenation polymerization of monolignols to produce the cross-linked amorphous chemical structure of lignin is known as the lignification process. The final chemical structure of lignin as produced through lignification can be best described as a highly branched macromolecular one. This structure consists of phenyl propyl (C9) units that have different functional groups (aliphatic and phenolic hydroxyls, carboxylic, carbonyl, and methoxy) at different sites. The C9 units are connected to each other through various (primarily ether and carbon-carbon) linkages (Figure 2-3). The relative amounts of any linkage in any lignin sample are heavily dependent on each particular monomer’s contribution to the lignification process. For example, conifer lignins (mainly composed of units) consist of more β-O-5, 5-5, and 4-O-5 linkages as compared to lignins that are composed primarily of units, mainly because the C5 position is available for coupling in units, whereas it is occupied by a methoxyl group in units building blocks. In general, the β-O-4 (ether) linkage is considered the most common within lignin and about 50% of the bonds formed during lignification are of this type.

Over the years, various characterization methods and spectroscopic techniques have been employed with the purpose of determining the chemical structures of different lignins. However, it is unlikely that the complete chemical structure of any lignin sample will ever be interpreted entirely, primarily due to the unrestrained nature of the lignification process. Nevertheless, it is widely accepted that the majority of linkages within the structure are formed through either the coupling of a monolignol to a growing oligomer, or the coupling of one oligomer to another. On the other hand, it is believed that monomer-monomer coupling is a much rarer event than the other previously described couplings.
during the lignification process due to the more challenging mechanism of monolignol dimerization.\textsuperscript{15,16}

Figure 2-3 Representative general chemical structure of lignin with the most common linkages of β-O-4 (A), α-O-4 (B), 5-5 (C), β-β (D), 4-O-5 (E), β-5 (F), and β-1 (I). Copyright 2014. Reproduced with permission from Elsevier Ltd.

Different conformations can be used to include various structural characteristics of the material according to the purpose of each report. Figure 2-4 shows some of the most common representative structures of lignin proposed in the literature.
Figure 2-4 Most common representative chemical structures of lignin.

2.1.3 Isolation of Lignin

Lignin is a by-product of the pulp and paper milling industries and its isolation methods, although different in technical aspects, all aim to degrade the chemical structure into smaller fragments that can be dissolved in the pulping media. The chemical and physical properties of the obtained lignin depend on the method of isolation used, as changing the isolation method will lead to a change in properties. Some key factors in measuring the effectiveness of a particular isolation method include the pH of the pulping system, the extent of participation of the solvent and solute in the fragmentation of lignin, and the solubility of the lignin fragments in the solvent employed. Sulfur plays a vital role when it comes to categorizing isolated lignins, whereof the four most common isolation processes in use today, sulfite and kraft processes yield sulfur-containing lignins, while organosolv and soda processes lead to non-sulfur-containing lignins. The lignins obtained from any of the mentioned processes are commonly referred to as technical lignins (technicalignins) and differ in properties such as purity levels, chemical structure, and molecular weight.

The sulfite process uses an aqueous solution of sulfite or bisulfite salts with ammonium, magnesium, calcium, or sodium countercations under heat. Approximately 1000 T of
sulfite lignin is currently being produced every year, worldwide.\textsuperscript{19} The obtained lignosulfonates (lignins obtained from the sulfite process) are water-soluble, a property that cannot be found in the other types of technical lignin. The soda process is mainly employed for the pulping of sugarcane bagasse, flax, and all types of non-woody biomass in general. During the soda process, the lignin in the lignocellulosic biomass is dissolved in sodium hydroxide and then recovered through precipitation in acid, maturation of the mixture, and filtration of the precipitate to obtain sulfur-free lignin.\textsuperscript{20} The third most common isolation process, the organosolv (Alcell) process, uses an aqueous-organic solvent mixture (e.g. ethanol, acetone, methanol, acetic acid, or formic acid) to simultaneously isolate hemicellulose, cellulose, and lignin in high purities (<1 wt.% carbohydrate content).\textsuperscript{21} However, Alcell has not been successfully incorporated as a large-scale industrial process to this date, mainly due to the non-optimized recovery methods for the used materials (e.g. solvents), leading to relatively higher process costs when compared to the other isolation processes.\textsuperscript{22}

Finally, the kraft process (kraft pulping) uses an aqueous solution of sodium sulfide and sodium hydroxide (white liquor) to fragment the lignin. The fragmentation is conventionally followed by a three-phase delignification process (separation of the lignin fragments from the cellulose and hemicellulose) occurring in the initial (150 °C, diffusion-controlled), bulk (150-170 °C, chemical reaction-controlled), and final/residual phases, with the most lignin removed during the bulk phase.\textsuperscript{23,24} Over the past decade, several improved separation processes have been reported for the kraft pulping which have significantly enhanced and optimized the global production of kraft lignin. Systems such as Westvaco, LignoBoost, and LignoForce are the major processes developed to improve the production of kraft lignin, among which the LignoBoost system is currently the one employed the most. The LignoBoost system can produce a high purity product through displacement washing. More precisely, in this system lignin is first precipitated in acidic media (CO\textsubscript{2}) and then filtered. Afterwards, the filtered cake is re-dispersed at pH = 2-4 to form a suspension, which is then filtered again and washed through displacement washing to obtain kraft lignin.\textsuperscript{25} On the other hand, The LignoForce system includes an oxidation stage, where the black liquor is oxidized under controlled conditions before acidification. As a result, the precipitated lignin in the acidic media has better filterability. The
LignoForce system has better potential in reducing the TRS (totally reduced sulfur) compounds in the obtained kraft lignin and can also reduce SO$_2$ emissions as compared to the other systems, which has led to increased interest from the pulping industries towards this system in the past few years.$^{26}$

### 2.1.4 Annual Production and Demand

It is estimated that approximately 56.8 BT of biomass is annually produced worldwide and while the primary target application for the produced biomass is the energy sector (biofuels), this production volume is deemed not enough to cover the entire energy needs for sustainable energy production. However, the produced biomass can perfectly cover the world’s chemical needs in terms of feedstocks and thus, developing methods for converting biomass and its resulting feedstocks into materials that are today produced from non-renewable sources is believed to be a viable approach towards sustainability, potentially capable of completely relieving the materials production industries of their dependence on fossil-based resources.$^{27-29}$ As one of the major constituents of biomass and simultaneously the least utilized, it is estimated that approximately 50-55 MT of technical lignin is produced worldwide, annually which highlights the role of lignin valorization (adding value) in sustainable development.$^{30,31}$

The major source of technical lignin in North America is kraft lignin. As the main internal source of energy, kraft pulp mills fire the black liquor into the recovery boiler to generate energy and recover the pulping chemicals, which has over the years helped them become completely independent and sustainable in terms of operational energy needs. However, over the past 30 years, the production at these mills have outgrown the increase in their need for energy, which has led to the recovery boiler becoming the production bottleneck and in constant need of offloading. Thus, these pulp mills have been removing tons of kraft lignin from the black liquor in their recovery boilers as means of offloading them. It has been determined that a typical kraft pulp mill can produce an additional ton of pulp for every ton of lignin that is removed from the recovery boiler$^{26}$ and given that the kraft pulp mills around the globe are producing more than 130 MT kraft pulp annually,$^{32}$ kraft lignin has now become part of the product stream at many kraft pulp mills, especially in North America. This highlights the importance of discovering viable methods to convert kraft
lignin into materials of higher value (value-added materials). Valorizing lignin as a sustainable bio-based material will not only have significant environmental benefits; it will also add a new and valuable stream of income to the kraft pulp mills and can eventually be the potential next leap into a sustainable future.

2.2 Valorization of Lignin

The current strategies for valorizing lignin can be best categorized based on whether or not chemical modification (by using in a chemical reaction) is implemented. Scheme 2-1 describes the general global algorithm for the valorization of lignin.

**Scheme 2-2-1** General scheme of current valorization processes employed for lignin.\(^{33}\) Copyright 2014. Reproduced with permission from Elsevier Ltd.

2.2.1 Valorization of Lignin Without Chemical Modification

Historically, valorization strategies in which lignin did not undergo chemical modification have been around for much longer than their counterparts (strategies that included chemical modification). Because of the unique physical and chemical properties of lignin, the first attempts at adding value to it focused on physical-blending as a bulk material. Properties such as hydrophobicity, thermal stability, thermo-oxidation resistance, UV-absorption, and having a highly crosslinked structure were the major ones that made lignin a suitable component as a filler in such blends.

Reports on utilizing lignin as a filler/additive material date back to mid-1900s, where in 1948 Keilen and Pollak\(^{34}\) incorporated up to 38.5% lignin into natural and synthetic rubber
latexes via coprecipitation and reported that the resulting rubbers showed superior abrasion resistance, tensile strength, and tear resistance in comparison rubber made with conventional inorganic fillers. Years later, in 1998, Baumberger et al.\textsuperscript{35} used lignin as a filler in starch films to overcome the high swelling degrees and the partial dissolution of starch films in a moist environment, caused by the hydrophilicity of starch. As a naturally hydrophobic filler, up to 30 wt.% lignin was incorporated into the starch films through mixing with wheat starch by extrusion and subsequent thermal molding or casting of the films. They reported that incorporation of up to 20% lignin into starch films at 58% relative humidity had a negligible effect on film elongation and stress at break, whereas it reduced the water affinity of the prepared films noticeably. However, they also determined that at 71% relative humidity and up to 30% incorporated lignin, the resistance to elongation decreased significantly in the prepared films, likely due to the incompatibility between the hydrophilic starch and the hydrophobic lignin components.

Lignin has also been probed as an excellent antioxidant filler/additive. Kosikova et al.\textsuperscript{36} reported that using up to 30 phr (parts per hundred parts of rubber) of lignin as a filler in natural rubber-based composites significantly improved their resistance to thermooxidative degradation. Moreover, they reported that other physiochemical properties such as elongation, 100% modulus, and the tensile strength at break were improved in the prepared composites by the addition of lignin. Moreover, Sadeghifar and Argyropoulos\textsuperscript{37} determined that the antioxidant properties of lignin are highly correlated with the number of phenolic hydroxyl groups on its chemical structure, due to the free radical-scavenging nature of these groups. They utilized lignin samples containing different amounts of phenolic hydroxyl functionalities as fillers in polyethylene blends to determine the effect of such combinations on the oxidation induction temperature (OIT\textsubscript{temp}) of the prepared blends. Consequently, they discovered that using acetone soluble softwood kraft lignin (ASKL) as the filler in the polyethylene blends increased their OIT\textsubscript{temp} by 50 °C in comparison to when acetone insoluble kraft lignin (AIKL) was used and concluded that lignin samples with higher phenolic hydroxyl contents showed significantly improved antioxidant properties, as the ASKL contained 54% more phenolic units than the AIKL sample.
Lignin has also been utilized as an additive in sun blocker materials. With chromophores such as phenolic and ketone functionalities present in its chemical structure, lignin can act as a natural broad-spectrum sun blocker. Qian et al.\textsuperscript{38} reported that lignin could be utilized as an additive (2-10 wt\%) suitable for commercial sunscreens. They blended dry, finely powdered lignin with pure creams (NIVEA Cream-N moisturizing cream and LIFE glycerin hand cream) at 1000 rpm for 24 h and measured the UV-transmittance and the sun protection factor (SPF) of the obtained lotions in comparison to commercial sunscreen (BIOTHERM and LIFE SPF 15, 30, and 50) lotions. They reported that while the blend of pure cream and 10 wt\% lignin had higher UV-transmittance than that of LIFE SPF 15 sunscreen, it showed lower transmittance in the UVA area (especially at 385-400 nm) which, along with the fact that the lignin sunscreen at 10 wt\% lignin contained less active ingredient than the SPF 15 sunscreen, were taken as evidence towards the suitability of lignin as a natural candidate for broad-spectrum sunscreens. Furthermore, they reported that adding 2 wt\% of lignin to SPF 15 commercial sunscreen lotions resulted in their sunscreen effect to increase to that of SPF 30 sunscreen lotions of the same brands. Finally, adding 10 wt\% lignin to the SPF 15 lotions resulted in them out-performing the sunscreen effects of SPF 50 sunscreen lotions of the same brands. Thus, they concluded that lignin can be probed both as the sole active ingredient or in parallel with other active ingredients in commercial sunscreen lotions. However, lignin has a dark-brown color, which can potentially have an adverse effect on developing a commercial lignin-mixed sunscreen lotion, which might very well be the main reason why lignin has not yet been utilized in sunscreen lotions on a commercial level.

Lignin has also been utilized as an additive to improve the flame retardance of target polymer networks. With an aromatic ring-rich chemical structure, lignin shows relatively high char formation and thus, has been utilized as an additive to promote char formation in different blends. De Chirico et al.\textsuperscript{39} examined the effect of using lignin as an additive in polypropylene blends on the final material’s flame retardancy. They used thermogravimetric analysis (TGA) to compare the char formation in polypropylene networks containing known flame retardants (melamine phosphate, ammonium polyphosphate, aluminum hydroxide, and poly(vinyl alcohol)) to the same values in polypropylene networks containing known mixtures of the mentioned flame retardants and
lignin. Thus, they discovered that adding up to 15 wt% lignin to polypropylene networks that already contain one of the mentioned flame retardants, improved the char formation in the resulting networks by up to 6 wt%. Furthermore, they reported that the networks that were only incorporated with lignin had higher char formations compared to when melamine phosphate, aluminum hydroxide, or poly(vinyl alcohol) was used, although they showed a lower char formation than when ammonium polyphosphate was used in the blends.

Although promising, the utilization of lignin into existing systems as an additive, without any chemical modification, has been proved to be somewhat limited in terms of target applications and/or only applicable to blends with relatively low sensitivity (e.g. rubbers) and not capable of adding refined properties to the final products. These limitations amplify the need to modify lignin into a compound that can be incorporated into the backbone of various systems in large percentages and thus, take full advantage of the unique properties and the rich chemical structure of this natural polymer.

2.2.2 Valorization of Lignin with Chemical Modification

Given the interesting chemical structure of lignin and the abundance of different functional groups on its structure, many researchers have examined modifying such groups by chemical reactions. Overall, the examined approaches toward chemically modifying lignin can be divided into 3 main categories of fragmentation/depolymerization, synthesis of new chemically active sites, and hydroxyl group modification which are herewith discussed in detail.

Fragmentation/Depolymerization

Lignin has been the subject of several fragmentation/depolymerization processes, e.g., hydrolytic/reductive/oxidative de-polymerization and pyrolysis, mainly to produce lower molar mass chemicals and/or fuels.

Among all the reported fragmentation processes for lignin, pyrolysis is the most studied. Generally, pyrolysis refers to the thermal treatment (thermolysis) of an organic compound in the absence of air, with or without a catalyst, to break down its molecular structure into
smaller substructures while disallowing further combustion to carbon dioxide. Depending on the pyrolysis conditions, the dominant product from pyrolyzing lignin can be a liquid (pyrolysis oil) or a gas and increasing the time and temperature of pyrolysis will typically cause lignin to fragment into components with lower molecular weights.\textsuperscript{40,41} Several factors such as feedstock’s origin\textsuperscript{42}, reaction temperature\textsuperscript{43}, heating rate, and the use of additives\textsuperscript{44} can affect the pyrolysis of lignin. Nonetheless, lignin pyrolysis produces gaseous hydrocarbons, carbon monoxide, carbon dioxide, volatile liquids (e.g. acetone, acetaldehyde, and methanol), monophenolic compounds (guaiacol, syringol, cathecol, and phenol) and several other polysubstituted phenols as its products.\textsuperscript{45} The liquid product of pyrolysis, i.e., pyrolysis oil, typically consists of about 20\% aqueous compounds (acetone, acetic acid, methanol, and water) along with about 15\% tar (condensed volatiles yielding phenolics, primarily).\textsuperscript{46} Figure 2-6 depicts the primary aromatic compounds that can be obtained from the pyrolysis of lignin.
Figure 2-5 Major products obtained from the pyrolysis of lignin: guaiacol (A), methyl guaiacol (B), syringol (C), methyl syringol (D), vanillin (E), syringaldehyde (F), vinyl syringol (G), vinyl guaiacol (I), and phenol (K).47

In addition to the mentioned products, a group of thermally stable fractions, commonly known as char, can also be obtained from the pyrolysis of lignin, the yield of which has a reverse correlation with the pyrolysis temperature.48

The pyrolysis of lignin happens over a wide range of temperatures (100-900 °C)49 with different products being yielded at different temperature ranges. For instance, formic acid, formaldehyde, carbon dioxide, carbon monoxide, and water can be obtained at a relatively lower temperature range (120-300 °C) as compared to the other pyrolysis products. The type of obtained products can be attributed to the type and subsequently the strength of the bonds being cleaved during the pyrolysis. For example, aryl ether linkages (e.g. β-O-4) are
relatively easier to break during thermal treatment than ether linkages at γ carbons, which are themselves more easily broken than methoxyl groups. While every lignin sample follows the same general trend in terms of the obtained pyrolysis products, the yield of each product and the specific temperature range for obtaining it differs depending on the type of lignin used. An excellent example for temperature-specific pyrolysis products was reported by Liu et al. where they used coupled TGA-FTIR to determine the temperature ranges of different pyrolysis products of birch and fir strong acid detergent fiber lignins. They reported that at around 100 °C, carbon dioxide and water were obtained as the first products of the pyrolysis process for both samples, followed by the production of carbon monoxide, formic acid, aldehydes, and phenols, along with more water and carbon dioxide at around 225 °C. The vast amounts of monomeric phenols obtained at 225 °C were attributed to the breakage of ether linkages and although the same products were obtained for both samples at this temperature, they observed a more obvious peak for carbon monoxide in birch lignin compared to fir lignin, which they took as evidence toward the lateral chains being more easily broken in the birch lignin samples. Furthermore, they reported that hydrocarbons (primarily methane), methanol, carbon monoxide, and carbon dioxide were the major products at temperatures around 375 °C for fir lignin and 425 °C for birch lignin.

Principally, pyrolysis can be categorized into conventional (slow) and flash pyrolysis in terms of the heating rate. In slow pyrolysis, such as the one reported by Liu et al. earlier, the heating process of lignin happens at a much lower rate than in flash pyrolysis, with a vapor residence time varying anywhere from 5 to 30 minutes and thus, the components in the vapor phase of slow pyrolysis will have more time to react with each other and subsequently, form more char and gaseous components. On the other hand, the higher heating rates of flash pyrolysis lead to vapor residence times of 2 seconds, which make it possible to produce a predominantly liquid product (bio-oil) with a yield of up to 75 wt%. The generated bio-oil and char can both be employed as fuels, and the produced gas can be recycled back into the process. However, despite the high interest in pyrolysis and especially flash pyrolysis over the past 20 years, it has yet to become a viable process for valorizing lignin, as the produced bio-oil contains significant amounts of oxygen-
containing compounds that are corrosive, less stable, and contain a lower heating value. Thus, pyrolysis has yet to become a valorization process capable of taking full advantage of the complex chemical structure of lignin in its entirety.43,53–55

Oxidation is another well-studied fragmentation method of lignin, primarily for obtaining phenolic compounds of lower molar masses, such as phenolic aldehydes (vanillin and syringic) and their acids (vanillic and syringic).56 Several oxidants have been reported to effectively oxidize lignin into such compounds, including oxygen, air, nitrobenzene, and metal oxides. Vanillin (4-hydroxy-3-methoxybenzaldehyde) is the most valuable product that has been obtained from the oxidation of lignin, and successful high-yield vanillin isolation has over the years been the primary goal of most lignin oxidation studies.57–61

Villar et al.62 reported that the oxidation of lignin at 170, 180, and 190 °C under oxygen pressures of 10, 12, and 15 atm yielded syringaldehyde, vanillin, syringic acid, and vanillic acid with the total yields ranging between 4-11%. They reported the maximum conversion of 11% at 190 °C under 12 atm of oxygen. Furthermore, they examined the same oxidation processes in the presence of Co(II) salen (N,N’-bis(salicynidene)ethylenediamine monohydrate), CuSO₄.5H₂O, CoCl₂, and two commercial platinum-alumina (referred to as A and B due to their proprietary compositions) catalysts and determined that the addition of catalysts did not have a positive effect on fragmentation into the mentioned products, particularly the aldehydes. Thus, they concluded that at the optimal oxidation conditions under oxygen, the maximum yield of aldehydes was approximately 4%. In a separate study, Villar et al.63 examined the oxidation behavior of kraft lignin using nitrobenzene and copper(II) oxide at 170, 180, and 190 °C for 10-120 minutes. For nitrobenzene, oxidant/lignin ratios (v/w) of 6.3, 9.4, and 12.6 were examined, and CuO/lignin molar ratios of 1.33, 2.66, and 3.99 were chosen for oxidations using copper(II) oxide. They reported that at the optimal oxidation conditions, lignin conversion was 23% with nitrobenzene (14% aldehyde yield) and 13% with copper(II) oxide (8% aldehyde yield). Optimal oxidation conditions as reported in this study were 40 min, 190 °C, and 6 (mL nitrobenzene/g lignin) for nitrobenzene and 70 min, 190 °C, and 4 (mol CuO/mol lignin) with copper(II) oxide.
While the oxidation of lignin has proven to be especially viable for producing aldehydes, specifically vanillin and syringaldehyde, both the total conversion of lignin to fragmented molecules and the aldehyde yields remain at relatively low percentages to date, which means that most of the lignin incorporated in such oxidation processes will still be depleted as process waste. Thus, much like pyrolysis, oxidation is yet to become a viable process capable of converting lignin to positive-value products at high conversion percentages.

More recently the Xu group developed highly efficient processes for depolymerization of kraft lignin based on hydrolytic reactions or reductive reactions. Kraft lignin was effectively depolymerized into polyols of moderately high hydroxyl number and yield with moderately low weight-average molecular weight (Mw) via direct hydrolysis using NaOH as a catalyst. The best operating condition was found to be at 250 °C, 1 h, and NaOH/lignin ratio ≈ 0.28 with 20 wt% substrate concentration, leading to <0.5% solid residues and ~ 92% yield of depolymerized lignin (aliphatic-hydroxyl number ≈ 352 mgKOH/mg and Mw ≈ 3310 g/mole), suitable for replacing polyols in polyurethane foam synthesis. In the latter work, formic acid was employed as an in-situ hydrogen donor for the reductive de-polymerization of kraft lignin. Under the optimum operating conditions, i.e., 300 °C, 1 h, 18.6 wt.% substrate concentration, 50/50 (v/v) water-ethanol medium with formic acid at a formic acid-to-lignin mass ratio of 0.7, kraft lignin (Mw ~10,000 g/mol) was effectively de-polymerized, producing de-polymerized lignin (Mw 1270 g/mol) at a yield of ~90 wt.% and <1 wt.% yield of solid residue.

Addition of New Chemical Active Sites

The generally high waste production and relatively low yield of the fragmentation processes, plus their inability to benefit from all that the chemical structure of lignin can offer, has led to much interest being placed on modifying the functional groups present on lignin, namely hydroxyl, methoxy, carbonyl, and carboxyl groups, with the purpose of increasing its reactivity. Adding/synthesizing new chemical active sites through modifying lignin can enhance its incorporation behavior in various systems. This section will address the studies on reactions employed for modifying the functional groups on the structure of lignin, namely alkylation/dealkylation, hydroxyalkylation, amination, and nitration.
Chemical modification of the hydroxyl groups of lignin will be discussed in a separate section, due to the relatively higher importance of this type of modification and its closer relevance to this thesis.

Alkylation of lignin is an essential class of chemical modification employed for this material. Most alkylation processes include pre-treatment of the lignin samples to fractionate their structure and decrease their molecular weight to increase their solubility in common organic solvents.\textsuperscript{66,67} On the other hand, a lignin dealkylation process was developed by Gaylord Chemical (US)\textsuperscript{68}, in which the demethylation of lignin is employed for producing dimethylsulfoxide (DMSO). During this process, lignin is reacted with molten sulfur in basic media, and dimethyl sulfide (DMS) is synthesized via the transfer of two methyl groups from lignin to sulfur, which is then oxidized using nitrogen dioxide to produce DMSO (Scheme 2-2).\textsuperscript{68}

\begin{center}
\includegraphics[width=\textwidth]{Scheme2-2.png}
\end{center}

**Scheme 2-2** Synthesis of dimethylsulfide (DMS) from lignin and molten sulfur and subsequent oxidation to dimethylsulfoxide (DMSO).\textsuperscript{69} Copyright 2010, reproduced with permission from John Wiley and Sons.

However, as shown in Scheme 2-2, the only remaining components of lignin in the final product of this process are the two methyl groups that were used in the synthesis of dimethylsulfide and DMSO, most of the lignin used in this process is still being disposed of as process waste. Other processes are today available that aim to utilize demethylated lignin in a less wasteful manner. For example, Liu \textit{et al.}\textsuperscript{70} prepared a formaldehyde-free water-resistant wood adhesive, using alkaline demethylated kraft lignin (DKL) and polyethyleneimine (PEI), where the demethylated lignin was prepared by suspending DKL
in water, dissolving the suspension in sodium hydroxide, and freeze-drying the obtained solution. DKL-PEI wood adhesive samples were then prepared through mixing the DKL (14-25 wt%) and PEI in water for 2-120 minutes. It was reported that at DKL percentages higher than 25 wt%, the resulting adhesives became too viscous to be uniformly applied to the wood’s surface. Afterwards, the adhesive was applied to two layers of maple veneer boards, lapped together, and cured using a hot-press. It was reported that at the DKL percentage of 20 wt% and PEI samples with $M_w = 750,000$, with the hot-press duration of 5 minutes at 120 °C, the shear strength of the bonded lap-shear specimens (calculated using the maximum force at breakage for each bonded lap-shear sample) was the highest (7 MPa). This report, despite the multi-step pre-treatment procedure for preparing the DKL and the relatively low weight percentages of the demethylated lignin in the adhesives, can be counted among the more viable approaches to valorizing lignin and can also highlight the potential benefit of chemically modifying lignin as the first step in incorporating it in different systems.

Apart from the several reports available on incorporating dealkylated lignin in wood adhesives, such as the one discussed earlier by Liu and Li,70 the abundance of phenolic groups on lignin has made it a desirable substitute for the non-renewable phenol in phenol formaldehyde (PF) resins and other adhesives.71–73 Notably, the structural similarities between lignin and phenol, along with their comparable reactivities with formaldehyde, marks an interesting potential application for lignin in the production of lignin-phenol-formaldehyde (LPF) resins.

Alonso et al.74 hydroxymethylated (methylolated) softwood ammonium lignosulfonate by dissolution in an 8% NaOH solution at 80 °C, followed by reaction with formaldehyde at room temperature (Scheme 2-3). They later probed the ability of this metholylated lignosulfonate to partially substitute the phenol in PF resins through a comparative study with a commercial resol phenolic resin. They determined that the maximum possible percentage of incorporated methylolated lignosulfonate in the PF resins that would keep different properties of the prepared LPFs (pH, free phenol, viscosity, and gelation time) comparable to the commercial PF resin, and in compliance with the requirements for utilization in plywood manufacture, was 35%.75
Amination is another class of reactions performed on lignin to introduce new chemical active sites on its structure. Principally, the amination of lignin is carried out using the Mannich reaction. Matsushita and Yasuda used 1-guaiacyl-1-<i>p</i>-hydroxyphenylethane as a model compound in a reaction with dimethylamine and formaldehyde in acetic acid and dioxane and reported that as resulted by the Mannich reaction, the aminoethyl group was introduced onto the model compound at the ortho position to the phenolic hydroxyl groups, giving 4 different products (Scheme 2-4), proposing that the same procedure can be utilized for the amination of lignin itself.

**Scheme 2-3** Hydroxymethylation of lignin as reported by Alonso <i>et al.</i>

**Scheme 2-4** Representation of the amination of the lignin model compound 1-guaiacyl-1-<i>p</i>-hydroxyphenylethane with formaldehyde and dimethylamine as reported by Matsushita and Yasuda. 4 products were isolated from this reaction: \( R_1 = -\text{CH}_2\text{N(CH}_3)_2 \), \( R_{2,3} = \text{H} \); \( R_{1,2} = \text{H} \), \( R_3 = -\text{CH}_2\text{N(CH}_3)_2 \); \( R_{1,3} = -\text{CH}_2\text{N(CH}_3)_2 \), \( R_2 = \text{H} \); \( R_{1,2,3} = -\text{CH}_2\text{N(CH}_3)_2 \). Yue <i>et al.</i> reacted fractionated softwood soda lignin (SL) with diethylenetriamine (DETA) and formaldehyde at a controlled pH of 11.5 to obtain lignin amine. Afterwards,
they examined the effects of using the lignin amine in the surface treatment of wood flour by preparing similar PVC composites from untreated wood flour (PVC-UWF), lignin amine-treated wood flour (PVC-LWF), and aminosilane-treated wood flour (PVC-KWF). They reported that the optimal lignin amine loading percentage for the PVC-LWF composites was at 2%, where the tensile strength in the PVC-LWF composites increased from 35 MPa to 42.5 MPa and their impact strength increased from 27.5 kJ/m² to 37.5 kJ/m². However, they determined that using aminosilane, the same increases can be achieved in the PVC-KWF composites at 1% loading. Furthermore, they evaluated the water absorption behaviors of the composites after 10 days immersion and determined that again, while both lignin amine and aminosilane decreased the weight gain of the composites by ca. 3%, similar decrease in weight gain was observed at 1% aminosilane loading and 2% lignin amine loading. Thus, based on the data reported by Yu et al. aminosilane outperforms lignin amine as the surface treatment agent. In addition to the low loading percentages generally reported for utilization of aminated lignin, to the best of our knowledge, no report is to date available on amination and utilization of non-fractionated lignin.

The last major class of chemical active site synthesis reactions of lignin is nitration. However, fewer reports are available on this method compared to the previously discussed ones.⁷⁹,⁸⁰ Zhang and Huang⁸¹ synthesized nitrolignin (NL) in the form of a “reddish-brown powder” through reacting alkali lignin with nitric acid and acetic anhydride and determined the nitrogen content of the synthesized NL sample to be at 6.32% using elemental analysis. They subsequently incorporated the synthesized NL samples at weight percentages from 1.4-6 wt% into polyurethane networks (PUNL) and determined that at 2.8% NL loading, both the tensile strength and the breaking elongation of the PUNL networks were 2 times higher than that of standard polyurethane (PU) networks. Loadings higher than 2.8% however, resulted in a decrease in the mentioned values. Much like amination, the product of the nitration of lignin is yet to be probed as a viable nitrogen-containing replacement with high loadings in materials.

The route of synthesizing new chemical active sites on lignin poses the potential for producing chemically modified lignin with the ability to be utilized in some common
industrial materials, mainly through dealkylation and hydroxymethylation of lignin,\textsuperscript{70,75} as these two reactions have shown promising results with acceptable loading percentages, while the other reported chemical modifications either use lignin as a source for functional groups and treat the rest of the macromolecule as waste,\textsuperscript{68} or fractionate the structure and employ only a fraction of it,\textsuperscript{78} or are not yet able to utilize the chemically modified lignin in the target systems at appreciable loading percentages.\textsuperscript{79–81} However, when it comes to chemical modification, there is another functionality within the chemical structure of lignin with considerably higher reactivity than the other functional groups discussed, the hydroxyl groups.

*Hydroxyl Group Modification*

As the most reactive sites within the structure of lignin, the various types of hydroxyl groups have been the subject of numerous studies.\textsuperscript{82–86} With the densely crosslinked structure of lignin and its subsequent relatively limited solubility in common organic solvents, such modifications can positively affect the solubility and thus, the reactivity of lignin in these solvents and provide potential routes for incorporating this sustainable natural polymer, in its entirety, in different supramolecular systems.\textsuperscript{12,87,88} Processes such as esterification, phenolation, and etherification, and the corresponding incorporations of the modified lignins in materials (where applicable) are discussed in this section as the most promising approaches toward this goal.

Esterification reactions are perhaps the most straightforward chemical modifications that can be employed for lignin, as these reactions often require relatively mild conditions to produce esterified lignin. Commonly, acyl anhydrides, acyl chlorides, and acidic compounds are used as the modifying reagents in these types of reaction, using a base (e.g. triethylamine, 1-methylimidazole, and potassium hydroxide) in catalytic amounts. As a result, numerous reports are available regarding the esterification of the hydroxyl functionalities within lignin.\textsuperscript{89–97}

Guo and Gandini\textsuperscript{98} esterified kraft lignin using two different diacyl chlorides, sebacoyl chloride (SC) and terphthaloyl chloride (TC), and triethylamine as the base, followed by copolymerization with polyethylene glycol (PEG) to obtain TC-based and SC-based lignin
copolymers. They conducted the first step of the process (esterification) at 0 °C for SC and 60 °C for TC, followed by the second step (copolymerization) at 120-130 °C for SC and 180-190 °C for TC (9 h) to obtain the corresponding copolymers with 15, 25, and 35 % lignin loadings. They reported that a portion of the obtained products was pyridine-soluble (used for extraction) and thus, that portion proved to be extremely difficult to remove and could not be included in the reported characterizations, as well as the reported yields. For the extracted portion, however, they reported that both the TC- and the SC-based copolymers containing 25 % lignin were stable at up to 300°C, which was higher than the same values for the initial lignin and the lignin copolymers without PEG (reported in a separate study by Guo et al.99). Furthermore, they reported that increasing the loading percentage of lignin in the initial reaction, increased the yield of the obtained copolymers, whereas the yields for the TC-based copolymers were 25, 45, and 60% using 15, 25, and 35% lignin loadings, respectively. The same values were 45, 60, and 65% yield for SC-based copolyester using 15, 25, and 35% lignin loadings, respectively. They also reported that increasing the lignin loadings increased the observed Tg’s for both the TC- and the SC-based copolymers.

In a report by Fang et al.,100 dimerized acid (DA, Scheme 2-5) was synthesized by dimerizing unsaturated fatty acids (derived from tall oil, cottonseed oil, and soybean oil). DA was then reacted at 50 °C with enzymatically hydrolyzed lignin (EHL) using a cation exchange resin (CER) as the catalyst for 5 h to synthesize a graft copolymer, DA-g-EHL, through an esterification reaction between DA and EHL.

![Scheme 2-5 Esterification of EHL with DA as reported by Fang et al.](image)

Afterward, they employed the DA-g-EHL graft copolymer in the preparation of modified phenolic aldehyde amine (PAA) as a partial replacement for phenol. Specifically, they
replaced 25 wt% of the phenol in the PAA samples with either the DA-g-EHL or the EHL to obtain two samples and used a commercial PAA sample as the control. Subsequently, they used the three samples in 20% loadings as curing agents for epoxy resins. The epoxy resins were thereafter cured at 86 °C for 3 h. They determined that the bending strength and the elongation at break were significantly higher in the epoxy resins cured by DA-g-EHL modified PAA than in the ones cured with EHL modified PAA, which they attributed to the long aliphatic chain, and subsequently several possible conformations of the DA, which when grafted onto the EHL can improve the flexibility of the cured epoxy resins. The bending strength at break observed for the epoxy resins cured with DA-g-EHL modified PAA (89.18 MPa) was almost equal to that of the ones cured with the commercial PAA (90.46 MPa) and higher than that of the ones cured with EHL modified PAA (73.87 MPa). The elongation at break, however, was higher in the case of DA-g-EHL modified PAA (7.75 %) than it was for the EHL modified PAA (5.77 %) and the commercial PAA (6.62 %). Moreover, they determined that, because of the long chains of DA reducing the rigidity of EA and the chain stiffness of the modified PAA, epoxy resins cured with DA-g-EHL modified PAA had a higher tensile strength (36.22 MPa) than the ones cured with EHL modified PAA (34.41 MPa), yet still lower than that of the ones cured with the commercial PAA (39.55 MPa).

Gordobil et al. esterified lignin with methacryloyl chloride (MC) using triethylamine as the catalyst at 14 °C (Scheme 2-6). Acyl chlorides are highly susceptible to homopolymerization, and thus, they used hydroquinone as an inhibitor to prevent such a phenomenon. They used 4 different lignin samples of organosolv eucalyptus lignin, kraft eucalyptus lignin, organosolv spruce lignin, and kraft spruce lignin in the mentioned esterification process and reported recovered mass yields of 118, 131, and 165% for organosolv eucalyptus lignin, organosolv spruce lignin, and kraft spruce lignin, respectively. They reported that yield for kraft eucalyptus lignin was lower than the amount of starting material, which they attributed to the high ash content (22.4%) of this sample. However, the high amounts of obtained final products can also be attributed to incomplete removal of unreacted reaction elements, as they reported solely using distilled water for isolating the final products.
Saito et al.\textsuperscript{102} reported an esterification reaction with dicarboxy-terminated polybutadiene (PBD-(COOH)\textsubscript{2}) with potassium hydroxide as the catalyst on three different lignin samples of hardwood kraft lignin (A-lignin), methanol extracted hardwood kraft lignin (W-lignin), and formaldehyde-crosslinked methanol extracted hardwood kraft lignin (F-lignin). The reaction was carried for 24 h at 100 °C to yield corresponding lignin-polybutadiene copolymers, isolated through precipitation in methanol (Scheme 2-7). They determined that a mixture of three products were possibly obtained from this reaction as lignin bridged with telechelic butadiene via covalent bonding (Scheme 2-7A), lignin bridged with telechelic butadiene via ionic bonding (Scheme 2-7B), and formation of an interpenetrating network (Scheme 2-7C) due to the presence of K\textsuperscript{+} cations.
Scheme 2-7 Esterification of lignin with dicarboxy-terminated polybutadiene yielding products with ester linkages (A), ionic linkages (B), and interpenetrating network (C).

They reported that the products of the esterification reactions gave copolymers in the forms of a free-standing solid for F-lignin-PBD(COOH)$_2$, a highly viscous liquid for W-lignin-PBD(COOH)$_2$, and a viscous liquid for A-lignin-PBD(COOH)$_2$. Thus, the F-lignin-PBD copolymer was taken as the only product capable of being utilized as a coating. They attributed this property to the fact that due to the methanol washing and the reaction with formaldehyde, the F-lignin samples contained a higher portion of a high molecular weight lignin fraction (as methanol extraction removes the lower molecular weight fractions and the reaction with formaldehyde crosslinks the phenolic groups) and thus, the F-lignin sample facilitated the formation of a more continuous network with the PBD bridges, while the broad distribution of molecular weight in A-lignin resulted in the formation of a less continuous network structure.
Among the various esterification reactions reported for lignin, one reaction, reported by Thielemans and Wool,\textsuperscript{103} is of particular interest to this thesis. They esterified kraft lignin using 5 different anhydrides: acetic anhydride (AA), propionic anhydride (PA), \(n\)-butyric anhydride (BA), maleic anhydride (MA), and methacrylic anhydride (MethA) to obtain corresponding esterified lignins. More specifically, they performed the reactions between lignin and AA, PA, and BA without solvents and using lignin:anhydride molar ratio of 1:2 with 1-methylimidazole (1MIM, 0.5 mL/g lignin). For MA and MethA however, they used 1,4-dioxane as the solvent. For MA, they used 1 mL of a 0.5 g of 1MIM in 10 mL solvent (per every gram of lignin), and for MethA, 6.1 mL of a 60-to-1 (MethA-1MIM by weight) mixture was used as a catalyst. lignin:solvent w/w ratios were 1:2 for MA and 1:4 for MethA (Scheme 2-8). They reported that, except for MA, all the anhydrides were able to successfully cap both the aliphatic and the aromatic hydroxyl groups on lignin. In the case of esterification with MA, they determined, through \(^1\)H NMR, that only the aliphatic hydroxyls were selectively capped, and the aromatic hydroxyls remained unreacted. The broad range of anhydrides employed in this report, along with the relatively straightforward reaction conditions, make it one of the most interesting studies on lignin modification. However, no discussion was made on quantifying the extents of the esterifications, determining the optimal reaction conditions, or peak assignments (on cases such as the one with MA as mentioned before) in this report.
Scheme 2-8 Synthesis of acetylated (A), propionated (B), butyrate (C), maleated (D), and methacrylated (E) kraft lignin through esterification with corresponding anhydrides.
Esterification, as discussed earlier, is perhaps the most clear-cut chemical modification process available for the hydroxyl groups on lignin. However, other hydroxyl-functionalization reactions are employed that aim to lead to new applications for lignin.

Phenolation (phenolysis) is another hydroxyl functionalization reaction for lignin. Phenolation can be carried out in acidic media, where lignin is reacted with phenol to condensate the aromatic rings and the side chains on its structure. This process is particularly essential for lignosulfonates, as it can serve to increase the number of phenolic hydroxyl groups and thus, increase their susceptibility to chemical modification.¹⁰⁴

Tan¹⁰⁵ used cardanol, a natural alkyl phenol derived from cashew nut shell liquid, as the phenol source in the phenolation of softwood kraft lignin to produce cardanol-lignin (CL) (Scheme 2-9). They then reacted the CL sample with 2,4-toluene diisocyanate (TDI) using stannous octoate as the catalyst to obtain cardanol-lignin-based polyurethane (CLPU) films. Furthermore, they determined that increasing the CL amount in the CLPU films (up to 30 wt%) increased the crosslinking density of the films, while loadings higher than 30 wt.% had an adverse effect on the crosslink density. They attributed this observation to the increase in the functionality level of the film formulations with increasing the CL amount. The decrease in the crosslink density after 30 wt% CL incorporations, however, was attributed to the increased solvent compatibility, caused by the long alkyl chains (R) on the CL samples at loading percentages higher than 30 wt%.

Scheme 2-9 Phenolation of kraft lignin using cardanol as reported by Tan.¹⁰⁵
Etherification of the hydroxyl functionalities on lignin with alkylene oxides is another well-studied modification process available. In the case the alkylene oxide in use is propylene oxide, the etherification of lignin is referred to as “oxypropylation,” which is the most well-known lignin etherification reaction and one of the most extensively-studied chemical modification reactions for lignin overall.\textsuperscript{106} The importance of oxypropylation is highlighted in the case of incorporating lignin into polyurethane, as this reaction has the ability to convert lignin into a polyol that is soluble in most organic solvents and thus can be used in preparing polyurethane.\textsuperscript{107–110} Sadeghifar \textit{et al.}\textsuperscript{111} oxypropylated kraft lignin through a reaction with propylene oxide (Scheme 2-10). They reported that using propylene oxide at milder conditions gave way to selectively masking the phenolic hydroxyls, while more extreme conditions would lead to simultaneous ring opening homopolymerization of the propylene oxide while masking both the aromatic and aliphatic hydroxyls. They further determined that 99\% of the phenolic hydroxyl groups on kraft lignin were successfully oxypropylated when excess amounts of propylene oxide (2.5 eq.) were used under the mild conditions.

\begin{center}
\textbf{Scheme 2-10} Oxypropylation of the phenolic hydroxyls on lignin with propylene oxide.\textsuperscript{111}
\end{center}

As discussed, modification of the hydroxyls on lignin poses excellent potential for valorizing lignin into materials used in various industries and substituting the fossil-based materials used in such industries with this abundant, renewable, bio-based resource.
2.2.3 Lignin-based Materials

Several reports are available today with the potential of converting lignin into a compound with the ability to be incorporated into materials in substantial percentages and/or offering unique properties. With the variety of functional groups present on lignin, comes the possibility of using those groups with the aim of adding certain properties to lignin. Depending on the end goal, such modifications can be tailored to give desired behavior to the natural polymer in target systems.

Scheme 2-11 Synthesis of lignin-Br ATRP macroinitiator and rosin polymer-grafted composites (lignin-g-(Rosin Polymer)): LGEMA (X = 2, Y = CH$_3$) using MAEDA (m = 2, Z = CH$_3$), LGEA (X = 2, Y = H) using AEDA (m = 2, Z = H), and LGBA (X = 4, Y =
H) using ABDA (m = 4, Z = H) as reported by Wang et al.\textsuperscript{112} Copyright 2011, reproduced with permission from John Wiley and Sons.

Wang et al.\textsuperscript{112} examined the possibility of preparing highly hydrophobic composites using lignin and gum rosin as two renewable materials via ATRP polymerization. They esterified lignin with 2-bromoiso-buteryl bromide (BiBB) using triethylamine as the catalyst to obtain lignin-Br. Afterward, they mixed tris[2-(dimethylamino)ethyl]amine (Me\textsubscript{6}-TREN), 2-methacryloyloxyethyl dehydroabieticcarboxylate (MAEDA), and lignin-Br (as the ATRP macroinitiator) with Cu(I)Br at 65 °C for 24 h to obtain the MAEDA polymer grafted lignin (LGEMA). They also prepared 2-acryloyloxyethyl dehydroabieticcarboxylate (AEDA) polymer grafted lignin (LGEA) and 4-acryloyloxybutyl dehydroabieticcarboxylate (ABDA) polymer grafted lignin (LGBA) samples using the same process, substituting the MAEDA with AEDA and ABDA, respectively (Scheme 2-11).

Separately,\textsuperscript{112} they synthesized dehydroabietic acid (DHA)-grafted lignin (Lignin-g-DHA), also referred to as rosin acid-grafted lignin, by reacting oxalyl chloride with DHA to obtain OA-Cl, which was then added to lignin using catalytic amounts of triethylamine (Scheme 2-12).
Scheme 2-12 Synthesis of rosin acid-grafted lignin (lignin-g-DHA) as reported by Wang et al.\textsuperscript{112}

They reported that the length of the alkyl side groups in the acrylic rosin polymer-grafted lignin composites had a significant effect on the $T_g$ values of the composites, as they reported $T_g$ values of 20, 70, and 95 °C for the LGBA, LGEA, and LGMA composites, due to the presence of butyl, ethyl, and methacrylic side chains, respectively. Furthermore, they examined the hydrophobicity of the unmodified lignin, lignin-Br, lignin-g-DHA, LGEMA, LGEA, and LGBA samples by spin-casting thin films of them on a glass substrate and measuring the water contact angles of the prepared films. Subsequently, they determined that while the water contact angles for lignin and lignin-Br spin-cast films were close (75° and 78°, respectively), the same values increased significantly in the spin-cast films of lignin-g-DHA (92°), LGEMA (91°), LGEA (89°), and LGBA (91°) because of incorporating hydrophobic rosin moieties into composites of the naturally hydrophobic
lignin. The reported values were close to that of hydrocarbon-based polystyrene (90°), which was taken as evidence towards the suitability of the prepared composites to act as bio-based substitutes for packaging materials made from hydrocarbon-based polystyrene.

Ismail et al.\textsuperscript{113} modified and incorporated sodium lignosulfonate (LS) in two series of bio-based epoxy resins (Scheme 2-13). The first resin (Ser1LSGLYPa) was prepared through esterification of equal amounts of LS and ethylene glycol (EG), and LS and glycerol (GLY) with excess succinic acid anhydride (SAH) using dimethylbenzylamine (DMBA) as the catalyst to synthesize corresponding ester carboxylic acid derivatives of LSEGPA and LSGLYPa, respectively, which were then used to crosslink glycerol diglycidyl ether (GLYDGE). The LSGLYPa percentages reported were at 0, 20, 40, 60, and 80%. The second resin (Ser2GLYDGE) was prepared by esterifying equal amounts of LS and GLY with excess SAH using DMBA as the catalyst to synthesize LSGYPa and subsequent crosslinking with a mixture of GLYDGE and ethylene glycol diglycidyl ether (EGDGE), where the GLYDGE/EGDGE mixture was varied at 0, 20, 40, 60, 80, and 100% (Scheme 2-13). They reported that the $T_g$ values for both epoxy resins were dependent on the GLY content in them, as the values increased with increasing the GLY content in the epoxy resins. They also determined that increasing the amount of LSGLYPa in Ser1LSGLYPa and the amount of GLYDGE in Ser2GLYDGE did not affect the thermal decomposition temperature ($T_d$) and the mass residue at 500 °C (MR$_{500}$) of the prepared epoxy resins, as they both showed $T_d$ values around 275 °C and MR$_{500}$ values of around 22%. Such epoxy resins, prepared using chemically modified lignin, can have applications as crosslinking agents for PF resins.
**Scheme 2-13** Preparation scheme for Ser1LSGLYPA and Ser2GLYDGE epoxy resins as reported by Ismail *et al.*\(^{113}\) Copyright 2009, reproduced with permission from John Wiley and Sons.

Feng *et al.*\(^{114}\) prepared acetic acid functionalized lignin-based hydrogels (LGA1, LGA2, and LGA3) by reacting acetic acid lignin (AAL) with acrylamide (AAm) and ethyleneglycol dimethacrylate (EGDMA) as the crosslinker, using hydrogen peroxide as the initiator (Scheme 2-14), alongside control hydrogels of AAm and EDGMA with 2,2-azobisisobutyronitrile (AIBN) as the initiator (PAAm). The AAL:AAm weight ratios used in the preparation of the AAL-incorporated hydrogels were 0.5:2.5, 0.5:3.5, and 0.5:4.5 for the LGA1, LGA2, and LGA3 hydrogels, respectively.
They reported that the incorporation of AAL into the hydrogels had minimal effect on their decomposition temperatures, as the 5% weight loss temperature remained at around 250 °C for the AAL, LGA2, and PAAm samples. However, the LGA2 hydrogels showed higher char formation at 700 °C (24 wt%) than the PAAm hydrogels (21 wt%) as a result of the incorporation of lignin as a char formation-promoting material. Moreover, they determined that decreasing the amount of incorporated AAL in the hydrogels increased their swelling degrees noticeably while increasing the amount of the hydrophobic AAL would cause slower diffusion of water in the hydrogels. Consequently, they reported that the LGA3 hydrogel reached the swelling ratio of 930% within 420 min, while the same value was approximately 700% and 820% for the LGA1 and LGA2 hydrogels, respectively within the same time frame. Finally, they determined that increasing the lignin content in the LGA hydrogels improved the dye-adsorption in them, as the LGA1 hydrogel was able to adsorb 29.65 mg of methylene blue per every gram of hydrogel. Such hydrogels have the potential to be utilized as a flocculent in waste-water treatment, an unconventional potential application of lignin considering its inherent hydrophobicity.\textsuperscript{114}

Jawerth\textit{ et al.}\textsuperscript{115} reported the first thiol-ene click reaction on fractionated kraft lignin, by allylating the ethanol-soluble fraction of the kraft lignin with allyl chloride in the presence of sodium hydroxide to selectively allylate the phenolic hydroxyl groups. They
subsequently synthesized lignin-thiol resins by adding the allylated fractionated kraft lignin to trimethylpropane tris(3-mercaptopropionate) at different thiol/allyl ether molar ratios in silicone molds, followed by heating at 120 °C for 24 h (Scheme 2-15). They reported unsuccessful attempts at UV-curing the lignin-thiol resins due to the strong UV absorbance of the allylated lignin sample and utilizing non-depolymerized kraft lignin in the same reaction was not discussed in the report.

Scheme 2-15 Synthesis of lignin-thiol resins from ethanol-soluble fractionated kraft lignin as reported by Jawerth et al.\textsuperscript{115}

2.3 Summary and Knowledge Gaps

Despite the aforementioned valorization techniques available for lignin today, it is evident by the large number of studies in this field and the sheer volume of yet underutilized lignin that is being disposed of as waste every day, that the knowledge gap in the area of lignin valorization still exists, which amplifies the need for novel, optimized, and target-specific approaches to the task of utilizing lignin or modified lignin, a relatively untapped, renewable, sustainable, and abundant natural polymer for high-value bio-based materials.

As the most promising studies in this research area either employ multi-step reactions with harsh reaction conditions, seek to depolymerize or fractionate lignin prior to utilization
dispose of a portion of the material, or have only been successful at utilizing low percentages of lignin (chemically modified or not) in the reported systems, finding simple, straightforward approaches with more gentle reaction conditions, capable of incorporating appreciable percentages of lignin in its entirety in the target systems can have a detrimental effect on enhancing the scientific discoveries in this field and play a crucial role in developing a sustainable future for many classes of materials that are currently developed from non-renewable fossil fuel-based sources.

2.4 References


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Chapter 3

3 UV-Curable Coatings of Modified Lignin

3.1 Introduction

“Lignocellulose” is the biomass derived from dry plant matter,\(^1\) which consists of three major components: cellulose, hemicellulose, and lignin. The importance of lignocellulose as a source of energy and commodity chemicals is hallmarked by the availability of this renewable, carbon-based resource to produce fuels and chemicals that currently, are obtained from petroleum-based sources.\(^2\) Among the main constituents of lignocellulose, lignin is considered the least utilized component. As a complex cross-linked, racemic\(^3\), amorphous heteropolymer that is primarily generated as a by-product of the pulp and paper industry, lignin is utilized on-site as a fuel for recovery boilers in the form of black liquor. However, given the large production volume of 50-55 million tons (and growing) per year, a great amount of the material is under-utilized, with only about 2% of the lignin by-product being utilized for onwards product development.\(^4,5\)

This natural polymer is can be best described as several phenylpropanoid (C9) units that are linked together via different linkages (mainly ether and carbon-carbon),\(^6\) leading to a macromolecular structure that is rich in functional group chemistry (e.g. -OH, -C=O, and -OMe),\(^7\) which can undergo a variety of chemical reactions aimed at converting this industrial by-product into a value-added one. Several processes have been reported in the past with the sole purpose of creating higher value products from this untapped resource such as fragmentation/depolymerization by thermal conversion,\(^8\)–\(^14\) with the purpose of breaking down the chemical structure of lignin to obtain commodity reagents (e.g. phenolic monomers). On the other hand, processes such as alkylation/de-alkylation,\(^15\)–\(^17\) hydroxyalkylation,\(^18\)–\(^20\) amination,\(^21,22\) and nitration\(^23\) focus on chemically modifying lignin in order to increase its reactivity towards post-modification through the introduction/addition of functional groups. Nevertheless, considering the abundance and diversity of hydroxyl groups within lignin’s structure\(^24\), chemical modification of the bulk material through conversion of the hydroxyl groups is considered a highly viable, if not the best, approach to converting lignin into a value-added material. Such modified lignin
will have the ability (depending on the type of modification) to undergo various post-modification processes specifically targeting value-addition. Chemical methods such as esterification,\textsuperscript{25–31} phenolation,\textsuperscript{32–36} etherification,\textsuperscript{37–40} and urethanization\textsuperscript{41} are among the commonly used processes reported for modification of lignin via the hydroxyl functional groups (Scheme 3-1).

\textbf{Scheme 3-1} Examples of compounds and materials produced by different lignin modification processes such as amination,\textsuperscript{21} fragmentation/depolymerization,\textsuperscript{42} and esterification.\textsuperscript{26}

Lignin has a significant Ultraviolet- (UV)-absorbing property and therefore, has been incorporated into formulations to impart this property into various systems,\textsuperscript{43,44} and therefore, the focus on lignin based coatings has centered on thermally-crosslinked thiol-ene networks\textsuperscript{45,46}, coatings\textsuperscript{47–49}, or polyurethane thermosets.\textsuperscript{50–53} Nevertheless, utilizing un-purified commercial lignin as a monomer into UV/eb-crosslinkable systems and determining the properties of the coatings remain rare and require the use of photosensitive moieties as grafted co-monomers, multi-step syntheses, or other additives to overcome the inherently UV-absorbing nature of lignin.\textsuperscript{54,55} Curing coatings by UV light is widely accepted as the most efficient method for the rapid generation of highly-crosslinked, polymer networks from liquid formulations at ambient temperature.\textsuperscript{56} In
comparison to other types of photopolymerization and thermally initiated polymerization, UV-curing has several significant advantages that include lower equipment costs, much lower energy consumption, rapid curing times, and a imparts a substantial improvement on workplace safety as light cured coatings are far less hazardous to manufacture.\textsuperscript{56,57}

In this context, kraft lignin (KL) was methacrylated using methacrylic anhydride to obtain methacrylated lignin (ML). The obtained ML was then incorporated into UV-curable formulations (up to 31 wt.% ML). The resulting formulations were cured on glass substrates as 15 µm thick coatings. Extent and efficiency of chemical modification of lignin were analyzed using multinuclear magnetic resonance (NMR) and Fourier transform-infrared (FT-IR) spectroscopies. The UV-cured coatings were also characterized and analyzed using FT-IR spectroscopy (Attenuated Total Reflectance; ATR), Thermogravimetric analysis (TGA), and Drop shape analysis (DSA); Pull-off strengths and cohesive/adhesive failures of the prepared coatings were obtained following ASTM D4541-09 standard test.

3.2 Experimental Section

3.2.1 Materials

Kraft lignin (KL) was provided by FPInnovations-Thunder Bay Bio-Economy Technology Centre. KL samples were crushed into powder using a mortar and pestle and heated at 45°C for 48 hours \textit{in vacuo} (762 mmHg; C9=180 g). Methacrylic anhydride (MethA; 94%, stabilized with 0.2% 2,4-dimethyl-6-tert-butylphenol), 1-methylimidazole (1MIM; 99%), and triphenylphosphine (Ph$_3$P; 99%) were purchased from Alfa Aesar and used as received. 2,2'-Dimethoxy-2-phenylacetophenone (DMPA; 99%), tetra(ethylene glycol)diacrylate (TEGDA; technical grade, contains 150-200 ppm monomethyl ether hydroquinone (MEHQ) as inhibitor; 100-150 ppm hydroquinone as inhibitor), methacrylic acid (MA; 99%, contains 250 ppm MEHQ as inhibitor), 2,2’-biphenol (99%), 4-methoxyphenol (99%), 4-(benzylxy)phenol (98%), benzyl alcohol (anhydrous, 99.8%), 3-phenyl-1-propanol (98%), 1,2-phenylenedicarboxylic acid (99%), biphenyl-4,4’-dicarboxylic (97%), triethylamine (NEt$_3$; ≥ 99.5%, and chlorodi(isopropyl)phosphine ([CH(CH$_3$)$_2$]$_2$PCl, 96%) were obtained from Sigma-
Aldrich Co. and used as received. 6,6’-methylenebis(2-methoxy-4-methylphenol) was prepared using a literature procedure.\(^5\) Tetrahydrofuran (THF) was purchased from Caledon and dried using an MBraun Solvent Purification System (SPS) for the methacrylation process. The same solvent was stored in the drybox over 4Å molecular sieves for the reaction with chlorodi(isopropyl)phosphine. Dichloromethane (CH\(_2\)Cl\(_2\)) and hexanes were purchased from Caledon Laboratories and used without further purification. Dimethylsulfoxide-\(d_6\) (DMSO-\(d_6\)) was purchased from Cambridge Isotope Laboratories and used without any further purification. 2-[bis(4-hydroxyphenyl)methyl]benzyl alcohol was purchased from Tokyo Chemical Industry Co., Ltd. (TCI America) and used as received.

### 3.2.2 Methacrylation of KL

KL (1.00 g, 5.50 mmol) was added to THF (10 mL) in a round-bottom flask under vigorous stirring. The mixture was stirred for 5 min to obtain a homogeneous dark-brown solution. A mixture of 1-methylimidazole (1MIM, 0.10 mL, 0.10 mg, 1.25 mmol) and methacrylic anhydride (MethA, 0.82 mL, 0.84 g, 5.50 mmol) were then added to the KL solution and stirred at 60 °C for 60 minutes under an N\(_2\) atmosphere. The brown mixture was then added dropwise into hexanes (100 mL) while stirring to precipitate a light brown powder. The obtained powder was dissolved in CH\(_2\)Cl\(_2\) (10 mL) and washed with deionized water (30 mL) to remove the catalyst and any unreacted reagents. The organic phase was then precipitated in hexanes (100 mL) and dried in vacuo to give 1.46 g (146 wt% recovered product yield) ML. \(^1\)H NMR (DMSO-\(d_6\), 600 MHz); \(\delta_H\) 1.89 (br, 3H, CH\(_3\)), 5.82 (br, 1H, vinyl), 6.14 (br, 1H, vinyl).

### 3.2.3 Product Characterization

Proton NMR spectra were recorded using a Varian INOVA 600 MHz spectrometer on solutions of ca. 5 mg of sample dissolved in 1 g of DMSO-\(d_6\). \(^31\)P\{\(^1\)H\} NMR spectra were recorded using a Varian INOVA 400 MHz spectrometer on solutions of 50 mg sample, dissolved in 1 g of THF (256 scans, relaxation delay = 5s). All \(^31\)P\{\(^1\)H\} NMR spectra were referenced using 85% phosphoric acid as an external standard (\(\delta_P = 0.0\)). FT-IR spectra of samples were performed on a PerkinElmer FT-IR Spectrometer using
the universal attenuated total reflectance mode (UATR), a diamond crystal, as well as the UATR sampling accessory (part number: L1050231). Thermogravimetric analyses (TGA) were carried out on samples weighing 5-6 mg, using the Ramp mode at temperatures increasing from 25 ºC to 600 ºC at the rate of 10 ºC/min on a Q600 SDT TA Instrument and analyzed using TA Universal Analysis. Water contact angles (WCA) were measured placing the coatings on the stage of a Kruss DSA100 Drop Shape Analyzer (KRUSS GmbH, Hamburg, Germany). A static drop shape analyzer was used to determine the water contact angle on the surface.

A modified version of the method reported by Argyropoulos\textsuperscript{24,59} was employed for quantitative \textsuperscript{31}P\{\textsuperscript{1}H\} NMR analysis of KL compared with ML to quantify the OH groups on the KL sample, quantitatively determined the extent to which the hydroxyl groups were successfully converted during the methacrylation process. KL (0.05 g, 0.28 mmol) was added to THF (2 mL) in a round-bottom flask under vigorous stirring in the dry-box. The mixture was stirred for 5 min. to obtain a dark brown solution. NEt\textsubscript{3} (0.19 g, 261 µl, 1.88 mmol) was added to the mixture under stirring and stirred for 1 min. and then chlorodiisopropylphosphine ([CH(CH\textsubscript{3})\textsubscript{2}]\textsubscript{2}PCl; 0.085 g, 90 µl, 0.56 mmol) was added and the reaction mixture was stirred for 10 min at ambient temperature. The reaction mixture was then precipitated in n-pentane (10 mL) and dried \textit{in vacuo} to obtain a light brown powder. The dried powder was swelled in THF (1.7 mL) and inserted into an NMR tube. 50 µL of a 0.05 molar solution of Ph\textsubscript{3}P (2.5×10\textsuperscript{-6} mol) in THF was transferred into a capillary tube, flame-sealed, and inserted into the NMR tube containing the product and \textsuperscript{31}P\{\textsuperscript{1}H\} NMR spectra were collected. The same procedure was carried out on the obtained ML sample and the extent of methacrylation was determined by comparing the integrations of the observed peaks in relative to the external standard. Different regions of the obtained \textsuperscript{31}P\{\textsuperscript{1}H\} NMR spectrum were identified using small molecule, model reactions of corresponding hydroxyl-containing substrates with [CH(CH\textsubscript{3})\textsubscript{2}]\textsubscript{2}PCl.

3.2.4 Small Molecule Reactions

A general procedure was employed for model the reactions of [CH(CH\textsubscript{3})\textsubscript{2}]\textsubscript{2}PCl with hydroxyl-containing small molecules. Briefly, one molar equivalent of each small molecule was dissolved in THF (5 mL) in the dry-box. Excess NEt\textsubscript{3} was added to the
stirred solution, followed by addition of 1 molar equivalent \([\text{CH(\text{CH}_3)_2}]_2\text{PCl}\). A sample of the reaction mixture was transferred to an NMR tube, and \(^{31}\text{P}\{^1\text{H}\}\) NMR spectra were obtained.

2,2’-biphenol, 4-methoxyphenol, and 4-(benzyloxy)phenol were used as model small molecules containing uncondensed phenolic hydroxyl groups. \(^{31}\text{P}\{^1\text{H}\}\) NMR (400 MHz): \(\delta_P = 148.3, [\text{C}_6\text{H}_4\text{OP(C}_3\text{H}_7)_2]_2; 149.4, \text{CH}_3\text{O(C}_6\text{H}_4)\text{OP(C}_3\text{H}_7)_2; 149.5, (\text{C}_6\text{H}_5)\text{CH}_2\text{O(C}_6\text{H}_4)\text{OP(C}_3\text{H}_7)_2\).

Benzyl alcohol and 3-phenyl-1-propanol were used to represent aliphatic hydroxyl groups. \(^{31}\text{P}\{^1\text{H}\}\) NMR (400 MHz): \(\delta_P = 154.6, (\text{C}_6\text{H}_5)\text{CH}_2\text{OP(C}_3\text{H}_7)_2; 151.5, (\text{C}_6\text{H}_5)\text{C}_3\text{H}_6\text{OP(C}_3\text{H}_7)_2\).

2-[bis(4-hydroxyphenyl)methyl]benzyl alcohol was used as a small molecule comprising both aliphatic and uncondensed phenolic hydroxyl groups on its structure. \(^{31}\text{P}\{^1\text{H}\}\) NMR (400 MHz): \(\delta_P = 154.9, [(\text{C}_6\text{H}_4)\text{OP(C}_3\text{H}_7)_2]_2\text{CH}[((\text{C}_6\text{H}_4)\text{CH}_2\text{OP(C}_3\text{H}_7)_2]; 148.1, [(\text{C}_6\text{H}_4)\text{OP(C}_3\text{H}_7)_2]_2\text{CH}[((\text{C}_6\text{H}_4)\text{CH}_2\text{OP(C}_3\text{H}_7)_2].

To assign where the chemical shifts for the product of the reaction of COOH groups with \([\text{CH(\text{CH}_3)_2}]_2\text{PCl}\) would be observed, biphenyl-4,4’-dicarboxylic acid was used. \(^{31}\text{P}\{^1\text{H}\}\) NMR (400 MHz): \(\delta_P = 144.4, [(\text{C}_3\text{H}_7)_2\text{POC(O)C}_6\text{H}_4)_2].

Finally, 6,6’-methylenebis(2-methoxy-4-methylphenol) was reacted with \([\text{CH(\text{CH}_3)_2}]_2\text{PCl}\) to assign the peak region of the product of condensed aromatic hydroxyl groups. \(^{31}\text{P}\{^1\text{H}\}\) NMR (400 MHz): \(\delta_P = 162.1, [\text{CH}_5(\text{C}_6\text{H}_5)(\text{OCH}_3)(\text{OP(C}_3\text{H}_7)_2])_2].

3.2.5 UV-Cured Coatings of Methacrylate Lignin (ML) and Characterization

Mixtures, containing different weight percentages of ML (0, 10, 20, and 31 wt.%), equal amounts of photoinitiator (2,2-Dimethoxy-2-phenylacetophenone; DMPA; 5 wt.%), reactive diluent (methacrylic acid; MA 38 wt.%, incorporated into the coating after curing) and corresponding amounts of crosslinker (tetra(ethyleneglycol)diacrylate; TEGDA) were weighed and combined in screw top vials and sonicated at room temperature for 20 min to obtain particulate- and haze-free formulations, which were then
cast on clean glass slides (or stainless steel sheets in case of the coating-to-surface adhesion tests) using a 12-inch meyer rod (AP-JR10, Jr. rod, 12 inch OA, 3/8 inch DIA, 8-1/2 inch EP, #10 wire). Photopolymerization was then performed using a modified UV-curing conveyor system, purchased from UV Process and Supply Inc. with a mercury bulb (UVA: 154 mW/cm², 189 mJ/cm², UVB: 74 mW/cm², 90 mJ/cm²; as determined by a PP2-H-U Power Puck II purchased from EIT Instrument Markets) by placing the formulation cast surfaces on a conveyor (5.8 cm/s speed) for 25 s and irradiating with UV light to give coatings of 15 µm thickness as measured by a micrometer. Coating-to-surface adhesion for pull-off strengths and cohesive/adhesive failures of the prepared coatings were performed were obtained following ASTM D4541–09ε standard test, using a PostiTest pull-off adhesion tester (type V, self-aligning), purchased from Defelsko Corporation USA on coatings of 15 µm thickness, applied and cured on 20×15 cm stainless steel sheets. The procedures for determining the coating properties were all carried out in triplicates to ensure reproducibility and the results are reported as mean ± standard deviation.

3.3 Results and Discussion

3.3.1 Methacrylation of KL

Methacrylic anhydride (MethA) and 1-methylimidazole (1MIM) were used as the methacrylating reagent and the catalyst, respectively (Figure 3-1a). Samples of the reaction mixture were evaluated by ¹H NMR spectroscopy and showed three new signals for the ML sample as compared to the KL sample. The broad signal at δ_H = 1.89 was assigned to the methyl functionality and the signals at δ_H = 5.82 and 6.14 were attributed to the olefinic protons of the methacrylate group and gave strong evidence for grafted methacrylate groups onto lignin (Figure 3-1b).

FT-IR spectroscopy was performed on KL and the prepared ML samples (Figure 3-1c). Spectra of ML repeatedly showed significantly decreased signals at 3200-3500 cm⁻¹ (s, br; O-H) relative to KL along with a corresponding increase in in the signal at 1735-1750 cm⁻¹ (C=O), providing corroborating evidence for the methacrylation of lignin and the transformation of hydroxyl to acrylate. After methacrylation, IR signals increased in the
ML product at 1000-1320 cm$^{-1}$ and 650-1000 cm$^{-1}$ corresponding to stretching vibrations of C–O (ester) and bending vibrations of the unsaturated C=C groups, respectively.

Taken together, these spectroscopic signatures provide strong evidence for the successful grafting of lignin.
Figure 3-1 Reaction scheme for methacrylation of lignin using methacrylic anhydride (MethA) and 1-methylimidazole (1MIM) (a), along with $^1$H NMR (b) and FTIR (c) spectra of KL (A) and ML (B).
3.3.2 Quantification of the Hydroxyl Functionalities of KL

Other studies have utilized $^{31}$P{$^1$H} NMR spectroscopy for quantifying the hydroxyl-based functional groups in lignin. Such work used 4,4,5,5-tetramethyl-1,2,3-dioxophospholane, in the presence of chromium(III) acetylacetonate (relaxation reagent) to probe a native lignin sample and the modified lignin.\textsuperscript{24,59,60} In this current work, chlorodi(isopropyl)phosphine ([CH(CH$_3$)$_2$]$_2$PCl) was used to install a phosphine functional group that aided in the identification and quantification of the hydroxyl environments within the structure of lignin. This is a good mimic for the phospholene employed by Argyropoulos, given the comparable steric and electronic demand. After dehydrohalogen coupling, the various locations of P throughout lignin gave separate signals for the different O-P functional group environments (aliphatic, $\delta_P = 160$-170; uncondensed aromatic, $\delta_P = 147$-150; condensed aromatic, $\delta_P = 150$-160; and carboxyl, $\delta_P = 136$-147). After the lignin has undergone methacrylation (i.e. to form ML), the chlorophosphine was again utilized as a probe for residual OH groups that remained unfunctionalized. To correlate $^{31}$P{$^1$H} signals arising from the dehydrocoupling reaction promoting the P-O linkage to lignin, benchmark reactions with small molecule models were completed. The stoichiometry of OH groups within lignin was correlated by referencing the signal integrations with a known internal standard, Ph$_3$P. In a typical reaction of either small molecule or lignin itself (Scheme 3-2), corresponding signals appear in the $^{31}$P{$^1$H} spectra that resulted from the dehydrohalogen coupling.
Scheme 3-2 Reaction scheme of kraft lignin and small molecules representing different hydroxyl functionalities with chlorodi(isopropyl)phosphine along with the corresponding chemical shifts of the products. Chemical shifts of the structural motifs were in good agreement with related molecular architectures.61,62,63

Spectroscopic signals ranging from $\delta_P = 136$-170 were observed and the obtained signals were used to identify four regions corresponding to different types of O-P environments and by extension we identified the -OH functionalities within the KL sample (Figure 3-2). The signal range was then integrated in comparison with the external standard (Ph$_3$P, $2.5 \times 10^6$ mols, $\delta_P = -4$) to determine the amount of -OP functionalities within 0.05 g of the sample.
Thus, it was determined that each C9 unit within the KL sample, contained an average of
1.5 ± 0.2 mol hydroxyl functionalities. This process was carried out in triplicate to
confirm its reproducibility. The breakdown of each of the regions gave the corresponding
amounts of each type of hydroxyl functionality. Once the corresponding regions were
identified, this information was used along with the Ph₃P standard to identify and
quantify the number and type of O-P functionalities within KL and ML samples per
gram of sample (Figure 3-3). The KL sample contained 0.86 mmol·g⁻¹ aliphatic hydroxyl
groups, 0.70 mmol·g⁻¹ uncondensed phenolic hydroxyl groups, 1.76 mmol·g⁻¹ condensed
phenolic groups, and 0.85 mmol·g⁻¹ carboxylic groups, on average. Furthermore, we
determined that on average, every C9 unit on lignin contains 1.5 ± 0.2 molar equivalents
of hydroxyl functionalities. The $^{31}$P{$^1$H} NMR spectroscopic analyses were performed in
triplicate.
Figure 3-2 $^{31}\text{P}\left\{^1\text{H}\right\}$ NMR spectra of products (Scheme 3-2) obtained from reaction of chlorodi(isopropyl)phosphine with kraft lignin (A), biphenyl-4,4'-dicarboxylic acid (B; $\delta_p = 144.4$), 6,6'-methylenebis(2-methoxy-4-methylphenol) (C; $\delta_p = 162.1$), 2-[bis(4-hydroxyphenyl)methyl]benzyl alcohol (D; $\delta_p = 154.9$, uncondensed aromatic, $\delta_p = 148.1$, aliphatic), 4-(benzyl oxy)phenol (E; $\delta_p = 149.5$), 2,2'-biphenol (F; $\delta_p = 148.3$), 4-methoxyphenol (G; $\delta_p = 149.4$), benzyl alcohol (H; $\delta_p = 154.6$), and 3-phenyl-1-propanol (I; $\delta_p = 151.5$).

### 3.3.3 Extent of Methacrylation

To determine the EOM, ML was reacted with [CH(CH$_3$)$_2$]$_2$PCl using the same reaction conditions and parameters as previously described for reacting the KL sample with [CH(CH$_3$)$_2$]$_2$PCl. Using the same amount of Ph$_3$P as the external standard while obtaining the $^{31}\text{P}\{^1\text{H}\}$ spectrum of the product of reacting ML with [CH(CH$_3$)$_2$]$_2$PCl (Figure 3-3), we were able to quantify residual hydroxyl functionalities on the ML sample that were not converted in the methacrylation process. Using this approach, the EOM was found to be $70 \pm 1.7\%$, where all carboxylic and uncondensed aromatic hydroxyls, as well as
72.27% and 44.48% of the aliphatic and the condensed aromatic hydroxyls were respectively determined to be completely converted through this process.

**Figure 3-3** $^{31}P$\{^1H\} NMR spectra of products obtained from the reaction of KL (A) and ML (B) with [CH(CH$_3$)$_2$]$_2$PCl.

### 3.3.4 Preparation and Characterization of UV-Cured Coatings Using ML

While the KL sample had a naturally dark brown color and limited solubility in common organic solvents, the ML sample had a much better solubility profile, making UV-curing more facile. ML-based UV-cured coatings were prepared using formulations consisting of ML, photoinitiator (DMPA; 5 wt.%), reactive diluent (MethAcid, 38 wt.%), and corresponding amounts of crosslinker (TEGDA) as described in Table 3-1. The cast formulations were applied onto the corresponding surfaces (glass or stainless steel) using a Meyer rod, and cured on a UV-curing conveyor (Scheme 3-3). Coatings containing more than 31 wt.% ML were not curable under UV-light, likely due to the decreased transparency of the formulations at higher ML loadings, preventing effective penetration of the coatings by UV light.
**Scheme 3-3** Process description for incorporating lignin into UV-cured coatings

(TEGDA = tetraethyleneglycoldiacrylate; MA = methacrylic acid; DMPA (photoinitiator) = 2,2-Dimethoxy-2-phenylacetophenone.

ATR-FTIR spectra were obtained before curing and for freshly prepared coatings after removing from the surface with a scalpel and grinding into powder form using a mortar and pestel. The peak corresponding to C=O (1720 cm\(^{-1}\)) was used as an internal standard as its intensity does not change upon curing. The C=C (1620-1680 cm\(^{-1}\)) peak was deconvoluted from the adjacent C=O peak in each spectra, the corresponding intensity was compared to the intensity of the internal standard peak before and after curing and the cure percentage was calculated as the percent decrease in the relative intensity of the C=C peak (Figure 3-4, 100% corresponding to zero C=C functionality remaining). Figure 3-5a shows the cure percentages corresponding to the wt.% of ML in each coating. It was determined that more of the liquid formulation was cured as the amount of ML was increased in the formulation (Table 3-1), which was attributed to the inherently high cross-link density provided by the ML.
Figure 3-4 ATR-FTIR spectra of the prepared liquid formulations and cured coatings, highlighting the reduction in the intensity of the C=C peak (corresponding to the cure percentage). Values shown on the deconvoluted peaks represent the peak intensities used for determining corresponding cure percentages.

Drop shape analysis was used to determine the effect of the ML-incorporation on the hydrophobicity of the prepared UV-cured coatings (Figure 3-5b). It was determined that the incorporation of ML into the prepared coatings increased their hydrophobicity, with the measured water contact angle of the ML₀ and the ML₃₁ being 43.1 ± 0.3 and 51.2 ± 0.6 degrees, respectively (Table 3-1). The obtained values are in-line with the hydrophobic nature of lignin or lignin derivatives²⁷,⁶⁴,⁶⁵ and comparable to the same values reported for thermally cured lignin-based coatings.⁶⁶

Pull-off adhesion tests were performed on the prepared coatings to determine the actual pull-off force (F) required to detach a steel dolly (14 mm diameter) that was affixed to coatings cured on stainless steel substrates (Figure 3-5c). Liquid formulations containing known amounts of ML (0, 10, 20, and 31 wt.%) were applied to clean steel plates (approximately 12 cm by 24 cm) and cured under UV-light. Coated surfaces were degreased using acetone to remove any oil, moisture, or dust. The surfaces were then
abraded using a 3M abrasive pad to promote adhesion between the surface and the dollies. A mixture of Loctite 907 Hysol adhesive was prepared according to the standard procedure and applied on the bases of 14mm dollies (approximately 50-100 microns thick); Dollies were then placed on the coated surfaces and pushed carefully to avoid any sliding. Any excess adhesive around the edges of the dolly was removed with cotton swabs. The dollies were then left for 3 days to let the adhesive cure completely. After 3 days, the dollies were connected to the tester using the provided actuator assembly. Following the manufacturer’s instructions, tests were carried out at MPa rates of 0.40, 0.70, 1.40, 2.00, and 2.50 to obtain the pressure required for pulling the attached dolly off the coated surface. Each coating was tested in triplicate and the results were reported as mean ± standard deviation. The required force for each coating was calculated according to the equation provided in the standard test procedure: 

\[ X = \frac{4F}{\pi d^2} \]

where \( X \) represents the greatest mean pull-off stress applied during a pass/fail test, or the pull-off strength achieved at failure (both have units of MPa); \( F \) represents the actual force applied to the test surface (N); and \( d \) stands for the equivalent diameter of the original surface area stressed (mm). It was determined that increasing the amount of ML in the coating formulation up to 31 wt.%, increased the force required for pulling the corresponding coating off the surface by 278 N (Table 3-1).
Figure 3-5 Determined cure percentages (a), water contact angles (b), actual pull-off forces (c), and adhesive/cohesive failures (d) of the ML-based UV-cured coatings.

Percentages of adhesive and cohesive failures were calculated by determining the respective areas on the dollies representing adhesive (area of the coating attached to the dolly after pull-off) and cohesive failures (area of the coating with visibly lost cohesiveness; Figure 3-5d). During the pull-off adhesion test, each tested surface was monitored to measure the cohesive and adhesive failures. Adhesive failure refers to the percentage of the coating which fails to stick on the surface, whereas the cohesive failure refers to the percentage of the coating that fails to keep its entirety and is divided into two layers (i.e. one sticking to the dolly and one to the surface). The area of the dolly was measured prior to the test (b). The area of coating attached to the dolly after the pull-off test was measured as well (a). Comparing these two recorded values gave the adhesive
failures as: \( A.F. = \frac{a}{b} \times 100 \). If the coatings were to detach from the surface in a non-cohesive manner (losing its entirety), the area of non-cohesive coating left on the dolly would be recorded \((c)\). Comparing this area with the total surface of the dolly \((b)\) would then give the cohesive failure as: \( C.F. = \frac{c}{b} \times 100 \). Each measurement was carried out in triplicate to ensure its reproducibility. As no cohesive failure was observed for any of the coatings, no mean and standard deviation was reported for the cohesive failures. While none of the coatings showed any signs of cohesive failure, the percentage of adhesive failure decreased as the amount of ML was increased in the coating formulation (Table 3-1).

**Table 3-1** Formulation details for ML-based UV-cured coatings.

<table>
<thead>
<tr>
<th>Composition or property</th>
<th>( \text{ML}_0 )</th>
<th>( \text{ML}_{10} )</th>
<th>( \text{ML}_{20} )</th>
<th>( \text{ML}_{31} )</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>ML</strong> (wt.%)</td>
<td>0</td>
<td>10</td>
<td>20</td>
<td>31</td>
</tr>
<tr>
<td>DMPA (wt.%)</td>
<td>5</td>
<td>5</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td>MA (wt.%)</td>
<td>38</td>
<td>38</td>
<td>38</td>
<td>38</td>
</tr>
<tr>
<td>TEGDA (wt.%)</td>
<td>57</td>
<td>47</td>
<td>37</td>
<td>27</td>
</tr>
<tr>
<td>Cure percentage (%)</td>
<td>43.4 ± 1.7</td>
<td>47.7 ± 2.3</td>
<td>66.2 ± 1.2</td>
<td>69.2 ± 1.1</td>
</tr>
<tr>
<td>Water contact angle (°)</td>
<td>43.1 ± 0.3</td>
<td>45.2 ± 0.6</td>
<td>48.2 ± 0.5</td>
<td>51.2 ± 0.6</td>
</tr>
<tr>
<td>Pull-off force (N)</td>
<td>407.7 ± 7.8</td>
<td>498.5 ± 8.7</td>
<td>607.2 ± 6.1</td>
<td>685.7 ± 5.1</td>
</tr>
<tr>
<td>Adhesive failure (%)</td>
<td>20.6 ± 3.1</td>
<td>20.6 ± 3.1</td>
<td>16.5 ± 2.6</td>
<td>7.0 ± 1.6</td>
</tr>
<tr>
<td>Cohesive failure (%)</td>
<td>0</td>
<td>0</td>
<td>0</td>
<td>0</td>
</tr>
<tr>
<td>( T_d ) onset (°C)</td>
<td>86.2 ± 1.7</td>
<td>140.5 ± 1.0</td>
<td>160.9 ± 0.8</td>
<td>165.3 ± 0.9</td>
</tr>
<tr>
<td>Wt.% at 600 °C</td>
<td>3.7 ± 1.3</td>
<td>13.3 ± 1.8</td>
<td>15.9 ± 1.5</td>
<td>21.9 ± 1.0</td>
</tr>
</tbody>
</table>
After casting and curing the formulations on glass slides, each coating was scraped off the glass surface using a scalpel and crushed using a mortar and pestle. Each crushed sample was heated at 40 °C in vacuo for 24 h before measuring its thermal properties. Thermogravimetric analyses were then performed on the crushed coating samples to examine the effect of ML-incorporation on the thermal stability of the coatings at high temperatures (Figure 5). It was observed that the methacrylation of lignin, along with the incorporation of ML into the prepared coatings, impacted the T_d onset temperatures of the coatings. The T_d onset values of the KL sample (99.2 ± 0.9 °C) increased by 60.6 °C upon methacrylation (159.8 ± 1.7 °C for ML). While increasing the amount of ML in the UV-cured coatings the T_d onset temperatures increased to 86.2 ± 1.7, 140.5 ± 1.0, 160.9 ± 0.8, and 165.3 ± 0.9°C for the ML0, ML10, ML20, and ML31 coatings, respectively. Similarly increasing ML amounts in the coatings increased the retained weight at 600 °C, i.e., increased char formation, resulting from the presence of lignin with inherently crosslinked structure that promotes char formation. 10,66

Figure 3-6 Thermogravimetric analysis of the KL(A) and ML(B) samples, along with the ML0 (C), ML10 (D), ML20 (E), and ML31 (F) coatings, representing the obtained
thermograms under N₂, as well as the onset decomposition temperatures (T₉₀ onset) and the retained weights (char) at 600 °C (wt.%) for each sample.

3.4 References

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Chapter 4

4 Methacrylation of Kraft Lignin for UV-Curable Coatings: Process Optimization Using Response Surface Methodology

4.1 Introduction

Lignin is a complex, racemic,\(^1\) and aromatic heteropolymer, composed of mainly three phenylpropanoid (C9) units: \(p\)-coumaryl alcohol, coniferyl alcohol, and synapyl alcohol containing zero, one, and two methoxy functionalities, respectively, linked mainly by ether bonds (\(\beta\)-O-4 or \(\alpha\)-O-4) and C-C (5-5) linkages.\(^2\,^3\) Lignin as the main component in lignocellulosic biomass is an abundant natural polymer second only to cellulose. It is currently produced in large quantities as technical lignin, such as kraft lignin (KL), byproduct from the pulping processes, which is however utilized mainly as low-value solid fuel for heat production and for regeneration of pulping chemicals in recovery boilers.\(^4\) As such, lignin is an under-utilized sustainable resource.

Lignin’s chemical structure, particularly the relative abundance of hydroxyl functionalities,\(^5\) provides the potential for the macromolecule to undergo various chemical modification processes for the production of high-value bio-based products.\(^6\,^8\) The chemical modification of hydroxyl functionalities has been the subject of many studies, aiming to convert lignin into a material that is suitable for post-modification and subsequent incorporation into various products that are currently produced from fossil-resources. Such chemical modifications have also proved to improve lignin’s limited solubility in common organic solvents and subsequently, increase the possibility of incorporating lignin into different supramolecular systems.\(^8\,^9\) Tan\(^11\) reported that phenolation of the aliphatic hydroxyl functionalities within lignin with excess amounts of cardanol, a natural alkyl phenol, for 6 h at 60 °C, resulting in the addition of phenolic hydroxyls in lignin and making it a more suitable polyol for the preparation of bio-based polyurethane coatings. Sadeghifar \textit{et al}.\(^12\) reported oxypropylation (i.e., etherification of phenolic hydroxyl functionalities with propylene oxide) of kraft lignin at 40 °C for 18 h, using excess amounts of propylene oxide. Wang \textit{et al}.\(^13\) reported that esterification of
lignin’s hydroxyl functionalities with 2-boromoisobutryl bromide (BiBB). They subsequently employed the synthesized lignin-Br material as an atom transfer radical polymerization (ATRP) macroinitiator with 2-methacryloxyethyl dehydroabieticcarboxylate (MAEDA), 2-acryloyloxyethyl dehydroabieticcarboxylate (AEDA), and 4-acryloxybutyl dehydroabieticcarboxylate (ABDA), respectively, to prepare corresponding MAEDA-, AEDA-, and ABDA-polymer grafted lignin materials with increased water contact angles as spin-cast films.\textsuperscript{13} In a separate study, Wang et al.\textsuperscript{14} reported conversion of lignin to lignin-Br, using excess amounts of BiBB for 25 h at 0 °C, which yielded ca. 65% lignin-Br material as an ATRP macroinitiator to synthesize lignin-graft-polystyrene (lignin-g-PS) and lignin-graft-poly(styrene-co-acryloyl benzophenone) (lignin-g-poly(styrene-co-ABP)) co-polymers. Thielemans and Wool\textsuperscript{15} reported that esterifying lignin with excess amounts of methacrylic anhydride for 18 h at 50 °C resulted in metacrylation of both the aliphatic and the phenolic hydroxyl functionalities within its chemical structure, yielding 55% of esterified lignin products as a suitable candidate for incorporation into unsaturated thermoset systems.

Although the reported studies on the chemical modification of lignin present very interesting chemistry, some major setbacks of the reported process of lignin modification are: (1) all were on a lab-scale; (2) the majority of the reported processes do not operate under optimized conditions; and (3) to the best of our knowledge, most chemical modification processes for lignin generally need excess amounts of reagents, as well as relatively long reaction times, yielding relatively low amounts of final products.\textsuperscript{15–17} While a shorter reaction time is always desirable for industrially-feasible processes, the use of excess reagent amounts can harm the process’s feasibility both in terms of its upfront costs, and the work-up costs of the process to recycle the unreacted reagent. Although lignin’s low cost and renewability make it an appealing starting material to produce high-value bioproducts, an expensive, complicated, and low-yield process would lead to expensive final products, regardless of the price of the starting material. To take full advantage of the sustainability that lignin offers, processes aiming at converting lignin into value-added products must be optimized, to achieve the highest possible efficiency for up-scale production.
Response surface methodology (RSM) comprises a set of mathematical and statistical techniques capable of obtaining an adequate functional relationship between a response and several associated input variables. Although unknown, this relationship can be approximately depicted depending on the complexity of the system. Among the available RSM designs, central composite design (CCD) is the most commonly employed. CCD starts with a first order design (fitting a first-degree model) and uses additional points to fit the second-degree model, which can be depicted by the following equation:\(^{18,19}\)

\[
Y = \beta_0 + \sum_{i=1}^{K} \beta_i X_i + \sum_{i=1}^{K} \beta_{ii} X_i^2 + \sum_{i=1}^{K} \sum_{j=i+1}^{K} \beta_{ij} X_i X_j + \varepsilon
\]

**Equation 4-1**

Where \(Y\) is the response in the study and \(X_i\) and \(X_j\) are independent variables, \(\varepsilon\) corresponds to the noise/error component in the response and \(\beta_0, \beta_i, \beta_{ii}, \) and \(\beta_{ij}\) represent the regression coefficients of the intercept, linear, quadratic and interaction parameters, respectively.\(^{19}\)

The present work aimed at transforming lignin - the under-utilized sustainable resource into a methacrylated lignin (ML) for UV-curable coatings. The process parameters (i.e. reaction time, reaction temperature, and the reagent/lignin molar ratio) for the methacrylation of kraft lignin (KL) were optimized using response surface methodology with a central composite design to maximize the mass yield of the synthesized methacrylated lignin (ML). The obtained ML material was then incorporated into a UV-curable formulation at 30 wt.% and cast as a 25 µm thick coating film on a glass slide subject to UV-curing. The obtained coating was then characterized with respect to its thermal stability, water contact angle, and crosslinking percentage.

### 4.2 Materials and Methods

#### 4.2.1 Materials

Kraft lignin (KL) was provided by FPInnovations, produced by FPInnovations-Thunder Bay Bio-Economy Technology Centre (\(M_w \sim 10,000\) g/mol, based on GPC-UV analysis;
0.57 wt.% ash; and 5.2 wt.% sulfur on a dry ash free basis). The KL samples were crushed into powder using a mortar and pestle and heated at 45ºC for 48 hours under vacuum (-762 mmHg; C9=180 g). Methacrylic anhydride (MethA; 94%, stabilized with ca. 0.2% 2,4-dimethyl-6-tert-butylphenol) and 1-methylimidazole (1MIM; 99%) were purchased from Alfa Aesar and used as received. Chlorodi(isopropyl)phospine ([CH(CH₃)₂]₂PCl, 96%) and 2-hydroxy-2-methylpropiophenone (HDMAP; 97%) were obtained from Sigma-Aldrich Co. and used as received. EBERCYL® 1360 (EB-1360, a hexafunctional silicone hexaacrylate resin from Allnex company) as a crosslinker for UV curing was purchased from Allnex Belgium SA. and used as received. Tetrahydrofuran (THF) was purchased from Caledon and dried using an MBraun Solvent Purification System (SPS) for the methacrylation process. The same solvent was stored in the drybox over 4 Å molecular sieves for the reaction with chlorodi(isopropyl) phospine. Dichloromethane and hexanes were purchased from Caledon Laboratories and used without further purification. Dimethylsulfoxide-d₆ (DMSO-d₆) was purchased from Cambridge Isotope Laboratories and was used without any further purification.

4.2.2 Characterization Methods

¹H NMR spectra were recorded using a Varian INOVA 600 MHz spectrometer (¹H, 599.5 MHz; ¹³C, 150.8 MHz; ³¹P, 242.6 MHz) and referenced to residual DMSO (2.5 ppm). ³¹P{¹H} NMR spectra were recorded using a Varian INOVA 400 MHz spectrometer (¹H, 400 MHz; ¹³C, 158 MHz; and ³¹P{¹H}, 162 MHz) and referenced using an external standard (85% H₃PO₄; δₚ = 0). FT-IR spectra of samples were performed on a PerkinElmer FT-IR Spectrometer using the universal attenuated total reflectance mode (UATR), a diamond crystal, as well as the UATR sampling accessory (part number: L1050231). Thermogravimetric analyses were carried out on samples weighing ca. 5-6 mg, using the Ramp mode at temperatures increasing from 25 ºC to 800 ºC at the rate of 10 ºC/min on a Q600 SDT TA Instrument and analyzed using TA Universal Analysis. Crosslinking percentages were calculated by comparing the signal intensity at 800 cm⁻¹, corresponding to the C=C bending vibrations in the FTIR spectra of the liquid UV-curable formulation (before curing) and the solid UV-cured coating samples. Water contact angles (WCA) were measured using a Kruss DSA100 Drop Shape Analyzer. UV-curing was performed using
a modified UV-curing conveyor system, purchased from UV Process and Supply Inc. with a mercury bulb (UVA: 154 mW/cm², 189 mJ/cm², UVB: 74 mW/cm², 90 mJ/cm²; as determined by a PP2-H-U Power Puck II purchased from EIT Instrument Markets).

4.2.3 Methacrylation of KL

The methacrylation processes were carried out in 10 mL round-bottom flasks by reacting KL with MethA in THF sovlen in the presence of 1MIM as catalyst at 40-80°C for 20-100 min. In a typical run (the center point of the CCD as shown in Table 1: 60 °C, 60 min and 1MIM/KL = 0.22), dried KL (200 mg, 1.1 mmol) was added to 2 mL THF in the round-bottom flask under vigorous stirring. The mixture was stirred for ca. 5 min to obtain a homogeneous dark-brown solution. A mixture of 1MIM (19 µL, 19 mg, 0.24 mmol) and MethA (165 µL, 171 mg, 1.1 mmol) was then added to the previous solution and stirred at 60 °C for 1 h under N₂ atmosphere. The brown mixture was then added dropwise to 20 mL hexanes under stirring to precipitate a light brown powder. The obtained powder was dissolved in DCM and washed with water to remove the catalyst and any unreacted reagent. The organic phase was then precipitated in 20 mL hexanes and dried under vacuum to give ML.¹H NMR (DMSO-d₆, 600 MHz): δH 1.89 (br, 3H, CH₃), 5.82 (br, 1H, vinyl), 6.14 (br, 1H, vinyl). The obtained powder was then weighed to determine the reaction yield. The recovered mass yield was calculated as follows:

\[
\text{Recovered Mass Yield (wt. %) } = \frac{m_{ML}}{m_{KL}} \times 100
\]

Equation 4-2

Where \(m_{KL}\) is the weight of KL as starting material and \(m_{ML}\) is the weight of the final product.

4.2.4 Experiment Design

A central composite design (CCD), incorporating 3 independent variables (i.e., reaction time (min), reaction temperature (°C), and KL/1MIM molar ratio) was used to optimize the KL methacrylation process. The design contained 8 cube points, 6 axial points, and 1 center point and consisted of 20 runs, including 6 replicate runs to examine the reproducibility of the center point conditions. Ranges for the three variables in this design
were chosen based on our preliminary experiments and the literature.\textsuperscript{15,20} The variables and their corresponding levels are detailed in Table 4-1. Response surface methodology was implemented, and the results were interpreted using MiniTab 17. The response surface design was analyzed to optimize the synthesis condition of ML and probe the relative or interactive effects of KL/1MIM molar ratio, reaction temperature, and reaction time on the amount of obtained product from the methacrylation reaction.

**Table 4-1** Independent variables and their levels in the central composite design.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Unit</th>
<th>Term</th>
<th>-1.682</th>
<th>-1</th>
<th>0</th>
<th>1</th>
<th>1.682</th>
</tr>
</thead>
<tbody>
<tr>
<td>Reaction temperature</td>
<td>°C</td>
<td>X₁</td>
<td>43</td>
<td>50</td>
<td>60</td>
<td>70</td>
<td>77</td>
</tr>
<tr>
<td>Reaction time</td>
<td>min</td>
<td>X₂</td>
<td>26</td>
<td>40</td>
<td>60</td>
<td>80</td>
<td>94</td>
</tr>
<tr>
<td>1MIM/KL</td>
<td>mol/mol</td>
<td>X₃</td>
<td>0.186</td>
<td>0.200</td>
<td>0.220</td>
<td>0.240</td>
<td>0.253</td>
</tr>
</tbody>
</table>

4.2.5 Preparation of ML-based UV-Cured Coatings

In 2 separate vials, two mixtures containing 0 and 30 wt.% ML, equal amounts of photoinitiator (5 wt.% HDMAP), corresponding amounts of the crosslinker (95 and 65 wt.% EB-1360), and 100 mg CH₂Cl₂ were mixed and sonicated for ca. 15 min to obtain particulate- and haze-free solutions. The obtained formulations were then cast on clean glass slides, using a 12-inch meyer rod (AP-JR30, 12-inch OA, 3/8-inch DIA, 8-1/2-inch EP, #30 wire) and placed on the UV-curing conveyer system and irradiated with UV-light to give coatings of ca. 76 µm thickness.

4.3 Results and Discussion

4.3.1 Modelling and Analysis of Variance

Detailed values of the performed reactions and their corresponding mass yields are reported in Table 4-2. The obtained results show that the amount of obtained product is dependent
on the amount of catalyst in use, reaction time, and reaction temperature. To find the best fitting model to the experimental responses, various functions such as linear, quadratic and cubic models were developed. A quadratic equation based on the coded values of the variables was determined to have the best fit to the experimental responses. Coefficient significances were evaluated according to a 95% confidence interval ($P$-value $> \alpha=0.05$). Any coefficient with $P$-value $< 0.05$ has a statistically significant effect on the responses and vice versa. The final model to predict the methacrylation recovered mass yield is as follows:

$$
Yield \ (wt. \%) = 141.16 - 14.34X_1 - 12.37X_2 + 3.62X_3 - 12.30X_1^2 - 8.76X_2^2 - 9.12X_3^2 - 4.81X_1X_2 - 8.56X_1X_3 + 12.06X_2X_3$$  \textbf{Equation 4-3}

Where $X_1$, $X_2$, and $X_3$ represent the coded values for reaction temperature, reaction time and 1MIM/KL molar ratio, respectively. A positive term is indicative of a synergistic effect on the response and a negative term indicates an antagonistic effect on the response. The complex model for the methacrylation mass yield shows that all the terms including linear, square and interaction factors are significant. The model also depicts that the linear factor of 1MIM/KL molar ratio ($X_3$) and the interaction term of reaction time and the 1MIM/KL molar ratio ($X_2X_3$) are the only two coefficients that have a positive effect on the response, while all other coefficients have negative effects on it. Thus, lower reaction times and reaction temperature, along with employment of more catalyst will lead to a higher mass yield for the methacrylation process of KL. However, the model predicts that decreasing the reaction temperature and reaction time will have a greater effect on the mass yield than increasing the catalyst amount, which can be worth considering in an industrial setting.
Table 4-2 The central composite matrix and output responses for methacrylation of KL.

<table>
<thead>
<tr>
<th>Std. Run #</th>
<th>X₁</th>
<th>X₂</th>
<th>X₃</th>
<th>Temp. (°C)</th>
<th>Time (min)</th>
<th>1MIM/KL (mol/mol)</th>
<th>Output responses</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>18</td>
<td>-1</td>
<td>-1</td>
<td>50</td>
<td>40</td>
<td>0.200</td>
<td>0.262 131</td>
</tr>
<tr>
<td>2</td>
<td>8</td>
<td>1</td>
<td>-1</td>
<td>70</td>
<td>40</td>
<td>0.200</td>
<td>0.26 130</td>
</tr>
<tr>
<td>3</td>
<td>9</td>
<td>-1</td>
<td>1</td>
<td>50</td>
<td>80</td>
<td>0.200</td>
<td>0.18 90</td>
</tr>
<tr>
<td>4</td>
<td>3</td>
<td>1</td>
<td>-1</td>
<td>70</td>
<td>80</td>
<td>0.200</td>
<td>0.143 71.5</td>
</tr>
<tr>
<td>5</td>
<td>6</td>
<td>-1</td>
<td>-1</td>
<td>50</td>
<td>40</td>
<td>0.240</td>
<td>0.268 134</td>
</tr>
<tr>
<td>6</td>
<td>10</td>
<td>1</td>
<td>-1</td>
<td>70</td>
<td>40</td>
<td>0.240</td>
<td>0.201 100.5</td>
</tr>
<tr>
<td>7</td>
<td>12</td>
<td>-1</td>
<td>1</td>
<td>50</td>
<td>80</td>
<td>0.240</td>
<td>0.286 143</td>
</tr>
<tr>
<td>8</td>
<td>5</td>
<td>1</td>
<td>1</td>
<td>70</td>
<td>80</td>
<td>0.240</td>
<td>0.177 88.5</td>
</tr>
<tr>
<td>9</td>
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<td>-1.682</td>
<td>0</td>
<td>43</td>
<td>60</td>
<td>0.220</td>
<td>0.265 132.5</td>
</tr>
<tr>
<td>10</td>
<td>4</td>
<td>1.682</td>
<td>0</td>
<td>77</td>
<td>60</td>
<td>0.220</td>
<td>0.16 80</td>
</tr>
<tr>
<td>11</td>
<td>7</td>
<td>0</td>
<td>-1.682</td>
<td>60</td>
<td>26</td>
<td>0.220</td>
<td>0.272 136</td>
</tr>
<tr>
<td>12</td>
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<td>0</td>
<td>1.682</td>
<td>60</td>
<td>94</td>
<td>0.220</td>
<td>0.193 96.5</td>
</tr>
<tr>
<td>13</td>
<td>20</td>
<td>0</td>
<td>0</td>
<td>1.682</td>
<td>60</td>
<td>0.186</td>
<td>0.227 113.5</td>
</tr>
<tr>
<td>14</td>
<td>2</td>
<td>0</td>
<td>0</td>
<td>1.682</td>
<td>60</td>
<td>0.253</td>
<td>0.234 117</td>
</tr>
</tbody>
</table>
Analysis of variance (ANOVA) and \( F \)-test were carried out on the regression model to evaluate its suitability and precision (Table 4-3).

**Table 4-3** ANOVA results of the quadratic model for methacrylation of KL\(^a\)

<table>
<thead>
<tr>
<th>Source</th>
<th>Deg. of freedom</th>
<th>Sum of squares</th>
<th>Mean squares</th>
<th>( F )-value</th>
<th>( P )-value (prob &gt; ( F ))</th>
</tr>
</thead>
<tbody>
<tr>
<td>Model</td>
<td>9</td>
<td>10783.5</td>
<td>1198.16</td>
<td>80.97</td>
<td>0.000</td>
</tr>
<tr>
<td>Lack of fit</td>
<td>5</td>
<td>85.1</td>
<td>17.03</td>
<td>1.36</td>
<td>0.373</td>
</tr>
<tr>
<td>Pure error</td>
<td>5</td>
<td>62.8</td>
<td>1.00</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>Total</td>
<td>19</td>
<td>10931.5</td>
<td>—</td>
<td>—</td>
<td>—</td>
</tr>
</tbody>
</table>

\(^a\) \( R^2 = 0.9865; \text{adjusted } R^2 = 0.9743; \text{predicted } R^2 = 0.9323 \)

The \( P \)-value < 0.05 for the regression model was taken as evidence for the significance and appropriateness of it for accurately predicting the reaction yields according to its
parameters. Moreover, The $F$-value of 1.36 and $P$-value of 0.373 (larger than confidence interval) for the “lack of fit” confirmed its insignificance in the developed model, further proving its good fitting. As well, the coefficient of determination ($R^2$) for the mass yield of methacrylation was 0.9865, meaning 98.65% of the mass yield variations can be explained by the independent variables based on the developed model, while the adjusted $R^2$ value was 0.9743, confirming that the model was not over-fitted. Additionally, the predicted $R^2$ value of 0.9323 showed a good agreement between the values predicted by the model and the real-time experimental responses. Moreover, the normal probability plot (Figure 4-1a) showed a normal distribution of errors, indicating a good prediction of the responses by the model, while the residual plot (Figure 4-1b), depicting the relationship between the residual values and the fitted values for each response, showed random scattering of the residuals around the zero line, further demonstrating the competence of the developed model.
Figure 4-1 The normal probability plot (a) and residual plot (b) of the obtained model.
4.3.2 Response Surface Plot and Optimization of Variables

The response surface plots (Figure 4-2a-c) and contour plots (Figure 4-2d-f) of the methacrylation of KL show a good agreement with the developed regression model. The developed plots and regression model were then employed to determine the maximum possible response of the reaction (recovered mass yield) obtainable from the minimum possible amounts of the reaction parameters. Following the predictions of the regression model and the depictions of the developed plots, it was determined that the response (mass yield) can reach values higher than 140 wt.% by keeping the reaction temperature between 50 °C and 60 °C, while having a 1MIM/KL molar ratio between 0.22 and 0.25 (Figure 4-2a and 4-2d).

Moreover, Figures 4-2b and 4-2e suggest that for the same response value range, the reaction time must be under 60 min, while Figures 4-2c and 4-2e give predictions in-line with the mentioned observations. On the other hand, the circular shape of the center of the contour plot, corresponding to mass yield (wt.%) > 140%, suggested that changes in each variable must be in a certain range and moving past said range will have a negative effect on the reaction mass yield (Figure 4-2d, 4-2e, and 4-2f). Thus, the process optimization was carried out via the response surface methodology to maximize the methacrylation mass yield, while minimizing the reaction parameters without negatively affecting the mass yield. The result suggested a methacrylation reaction run at 54 ℃ for 54 minutes, with 1MIM/KL molar ratio of 0.225. The predicted mass yield for such reaction was 148.13 wt.%.

Given the recommended reaction conditions, a new reaction was carried out, which yielded 146.5 wt.% ML and thus, the predicted values and experimental results were proven to be in a very good agreement. Considering the maximum theoretical yield of 150.25 such a high product mass yield is significant.
Figure 4-2 Response surface plots and contour plots of different variable values vs. recovered mass yield: effects of 1MIM/KL molar ratio and reaction temperature (a, d); effects of 1MIM/KL molar ratio and reaction time (b, e); and effects of reaction time and reaction temperature (c, f).
4.3.3 Optimized Product Characterization

The ML obtained via the optimized reaction conditions (Figure 4-3-I) was characterized in comparison with KL by $^1$H NMR and FTIR (Figure 4-3-II). Three new broad signals were observed in the $^1$H NMR spectrum of ML as compared to KL, with chemical shifts of $\delta_H = 1.89, 5.82,$ and $6.14$ which were attributed to the methyl functionalities and the methacrylate group’s two olefinic groups. Moreover, the FTIR spectrum of ML showed a decrease at $3200-3500$ cm$^{-1}$ (s, br) relative to KL, corresponding to significant removal of hydroxyl groups, as well as increases in the esters’ C=O stretching vibrations ($1735$-$1750$ cm$^{-1}$), C-O stretching vibrations ($1000$-$1320$ cm$^{-1}$) and C=C bending vibration regions ($650$-$1000$ cm$^{-1}$), suggesting successful grafting of methacrylate groups on lignin’s structure. Finally, the extent of methacrylation was determined by a modified version of the method reported by Argyropolous$^{22}$ in 1994 for the quantification of lignin’s hydroxyl groups. The most significant alterations were the change in the halo-phosphine reagent ([CH(CH$_3$)$_2$]$_2$PCl), the base (N(Et)$_3$), and the use of an external standard (PPh$_3$) for quantification purposes. Equal amounts of KL and ML were reacted with [CH(CH$_3$)$_2$]$_2$PCl separately in presence of excess amounts of N(Et)$_3$ at room temperature. The extent of methacrylation was then determined by comparing the integrations of resulting -OP regions in $^{31}$P{$^1$H} NMR spectra to an external standard solution of PPh$_3$ (Figure 4). Based on the obtained results, we determined that on average ca. 70% of the hydroxyl groups on KL were converted to methacrylate groups in ML.
**Figure 4-3** I) The methacrylation scheme under optimized reaction conditions and II) $^1$H NMR and FTIR spectra of (A) KL and (B) ML synthesized under the optimized reaction conditions.
Figure 4-4 $^{31}\text{P}$-$^1\text{H}$ NMR spectra of products obtained from the reaction of (A) KL and (B) ML with [CH(CH$_3$)$_2$]$_2$PCl. All peak regions were integrated according to peak at -4.6 ppm, corresponding to the external standard used in this procedure (PPh$_3$).

4.3.4 Characterization of the ML-based UV-Cured Coatings

ML (30 wt.%) was incorporated into a UV-cured coating sample as discussed before, containing EB-1360, a hexafunctional siloxane-based crosslinker (65 wt.%), HDMAP (5 wt.%) as the photoinitiator. To determine the effect of such incorporation, a UV-cured coating sample containing no ML was prepared as a reference through the same process (with 95 wt.% EB-1360 and 5 wt.% HDMAP) and the coatings were characterized in conjugate with each other.

Drop shape analysis was employed to evaluate the effect of introducing ML as a hydrophobic component into the coating system, as siloxane-based systems are known for their relatively low hydrophobicity and lignin was known to reduce the wettability of the system in which it is incorporated.$^{16,23,24}$ It was determined that while the ML-free coating sample had a water contact angle of 60.4 ± 1.8 degrees, such value was increased to 82.6
± 1.9 degrees in the ML-based coating. This increase in the hydrophobicity of the coating was attributed to the incorporation of ML into the coating system as a hydrophobic moiety.

To determine the crosslinking percentages of the coating, the FTIR spectrum of each cured coating sample was compared to that of the corresponding liquid formulation of the said coating before UV-curing, particularly with regards to the signal intensity at ca. 800 cm\(^{-1}\) (C=C bending vibrations). It was determined that while the crosslinking percentage of the ML-free coating was ca. 35.8%, the same value for the ML-based coatings as ca. 64.1%, suggesting better crosslinking in the coating system with the incorporation of the ML moiety. Finally, to compare the thermal stabilities of the prepared coatings, each cured coating sample was scraped off the glass surface using a surgical-grade scalpel and fully crushed by a mortar and a pestle. The obtained crushed coatings were then heated \textit{in vacuo} at 40 °C for 24 h, at which point, thermogravimetric analyses were performed on the samples. Moreover, the thermal stability of the KL and ML samples were also evaluated for references. It was determined that KL had an onset decomposition temperature of 200 °C, and it was lowered to 189 °C in ML and similarly, ML’s residual weight percent at 800 °C was ca. 36.7 wt.%, slightly lower than that of KL (40.1 wt.%). Consequently, it was determined that the ML-free and the ML-based coatings decomposed at 239 and 246 °C, respectively, while the residual weight percentage at 800 °C was 3.3 wt.% and 13.8 wt.% for the ML-free and the ML-based coatings, respectively. The above results suggest that the introduction of ML into the UV coating system had a positive effect on the coating’s thermal stability (Figure 4-5), likely attributed to the improved crosslinking percentages of the coating by the ML, which aligns well with the resulted reported in many literature studies on the positive effects of lignin on char formation.\textsuperscript{25–27}
Figure 4-5 Thermogravimetric analysis of KL (A), ML (B), plus ML-free (C) and ML-based (D) UV-cured coatings.

4.4 Conclusions

Kraft lignin was chemically modified via a methacrylation process. The process parameters (reaction time, reaction temperature, and 1MIM/KL molar ratio) for the lignin methacrylation were optimized through central composite design and response surface methodology to maximize the methacrylated lignin (ML) product yield. The optimum conditions for the lignin methacrylation, determined from the study, are as follows: 54 minutes at 54 °C with 0.225 1MIM/KL molar ratio, resulting in an ML mass yield of 146.5 wt.%. The obtained ML demonstrated to be curable in a UV-cured coating system, with positive effects on the hydrophobicity, crosslinking percentage, and thermal stability of the coating system in which it was incorporated. Thus, this work realized optimized chemical modification of lignin by methacrylation, demonstrated promising application of the methacrylated lignin as an environmentally friendly UV-curable material for UV-cured coatings.
4.5 References


Chapter 5

5 “Lignophines”: Lignin-based Tertiary Phosphines with Metal-Scavenging Ability

Lignin is a biopolymer that is a natural component of wood and a by-product from the pulp and paper industry. This material is naturally present in the secondary cell walls of plants, resulting from an in vivo enzyme-mediated dehydrogenation polymerization (lignification) of three hydroxycinnamyl alcohol monomers that include p-coumaryl, sinapyl, and coniferyl alcohol units (monolignols; Scheme 5-1a). Of the almost 50 million tons per year of lignin produced globally, only 2% is utilized for conversion to specialty products, while the rest is used either as low value heating fuel at the site of manufacture or it is disposed of as waste (published data from 2010). New outlets for up conversion and repurposing of this biopolymer need to be established. Lignin is a renewable resource that could be leveraged for a broad range of materials applications given the rich functional group chemistry present within the biopolymer, including hydroxyl, ether, carbonyl, and carboxylic acids, that can be accessed and transformed into value-added products.

Both the aromatic and aliphatic hydroxyl groups on lignin have been utilized as key reactivity outlets for valorization. For example, lignin’s phenolic hydroxyl groups have been selectively methylated using dimethyl sulfide or oxypropylated with propylene oxide, while acyl anhydrides and acyl chlorides have been employed to cap both aliphatic and aromatic hydroxyl groups. The key outlets of modified lignins include bio-based epoxy resins, thermo- and UV-cured graft copolymers, and flexible and rigid polyurethanes. However, the strategies for up-converting lignin have shifted towards developing novel materials with specific applications in recent years, such as cathode-active materials in primary lithium batteries and non-toxic bio-degradable nanoparticles with potential applications in drug delivery and pharmaceuticals.
Scheme 5-1 a) The three standard monolignols of $p$-coumaryl (A), coniferyl (B), and sinapyl (C) alcohol as the building blocks of lignin (KL, representative structure of lignin) and b) the reaction scheme for grafting lignin with hydrolysed MMA/MAPC1 copolymer P(MMA-co-MAPC$_1$(OH)$_2$) as reported by Ferry et al.

Despite the headway made in upconverting lignin, the elemental diversity installed into the material is woefully narrow and therefore, limits other potential outlets for an inexpensive, sustainable, and abundant biopolymer.

Polymerizable fragments (e.g. olefins, amides, and carboxyls), which contain unsaturated functional groups have been installed into lignin’s structure and they can be exploited to build elemental diversity into the material, offering further opportunity in creating a usable, value-added product. Because olefins are easily modified via E-H bond addition, impressive opportunities exist for adding B-H, Si-H, S-H, and P-H across the double bond, creating Lewis acidic or basic bio-based polymers.

Our group has been particularly interested in utilizing transformations of P-H bonds to prepare highly functional phosphorus-based polymeric materials, including self-healing
polyphosphonium ionic networks, P-based interpenetrated polymer networks (IPNs),
and highly crosslinked phosphorus photopolymer networks, which have been applied as
effective antimicrobial agents, targeted anion self-assembly, and in metal scavenging
applications.

Reported procedures on grafting phosphorus-containing compounds onto lignin’s
structure, to this date, have solely focused on exploiting the hydroxyl functionalities on
lignin for the formation of P-O functionalities along the macromolecule’s structure.
Argyropolous et al. have studied such exploitation in detail to develop a procedure capable
of quantifying lignin’s hydroxyl functionalities through grafting these P-O
functionalities. Ferry et al. reported a graft co-polymer of lignin using hydrolyzed
methylmethacrylate/dimethyl(methacryloxy)-methyl phosphonate (MMA/MAPC1)
copolymer (Scheme 1b), which introduced phosphorus atoms onto lignin via P-O linkages.
They then blended the synthesized copolymer with polybutylene succinate (PBS) pellets
using compression molding to improve the fire behavior of PBS. However, there have
been no reports on the preparation of polymer networks that contain lignin as the backbone
and phosphorus as a substituent bound to the network via P-C linkages, which can expand
lignin’s utility by exploiting the diverse chemistry of phosphorus in derivatizing the
network after polymerization. In this context, we have utilized methacrylated lignin in a
phosphane-ene click reaction with PH$_3$(g). The resulting product is rich in P-H bonds,
which can be used in subsequent phosphane-ene chemistry with substituted olefins. This
creates an array of tertiary alkyl phosphines on the lignin scaffold, which behaves as a
polyphosphine and is amenable to metal coordination and metal scavenging. These results
represent a major shift in the immense, unlocked potential for lignin as a commodity
chemical, rather than negative value waste.

Methacrylated lignin (ML, 200 mg, synthesized and quantified following the same
procedure as reported in chapter 4) was swelled in CH$_3$CN (100 mL). Azobis(isobutynitrile) (AIBN; 6 mg, 3 wt.%) was added to the swelled ML and the
mixture was transferred to a 300 mL autoclave reactor. The autoclave was then degassed
by purging N$_2$ for ca. 10 min and pressurized with phosphine gas (PH$_3$, 80 psi). The vessel
was then stirred for 1 h, at which point it was re-pressurized with PH$_3$ and heated to 50 °C
under stirring for 24 h. Afterward, the pressurized PH$_3$ was released in a controlled environment, where it was ignited and burned. Subsequently, the vessel was brought into the glovebox, opened, and the mixture (swelled light-brown substance in solvent) was centrifuged and dried in vacuo to obtain MLPH$_2$ in the form of a light-brown powder (112.5 % recovered mass yield). A sample of the product was re-swelled in CH$_3$CN and the $^{31}$P{$^1$H} NMR spectrum of the sample (Figure 5-1) showed a signal at $\delta_P = -148$ consistent with molecular primary phosphines and the FT-IR spectrum of the same sample showed the clear presence of P-H functional groups (2300 cm$^{-1}$; P-H stretch; Figure 5-1) and therefore the product was assigned as the primary phosphine (MLPH$_2$), with 66% of the acrylate groups on ML successfully functionalized.

$^{31}$P{$^1$H} NMR and FTIR spectra of MLPH$_2$ (A), MLP$_{\text{Hex}}$ (B), MLP$_{\text{RFn}}$ (C), and the $^{31}$P{$^1$H} NMR spectrum of MLP$_{\text{Ag}}$.

The hydrophosphination percentage (EOHP) was calculated using a modified version of the quantification process as reported by Argyropoulos.$^{36}$ MLPH$_2$ (20 mg) was swelled in CH$_3$CN (2 mL) and transferred into an NMR tube. Separately, 40 µl of a 0.025 mol/L solution of PPh$_3$ in CH$_3$CN was transferred to a capillary tube, which was then flame sealed.
and inserted into the NMR tube containing the \textbf{MLPH}_2 sample and the $^{31}\text{P}\{^1\text{H}\}$ NMR spectrum of the mentioned tube was recorded (Figure 5-2).

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure5-2.png}
\caption{$^{31}\text{P}\{^1\text{H}\}$ NMR spectrum of \textbf{MLPH}_2 in comparison with PPh$_3$P as external standard ($\delta_P = -4$ ppm).}
\end{figure}

Another peak was observed at $\delta_P = -82$ ppm, consistent with molecular secondary phosphines, which was attributed to the result of two methacrylate groups linking to one phosphorus atom and forming a secondary lignophine (\textbf{MLPH}_1). The amount of the \textbf{MLPH}_1 formed was determined to be ca. 3\%, which was considered negligible in
determining the obtained material as primary lignophine (MLPH$_2$). The EOHP was determined as described below:

Considering ca. 70% of the -OH functionalities were methacrylated in ML:

$$1\text{mol } K L \times 1.5 \frac{\text{mol OH}}{\text{mol } 1} \times 70\% = 1.05 \text{ mol methacrylate on one C9}$$

$$\text{mod. C9} = 180\, \text{g} + \left( 1.05 \text{ mol meth.} \times 85.03 \frac{\text{g}}{\text{mol}} \right) - \left( 1.05 \text{ mol H} \times 1.008 \frac{\text{g}}{\text{mol}} \right)$$

$$= 268.24 \, \text{g}$$

Thus, approximately 0.6727 mol of the methacrylate groups on ML were capped.

Considering the integration of the peak at $\delta_P = -148$ ppm:

$$10^{-3}\text{mmol } PPh_3 \times \frac{3.01}{0.06} \times \frac{268.24}{20} = 0.6727 \, \text{mol}$$

Thus, approximately 0.6727 mol of the methacrylate groups on ML were capped.

Considering the integration of the peak at $\delta_P = -82$ ppm:

$$10^{-3}\text{mmol } PPh_3 \times \frac{0.05}{0.06} \times \frac{268.24}{20} \times \frac{2 \text{ mol meth.}}{1\text{mol MLPH}_1} = 0.0222$$

Thus, approximately 0.0222 mol of the methacrylate groups on ML were capped.

Finally, to determine the EOHP:

$$EOHP(\%) = \frac{0.6727 + 0.222}{1.05} \times 100 = 66\%$$

The resulting MLPH$_2$ was then heated with excess 1-hexene or 1H,1H,2H-perfluoro-1-hexene in the presence of AIBN at 40 °C to obtain tertiary alkyl or fluoroalkyl lignophine (Scheme 5-2), respectively (MLPH$_{\text{Hex}}$; MLPH$_{\text{RF4}}$). Both reaction mixtures were filtered, the precipitates washed and then dried in vacuo to give light brown solids in all cases. $^{31}\text{P}^{[1\text{H}]}$ NMR spectra of the swelled isolated materials each showed one signal at $\delta_P = -33$ and -29, respectively and consistent with the preparation of tertiary phosphines. Moreover, the FT-
IR spectra of the isolated material showed no P-H vibration confirming the absence of P-H functional groups (Figure 5-1).

Scheme 5-2 Reaction schemes for the synthesis of MLPH$_2$, MLP$^{RF_n}$, MLP$^{Hex}$, and coordination of MLP$^{Hex}$ to AgOTf.
Thermal gravimetric analysis of ML, MLPH₂, MLPHex, and MLPRFn showed that the MLPH₂ and MLPHex samples retained more of their respective weights at high temperatures in comparison with the ML sample (43 wt.% for MLPHex, 40 wt.% for MLPH₂, and 34 wt.% retained weight at 900 °C for ML), whereas the retained weight for the MLPRFn sample was almost equal to that of the ML sample (Figure 5-3).

![TGA thermograms of ML (A), MLPH₂ (B), MLPHex (C), and MLPRFn (D).](image)

**Figure 5-3** TGA thermograms of ML (A), MLPH₂ (B), MLPHex (C), and MLPRFn (D).

To ascertain whether or not the Lignophine materials are amenable for metal coordination, a sample of MLPHex was then added to a solution of AgOTf in CH₃CN (Scheme 2). The resulting mixture was then filtered, washed with CH₃CN and dried *in vacuo* to obtain a light brown powder. A swelled sample of the isolated material showed a new signal at δₚ = -4 in the ³¹P{¹H} NMR spectrum, a Δδₚ of 31 downfield from MLPHex that was consistent with metal coordination at phosphorus, giving MLPAg (Figure 5-1). The ¹⁹F{¹H} NMR spectrum of MLPAg was considerably broadened at δ_F = -77.5 and is consistent with an associated triflate ion (Figure 5-4).
Figure 5-4 $^{31}$F-$^1$H NMR spectra of AgOTf (A) and MLP$_{Ag}$ (B).

Given the ability of the lignophine to bind Ag$^+$, we considered that the material could be readily utilized as a metal-scavenger in catalytic reactions to sequester homogeneous catalysts from solution. As a proof of principle and using the methodology reported by Cuthbert et al.,$^{31}$ scavenging of the homogeneous metathesis catalyst Grubbs I (GI) was carried out (Figure 5-5a). Under the standard ring closing metathesis (RCM) conditions$^{37}$ GI converts 76% of diethyl diallylmalonate (2) to product 3 within 35 min (Figure 5-5b, Trace A). In separate reaction vessels, GI was first stirred with excess MLP$_{Hex}$ (5 mg/0.5 mg GI and 10 mg/0.5 mg GI) for the incubation periods of 1 h and 24 h. These suspensions were then treated with RCM substrate 2, where conversions of 11% and 8% were observed for the 5 mg/0.5 mg (MLP$_{Hex}$/GI) incubations for 1 h and 24 h, respectively, while the same values for the 10 mg/0.5 mg (MLP$_{Hex}$/GI) incubations of 1 h and 24 h were 5% and 0%, respectively (Figure 5-5b). The arrested catalytic conversion shows that GI is successfully sequestered from solution by MLP$_{Hex}$ and that maximum sequestration of GI can be achieved by pre-incubation with MLP$_{Hex}$ (10 mg/0.5 mg GI) for 24 h. In a separate experiment, MLP$_{Hex}$ was pre-functionalized with GI and the resulting solid was isolated and used as the catalyst in the RCM of 2. No RCM product was observed, which indicates
both that the sequestered metal is not catalytically active and that the GI is not released into the solution in appreciable amounts to initiate catalysis. This also confirms that the minor RCM conversions at short incubation times and less MLpHex amounts (B-D, Figure 5-5b) were due to incompletely sequestered GI.

![Reaction scheme of the RCM of 2 with GI without metal-scavenger (A) and with pre-incubated GI with MLpHex (B-E) and b) conversion (%) of 2 in the RCM reaction in the presence of GI without metal-scavenger (A), GI pre-incubated with 5 mg MLpHex for 1 h (B) and 24 h (C), and GI pre-incubated with 10 mg 1PHex for 1 h (D) and 24 h (E).](image)

**Figure 5-5** Reaction scheme of the RCM of 2 with GI without metal-scavenger (A) and with pre-incubated GI with MLpHex (B-E) and b) conversion (%) of 2 in the RCM reaction in the presence of GI without metal-scavenger (A), GI pre-incubated with 5 mg MLpHex for 1 h (B) and 24 h (C), and GI pre-incubated with 10 mg 1PHex for 1 h (D) and 24 h (E).

In conclusion, tertiary lignophine was successfully synthesized through the reaction of methacrylated lignin (ML) with phosphine gas and subsequent reaction with 1-hexene.
(resulting in $\text{MLP}^{\text{Hex}}$) and $1\text{H},1\text{H},2\text{H}$-perfluoro-1-hexene (resulting in $\text{MLP}^{\text{RFn}}$). The obtained primary and tertiary lignophines were characterized by $^{31}\text{P}\{^{1}\text{H}\}$, $^{19}\text{F}\{^{1}\text{H}\}$ (for $\text{MLP}^{\text{RFn}}$), and FT-IR spectroscopy. Thermal gravimetric analysis was then performed on $\text{ML}$, $\text{MLPH}_2$, and $\text{MLP}^{\text{Hex}}$ and it was concluded that the lignophines exhibited higher retained weight and more char formation at 900 °C in comparison with the $\text{ML}$ sample. The ability of the obtained $\text{MLP}^{\text{Hex}}$ sample to coordinate to transition metals was probed using AgOTf and it was concluded (via $^{31}\text{P}\{^{1}\text{H}\}$ $^{19}\text{F}\{^{1}\text{H}\}$ NMR spectroscopy) that the synthesized tertiary lignophine has the ability to coordinate to the silver atom in AgOTf. The addition of lignophine to RCM reactions catalyzed by GI shows nearly complete catalyst quenching. This indicates that the polymer is an effective metal scavenger and opens a new door for opportunities with lignin a renewable bioresource.

5.1 References


(18) Cateto, C. A.; Barreiro, M. F.; Rodrigues, A. E.; Brochier-Salon, M. C.; Thielemans, W.; Belgacem, M. N. Lignins as Macromonomers for Polyurethane Synthesis:


Chapter 6

6 “Lignophoniums”: Stimuli-Responsive Lignin-based Polyphosphonium Networks for Controlled Drug Delivery

6.1 Introduction

Lignin is the world’s second most abundant polymer, after cellulose. Residing mainly in the secondary cell wall, lignin acts as a filler vital to plant life, as it is responsible for giving stiffness, as well as facilitating the transport of water and other nutrients throughout the plants.\textsuperscript{1,2} This complex natural polymer originates from the connection of three phenylpropanoid (C9) units (\(p\)-hydroxyphenyl, H; guaiacyl, G; and syringyl, S) through various, primarily arylglycerol ether, linkages.\textsuperscript{3} The C9 units and their respective linkages are resulted from an \textit{in vivo} enzyme-mediated dehydrogenation polymerization (lignification) of three hydroxycinnamyl (\(p\)-coumaryl, sinapyl, and coniferyl) alcohol monomers called monolignols (Figure 6-1a).\textsuperscript{4–6}

As a sustainable resource with the annual global production of 50–55 MT, more than 98% of the produced lignin is used at its manufacturing plants as a fuel for heat or buried in landfills as waste.\textsuperscript{7} Such under-utilization, along with a functional group-rich chemical structure has turned the research field of lignin valorization into a highly attractive area for many researchers.\textsuperscript{8,9} The up-conversion of lignin has been widely targeted by chemically modifying and incorporating it into different materials such as epoxy resins,\textsuperscript{10–13} coatings,\textsuperscript{14–16} polyurethanes,\textsuperscript{17–20} and adhesives.\textsuperscript{21–24} Approaches to lignin valorization strategies have gained substantial interest over the past few years, which aim at converting lignin into materials with more unique applications. For example, lignin has been chemically modified to produce carbon fibers, nanofibers, and highly hydrophobic ATRP-polymerized graft copolymers,\textsuperscript{25–33} all hallmarks of the immense potential lignin has in areas such as biomedicine and medicinal chemistry. Lignin has been probed as a component in systems capable of controlled drug release, mainly in the form of nanocarriers\textsuperscript{34–37} and hydrogels.\textsuperscript{38,39} Such new streams of up-conversion strategies can play
an important role in the future of lignin valorization and become the next step in turning lignin from a negative-value material to a positive-value one.

Figure 6-1 The three standard monolignols of \( p \)-coumaryl (A), coniferyl (B), and sinapyl (C) alcohol and their corresponding \( p \)-hydroxyphenyl (H), guaiacyl (G), and syringyl (S) C9 units as the building blocks of lignin (KL, representative structure of lignin).

Phosphorus chemistry has yet to find its own stream in lignin valorization. While chemical modification of lignin with phosphorus-containing compounds, particularly halophospholanes, has been used extensively since mid-1990s for the purpose of quantifying lignin’s hydroxyl functionalities,\(^40\)\(^-\)\(^42\) reports on utilizing lignin-based phosphorus containing materials remain rare and generally focus on using phosphates in conjunction with lignin for improved surface area or flame retardancy in the prepared materials.\(^43\)\(^-\)\(^46\) Regardless of aiming at valorizing lignin or not, these reports focus on utilizing -O-P functional groups within the structure of lignin, where our group remains the only one examining lignin modification processes through the introduction of -C-P functionalities. A general research stream of our group relies in P-H bond addition reactions to olefinic groups to synthesize highly functional phosphorus-rich materials via phosphane-ene chemistry that have metal and oxygen scavenging abilities,\(^47\),\(^48\) self-healing networks,\(^49\) antimicrobial properties,\(^50\)\(^-\)\(^52\) and act as ceramic precursors.\(^53\) Recently, we
have synthesized lignophines, lignin-based tertiary polyphosphines, that show effective metal-scavenging ability.

In this context, we have synthesized lignophonium, a lignin-based polyphosphonium salt, from tertiary alkylated lignophine, \( \text{MLP}^{\text{Hex}} \). The isolated lignophonium was then incorporated into siloxane-crosslinked UV-cured thin films, which showed high cure percentages, water contact angles and surface charge density value well over the necessary threshold for antimicrobial films. The swelling degrees were also determined to be suitable for sustained drug release.

### 6.2 Experimental

#### 6.2.1 Materials and Methods

Lignophine (\( \text{MLP}^{\text{Hex}} \)) was synthesized according to the procedure reported in chapter 5 of this thesis. Solvents were purchased from Caledon and dried using an MBraun Solvent Purification System (SPS). EBERCYL 1360 (EB-1360; hexafunctional siloxane-based crosslinker) was purchased from Allnex Belgium SA and used as received. 2-hydroxy-2-methylpropiophenone (HDMAP, 97%), 4-vinylbenzyl chloride (90%), cetyltrimethylammonium chloride (CTAC; 25 wt.% in \( \text{H}_2\text{O} \)), and fluorescein (sodium salt) were purchased from Sigma-Aldrich Co. and used as received. Phosphate buffered saline (PBS) (0.15 M) was prepared following a previously reported procedure.\(^{54}\)

\(^{31}\text{P}\{\text{H}\} \) NMR spectra were recorded using a Varian INOVA 600 MHz spectrometer (\( ^1\text{H}, 599.5 \text{ MHz}; ^{13}\text{C}, 150.8 \text{ MHz}; ^{31}\text{P}, 242.6 \text{ MHz} \)) and referenced using an external standard (85\% \( \text{H}_3\text{PO}_4; \delta_P = 0 \)). FT-IR spectra were recorded using a PerkinElmer FT-IR spectrometer using the universal attenuated total reflectance mode (UATR), a diamond crystal, as well as the UATR sampling accessory (part number: L1050231).

#### 6.2.2 Synthesis of Lignophonium Salt (\( \text{MLP}^+ \))

\( \text{MLP}^{\text{Hex}} \) (0.600 g) was added to MeCN (30 mL) under stirring in a 100 mL pressure round-bottom flask in the dry-box and swelled for 10 min. 4-vinylbenzyl chloride (0.300 g, 1.965 mmol) was added to the mixture, the flask was sealed, moved out of the dry-box, and heated in an oil bath at 45 °C for 18 h. The pressure flask was cooled, brought back into the glove
box, and the mixture was centrifuged to obtain a dark brown swelled powder at the bottom of the centrifuge tube. The swelled powder was then washed three times with MeCN (20 mL), centrifuged after each wash, and the solvent was disposed of. Afterward, the swelled powder was transferred into a 50 ml Schlenk round-bottom flask and dried in vacuo at room temperature to obtain a light brown powder (109 wt% recovered mass yield).

6.2.3 Preparation of MLP+ UV-Cured Films

MLP+ (15 wt.%), EBERCYL 1360 (EB-1360, hexafunctional siloxane-based crosslinker, 80 wt.%), and 2-hydroxy-2-methylpropiophenone (HDMAP, 5 wt.%) were combined and diluted to 50% in N,N-dimethylformamide (DMF). 50 µL of the resulting mixture was then deposited on a clean glass slide. A second glass slide, using ~ 170 µm thick electrical tape as a spacer to define the film’s thickness, was then placed on top. The slides were passed 5 times at the speed of ~ 0.06 m•s−1 through a modified UV-curing system purchased from UV Process Supply Inc. (Chicago, IL, USA) with a mercury bulb (UVA: 154 mW•cm2, 189 mJ•cm2, UVB: 74 mW•cm2, 90 mJ•cm2; as determined by a PP2-H-U Power Puck II purchased from EIT Instrument Markets). After irradiation, the glass slides were separated, and the resulting film was dried in vacuo overnight at 40 °C, resulting in an MLP+ UV-cured film of ~ 150 µm thickness as measured by a micrometer.

6.2.4 Measurement of Cure Percentage

ATR-FTIR spectra were obtained from the liquid formulation before curing and the MLP+ UV-cured film after drying overnight in vacuo at 40 °C. The signal corresponding to C=O (~ 1720 cm−1) was used as an internal standard to normalize the obtained spectra, as its intensity does not change after curing. The intensity of the signal corresponding to C=C (~ 810 cm−1) was then compared between the liquid formulation and the UV-cured film. The cure percentage was calculated as the decrease (%) in the relative intensity of the C=C signal (100% corresponding to no C=C functionality remaining in the film’s ATR-FTIR spectra). The cure percentage was measured in triplicate and the results are reported as the mean ± standard deviation.
6.2.5 Measurement of Gel Content

The MLP+ UV-cured film from the cure percentage analysis were used to calculate the gel content. The MLP+ UV-cured film was accurately weighed and immersed in excess MeCN (10 mL) for 18 h. The solvent was removed, and the film was dried in vacuo at 40 °C for 18 h and re-weighed. Gel content was then calculated as the percentage of mass (%) remaining relative to the initial mass of the film. The gel content was calculated in triplicate and the results are reported as the mean ± standard deviation.

6.2.6 Measurement of water contact angle

The MLP+ UV-cured film from the cure percentage analysis were used to measure the water contact angle. The MLP+ UV-cured film was placed on a glass slide and placed on the stage of a Kruss DSA100 Drop Shape Analyzer (KRUSS GmbH, Hamburg, Germany). Afterward, a 5 µL drop of deionized water was placed on the surface and a static drop shape analyzer was used to measure the water contact angle on the film’s surface. Water contact angle was measured in triplicate and the result is reported as the mean ± standard deviation.

6.2.7 Charge density determination

The charge density was determined following a previously reported procedure. After incubating the MLP+ UV-cured film in MeCN (as described above for the measurement of gel content) and drying the film in vacuo at 40 °C for 18 h, the MLP+ UV-cured film (0.28 cm²) was immersed in 10 mL of a 1% w/v aqueous sodium fluorescein solution for 18 h. The film was then immersed in 10 mL deionized water for 4 h and immersed in 10 mL of a 0.1% v/v aqueous cetyltrimethylammonium chloride (CTAC) solution. At different time points, the film was transferred to a fresh CTAC solution, while 10 mM, pH 8 phosphate buffer (100 µL) was added to the previous CTAC solution and its absorbance at 501 nm was measured. The concentration of accessible charges on the surface was then determined according to the extinction coefficient of fluorescein (ε = 54,760 M⁻¹⋅cm⁻¹), assuming that each fluorescein molecule binds to one accessible cation. The charge density was measured in triplicate and the result is reported as the mean ± standard deviation.
6.2.8 Measurement of Mass Swelling Ratio in Water

The MLP+ UV-cured film from the cure percentage analysis were used to measure the mass swelling ratio in water by immersing the film in 10 mL deionized water for 24 h. Afterward, the film was removed from the water, the excess water on the film was removed with Kimwipe, and the film was weighed. The value of \( \frac{\text{final mass} - \text{initial mass}}{\text{initial mass}} \times 100\% \) was calculated and reported as the mass swelling %. Mass swelling ratio in water was determined in triplicate and the value was reported as the mean ± standard deviation.

6.1 Results and Discussion

Tertiary alkylated lignophine (MLPHex, 0.60 g) was synthesized and isolated (See chapter 5), swelled in CH3CN (100 mL), and heated with excess 4-vinylbenzyl chloride at 45 °C for 18 h (Figure 6-2a). The mixture was filtered, the precipitate was washed and dried in vacuo to give a light brown powder (109 wt% recovered mass yield). The \(^{31}\text{P}\{^1\text{H}\}\text{NMR}\) spectrum of the swelled, isolated material showed one signal at \( \delta_{\text{P}} = 32 \), consistent with the preparation of a phosphonium cation (Figure 6-2b) and the product was assigned as the “lignophonium” (MLP+; lignin-based phosphonium salt). A mixture of MLP+ (15 wt.%), EB-1360 (crosslinker, 80 wt.%), 2-hydroxy-2-methylpropiophenone (HDMAP, photoinitiator, 5 wt.%) was diluted 2-fold with \( N,N\)-dimethylformamide (DMF), cast on a glass slide, and cured using a UV-curing conveyor, producing MLP+-incorporated cured networks as free-standing films (MLP+\text{N}) that were ca. 150 µm thick, with a surface area of 0.28 cm\(^2\) (Figure 6-2c).
Figure 6-2 a) Reaction scheme for the synthesis of MLP⁺ from MLP\textsuperscript{Hex} and b) $^{31}\text{P}\{^1\text{H}\}$ NMR spectrum of MLP⁺, and c) process scheme for the preparation of MLP⁺N UV-cured films.

Surface-accessible charge density was calculated using a procedure reported by Murata et al.\textsuperscript{55} through immersing the dried MLP⁺N film in a 1% w/v aqueous fluorescein sodium solution (Fl.; 5 mL) to allow for the replacement of the chloride anions with fluorescein via anion exchange. The film was then washed with water to remove any unbound fluorescein, then placed in a 0.1% v/v aqueous cetyl trimethyl ammonium chloride (CTAC, 5 mL) solution to release the surface-bound fluorescein into the solution. Phosphate buffer (10 mM; pH = 8) was then added to the CTAC solution and the absorbance at 501 nm was
measured (Figure 6-3a), which corresponds to the absorption of the conjugated π system in fluorescein. The concentration of accessible charges on the surface was then determined according to the extinction coefficient of fluorescein ($\varepsilon = 54,760 \text{ M}^{-1}\text{cm}^{-1}$), with the assumption that each fluorescein molecule binds to one accessible cation and it was determined that the MLP+N film has $5.33 \pm 0.38 \times 10^{18}$ cations per cm$^2$ (Figure 6-3b), a value well above the suggested minimum surface accessible charges required for antimicrobial activity.\textsuperscript{56}

**Figure 6-3** a) Process scheme for the measurement of surface charge density and b) UV-Vis of the CTAC solution containing the MLP+N at 12 h (A), 60 h (B), and 180 h (C).
Drop shape analysis was used to measure the water contact angle (WCA) and it was determined that the MLP$^+\text{N}$ film had a WCA value of $88 \pm 1.2^\circ$, which was more than twice the same value for standard siloxane-based UV-cured films prepared without MLP$^+$ in the formulation (Figure 6-4a). ATR-FTIR was used to determine the cure percentage for the MLP$^+\text{N}$ film, which was calculated at $84.7 \pm 0.3\%$. The same value for the standard film was calculated at $36.1 \pm 0.2\%$, showing improved crosslinking within the network because of the MLP$^+$-incorporation (Figure 6-4b). Moreover, it was determined that the MLP$^+\text{N}$ film swelled to $55.5 \pm 3.8\%$ its original weight after immersion in water for 24 h. The hydrophobicity of the film, along the degree of swelling make the films potential candidates for slow, controlled release of ionic molecules in aqueous media.

**Figure 6-4 a)** Drop shape images for standard siloxane-based films and the MLP$^+\text{N}$ film and **b)** ATR-FTIR spectra for both the liquid formulations and cured films of standard siloxane-based films and the MLP$^+\text{N}$ film showing the intensity reduction for the C=C signal.

The controlled drug release ability of the MLP$^+\text{N}$ films was probed by incubating the film in a solution containing 1 wt% diclofenac sodium salt (10 mL) for 24 h, followed by washing with deionized water to remove any unbound molecules and submerging into a PBS solution (10 mL) to release diclofenac. The concentration of released diclofenac into
the PBS solution from the film was then determined according to the extinction coefficient of diclofenac in PBS ($\varepsilon = 4245.8 \, \text{M}^{-1} \text{cm}^{-1}$), assuming that each diclofenac molecule binds to one accessible cation and it was determined that the MLP$^+$N film released $6.23 \pm 0.19 \times 10^{18}$ diclofenac molecules per cm$^2$ over the timespan of 173 h, while complete release was observed at 157 h timepoint (Figure 6-5), which was in close agreement with the surface charge density as measured earlier.

![Graph](image)

**Figure 6-5** Release profile of the MLP$^+$N film.

### 6.2 Conclusion

Lignin-based polyphosphonium salt (Lignophonium; MLP$^+$) was successfully synthesized through the reaction of lignin-based tertiary alkylated polyphosphine (tertiary alkylated lignophile; MLP$^\text{Hex}$) with 4-vinylbenzyl chloride and characterized by $^{31}$P$^1{\text{H}}$ NMR spectroscopy. The lignophonium was then incorporated into a UV-crosslinked free-standing film (MLP$^+$N) using a siloxane-based crosslinker. The obtained films showed high cure percentages and water contact angles, along with a surface charge density value well over the necessary threshold for antimicrobial films and swelling degrees suitable for controlled drug release. The films were probed for their controlled drug release ability
using diclofenac as the loaded drug and examining the release in PBS and showed a promising release profile.

6.3 References


Chapter 7

7 Conclusions and Recommendations for Future Work

7.1 General Conclusions

The goal of this thesis was to conduct novel research on lignin valorization using the chemical modification of the hydroxyl functionalities of lignin. These groups are the most reactive sites within the macromolecule, and the most practical in terms of susceptibility to chemical modification. The chemical modification of these groups provides several possibilities regarding post modification processes. To graft onto the macromolecule, vinyl groups were chosen, primarily because of their crucial role in materials science, particularly in free radical polymerization. This thesis aimed at finding a chemical modification capable of grafting kraft lignin with vinyl-containing functionalities in the most facile fashion, using mild reaction conditions contrary to the other processes typically used for modifying lignin modification. The grafted macromolecule was probed with regards to its ability to be used in applications such as UV-curable coatings, and advanced materials with more complex and higher-value applications such as metal-scavengers and drug release vehicles.

7.2 Detailed Conclusions

Despite the inherently UV-absorbing nature of lignin, grafting of methacrylate groups in place of its hydroxyl functionalities successfully modified lignin into a UV-curable material. The modified lignin was quantitatively analyzed using a more readily available quantification reagent than the ones reported in literature. The modified lignin was crosslinkable in UV-cured coating samples at percentages up to 31 wt.%. The effect of crosslinking the modified lignin in the coating systems was evaluated regarding different characteristics of the UV-cured samples, particularly crosslinking percentage, hydrophobicity, surface adhesion, thermal stability, and retained weight at high temperatures. The chemical modification process developed for lignin (methacrylation) was then optimized using response surface methodology and central composite design. The developed mathematical model for the methacrylation process optimized reaction parameters to obtain the highest possible yield of modified lignin, while minimizing the
reaction temperature, time, and lignin/catalyst ratio. The optimized modified lignin was then examined at 30 wt.% in a siloxane-based UV-cured coating system as a proof of concept to determine the practicality of the developed optimization process.

The methacrylated lignin was utilized in a hydrophosphination reaction to report the first ever grafting of -C-P functionalities onto lignin, yielding lignin-based primary polyphosphines, referred to in this thesis as “primary lignophines”, which were then converted into lignin-based tertiary polyphosphines, referred to in this thesis as “tertiary lignophines,” before examining them in metal-scavenging settings. It was determined that the tertiary lignophine can coordinate to transition metals (e.g. silver). The tertiary lignophine was also probed as an effective transition-metal scavenger in lab-scale settings. It was determined that the tertiary lignophine had the ability to effectively sequester a catalyst (Grubbs I) in a ring closing metathesis reaction of diethyl diallylmalonates. This is the first report of lignin being used as a catalyst scavenger under normal catalytic conditions and an interesting potential new application for lignin.

Once the lignophines were synthesized and fully characterized, they were utilized in a quarternization process to yield lignin-based polyphosphonium salts, referred to in this thesis as “lignophoniums,” which were then crosslinked with a siloxane-based crosslinker to give free-standing lignophonium films. Moreover, the films were examined with regards to their surface accessible charge density, hydrophobicity, swelling degree, and cure percentage, showing excellent surface charge density values, well over the threshold required to create antimicrobial films. The films were also probed as effective controlled drug release materials by examining the release profile of diclofenac-loaded films in PBS.

7.3 Contributions and Novelty

The main novelties and contributions of the research reported in this thesis can be summarized as follows:

- First report of crosslinked UV-cured networks made from modified lignin (31 wt.%).
- First report on the optimization of lignin’s methacrylation process, capable of being utilized in scale-up production.
First report of hydrophosphination of a lignin derivative and subsequent utilization in phosphane-ene chemistry.

First report of developing a lignin-based metal/catalyst scavenger.

First report of synthesis and preparation of polyphosphonium salts and subsequent networks (free-standing films) from lignin, with potential applications as antimicrobial films and effective controlled drug delivery vehicles.

### 7.4 Recommendations for Future Work

#### 7.4.1 Tunable Lignin-based UV-cured Coatings and Up-scale Production

Although the effect of incorporating chemically modified lignin into UV-curable lignin-based coatings was determined in this thesis, the effect of different crosslinkers in such systems remains unexamined. Utilizing different crosslinkers can give the ability to tune the coatings’ properties and their behavior on different substrates according to the end-use in mind and be an important step towards possible commercialization of the lignin-based UV-curable coatings. Moreover, while scaled up methacrylation processes were performed by our research group in a scale up to 1L (reactor size), larger scale production of methacrylated lignin can be an important step in further proving the practicality of the optimized methacrylation process and make methacrylated lignin an interesting starting material for different target biomaterials. Preparing a mixed package comprised of up-scale production of methacrylated lignin can be an interesting package, capable of being proposed to coating industries.

#### 7.4.2 Design of Experiments for Up-scale Production of Primary Lignophines

Design of experiment, like the central composite design used in chapter 4 of this thesis can be utilized to optimize the hydrophosphination of lignin as reported in chapter 5 of this thesis. Optimizing this process will give the ability to produce primary lignophines in much larger amounts, which can be greatly impactful on potential industrial applications of these materials. Moreover, the same approach can be utilized for the synthesis of tertiary
lignophines to prepare a complete synthetic package, capable of synthesizing each material in desired yields using optimized reaction settings.

### 7.4.3 Tunable Lignin-based Tertiary Polyphosphines

Changing the capping reagents in the synthesis of tertiary lignophines can give the ability to tune some properties of the obtained material such as swellability. As the lignophine materials are not solvent-soluble and both their characterization and utilization in metal scavenging settings relied on their swellability in common organic solvents, making more swellable materials by changing the capping reagent might enhance such properties in the final products. Capping reagents such as 4-penten-1-ol can be reacted with the primary lignophine \((\text{MLPH}_2)\) to give corresponding tertiary lignophine \((\text{MLP}^{\text{OH}})\). This can increase the swellability of the tertiary lignophine in common organic solvents considerably and affect the metal-scatenging ability of the material (Scheme 7-1).

![Scheme 7-1 Proposed reaction scheme for the synthesis of MLP^{OH}.

Moreover, less hindered capping reagents such as ethylene gas can also be utilized in the synthesis of tertiary lignophines (Scheme 7-2). The synthesized ethylated lignophine \((\text{MLP}^{\text{Et}})\) will consequently have hindrance around the phosphorus atoms. Such alteration can in turn induce the ability to scavenge oxygen, which can be of great interest as a highly novel packaging material.
7.4.4 Synthesis of Metal-Phosphides from Lignin-based Tertiary Polyphosphines

Tertiary lignophines were probed for their ability to coordinate to transition metals in this thesis. The next step in this research area can focus on coordinating the tertiary lignophines to different metal carbonyls such as Mo(CO)₆, W(CO)₆, and Cr(CO)₆. The resulting metal-coordinated networks can then be pyrolyzed to yield corresponding metal-phosphides of MoP, PW, and CrP (Scheme 7-3). These are promising candidates for lignophines to act as ceramic precursors, which have applications in several industries such as batteries and energy storage.

Scheme 7-2 Proposed reaction scheme for the synthesis of MLP_{Et}.

Scheme 7-3 Proposed scheme for the synthesis of metal-phosphides from MLP_{Hex} through coordination to different metal carbonyls (M = Mo, W, and Cr) and subsequent pyrolysis.

7.4.5 Studying the Tunability and Further Utilization of Lignin-based Polyphosphonium Salts

Using different halides as the quarternization reagents for the synthesis of the lignin-based polyphosphonium salts (“lignophoniums”), as well as different crosslinkers for the
preparation of the free-standing films can enhance the tunability of the material with regards to its wettability, cure percentage, and swelling degrees, which can enhance its usefulness in controlled drug delivery systems.

Moreover, antimicrobial studies need to be performed on the lignophonium networks to further demonstrate their antimicrobial performance in real-world or lab settings. Like the catalyst sequestration studies for the lignophines, collaborating with research groups active in the field of biology and microbial studies would lead to promising results for the lignophoniums.
Appendices

Appendix 1. Supporting Information for Chapter 3

UV-Vis spectra

UV-Vis spectra were obtained of the KL, ML, and DMPA (photoinitiator) samples to evaluate the effect of the methacrylation reaction on the UV-curability of lignin (Figure S3-1). It was determined that while the KL sample had a complete overlap in absorbance with the photoinitiator (hence the non-UV-curable nature of lignin), the absorbance is significantly reduced in the ML sample, proving the detrimental effect of the chemical modification on converting lignin into a UV-curable material.

Figure S3-1 UV-Vis spectra of kraft lignin, methacrylated lignin, and the photoinitiator (DMPA) used for the preparation of the lignin-based UV-cured coatings.
Reaction of Lignin with Chlorodi(isopropyl)phosphine

KL was reacted with chlorodi(isopropyl)phosphine ([CH(CH₃)₂]₂PCl) to determine the types and amounts of hydroxyl functionalities within the native sample (Scheme S1).

![Reaction Scheme](image)

**Scheme S3-1** Reaction of KL with chlorodi(isopropyl)phosphine ([CH(CH₃)₂]₂PCl).

Upon examining the $^{31}\text{P}[^{1}\text{H}]$ NMR spectrum of the obtained product, signals corresponding to the main product (KL-Pi, $\delta_{\text{P}} = 136.0-170.0$) and two bi-products, chlorodi(isopropyl)phosphine oxide (O-Pi, $\delta_{\text{P}} = 52.0$) and di(isopropyl)phosphino chlorodi(isopropyl)phosphonium chloride $^1$ (P-P$^+$; $\delta_{\text{P}} = 65.0, 64.0$, d, $^{1}\text{J}_{\text{P-P}}=275.3$ Hz, $^1[(\text{CH}_3)_2(\text{CH})]P^+(\text{Cl})-P[\text{CH(CH}_3)_2]_2]$, $\delta_{\text{P}} = -11.3, -13.0$, d, $^{1}\text{J}_{\text{P-P}}=275.3$ Hz, $^1[(\text{CH}_3)_2(\text{CH})]P^+(\text{Cl})-P[\text{CH(CH}_3)_2]_2]$) were observed (Figure S2).
Figure 0-2 $^{31}$P{${}^1$H} NMR spectrum of the product of the reaction of KL with chlorodi(isopropyl)phosphine.

O-Pi was determined to be the product of the reaction of chlorodi(isopropyl)phosphine with residual water present in the native KL sample and the formation of P-P$^+$ was attributed to the reaction of two molecules of chlorodi(isopropyl)phosphine.

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