

Proceedings



A preparation method of softwood lignin derivatives: US9347177B2 patent evaluation ⁺

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Abstract: This study, in the form of a patent evaluation, which is a technique for studying the infor-11 mation present within and attached to patents, describes the state of the art by introducing what has 12 been patented in relation to softwood lignin derivatives. Inventors have described and claimed, 13 through the US9347177B2 patent, a method for the preparation of derivatives of native lignin from 14 softwood sources having a certain aliphatic hydroxyl content. The invention covered by the patent 15 concerns the processes of treating or compounding macromolecular substances and compositions 16 of lignin-containing materials as well as lignin and products derived therefrom. To prove the con-17 cept of this invention, recovery of lignin derivatives has been carried out from three softwood spe-18 cies. Samples of each softwood biomass feedstock were treated using an acid catalyzed ethanol or-19 ganosolv pulping process under different conditions. As a result, the recovered lignin derivatives 20 may have an aliphatic hydroxyl content of 2.5-7 mmol.g⁻¹, a phenolic hydroxyl content of 2-8 21 mmol.g⁻¹, a molecular weight that varies in the range of 200-4000 g.mol⁻¹, and any suitable polydis-22 persity of 1-5. These features result in a product with more consistent antioxidant activity. Further-23 more, these recovered lignin derivatives may be advantageous when formulating such composi-24 tions, making these materials highly desired for wide applications. 25

Keywords: softwood; lignocellulosic material; lignin; polymer; patent.

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1. Introduction

Lignin is one of the three major biopolymers of the lignocellulosic biomass, account-29 ing for 10–25% w/w of its composition and about 30% w/w of organic carbon in the bio-30 sphere [1,2]. It consists of inter-unit linkages of three repeating phenyl-propane mono-31 mers, termed sinapyl alcohol (S), coniferyl alcohol (G), and p-coumaryl alcohol (H) (Fig-32 ure 1) [3]. Other monolignols may be present in infinitesimal concentrations [4]. The pro-33 portions of these monomers may vary depending on the type of the plant species (e.g., 34 hardwoods, softwoods, annual plants, etc.), and environmental conditions [1,5]. This en-35 dows lignin macromolecules with a diverse range of molecular weights and polydisper-36 sity [6]. Due to its composition and structure, lignin can serve as an environmentally 37 friendly, biodegradable, antimicrobial, and antioxidant substance to develop high-value 38 products with potential applications in the chemical, pharmaceutical, cosmetic, and food 39 fields [7]. 40

Lignin can be isolated in various forms by different extraction processes, which can 41 be classified into two major categories, namely sulphur and sulphur-free processes (Figure 2) [3]. Generally, extracting native lignin from lignocellulosic biomass results in lignin 43 fragmentation into numerous mixtures of irregular components [8,9]. Several pre-

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treatment methods classified into chemical, physicochemical, and enzymatic pre-treatments have been developed to investigate and allow the isolation and recovery of lignin from wood [10]. In addition, a number of alternative wood-lignin extraction processes have been developed but not yet industrially introduced. Among them, four major organosolv pulping methods tend to produce highly-purified lignin mixtures and are defined as [11]: 50

- 1. Alcell[®] process: This method involves the use of ethanol and solvent pulping;
- ASAM process: This method involves pulping with alkaline sulphite anthraquinone
 methanol;
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- Organocell process: This method uses methanol pulping followed by methanol, 54 NaOH, and anthraquinone pulping; 55
- Acetosolv process: This method uses acetic acid, hydrochloric acid, or formic acid 56 pulping. 57



Figure 1. Chemical structures of different lignin monomers: sinapyl alcohol (S), coniferyl alcohol59(G), and p-coumaryl alcohol (H) (Adapted from Tribot *et al.*, 2019 [3], with the permission of Elsevier, published under license. Copyright © 2019 Elsevier Ltd.).60



Figure 2. Different processes for the isolation of lignin from lignocelluloses wood matrix (Adapted63from Tribot *et al.*, 2019 [3], with the permission of Elsevier, published under license. Copyright ©642019 Elsevier Ltd.).65

These four commercially registered organosolv pulping methods (i.e., Alcell[®], 66 ASAM, Organocell, and Acetosolv) can yield lignin derivatives with excellent performance characteristics such as structural optimization, low inorganic impurities, and low 68 molecular weight, thereby opening up new potential applications [11,12]. The commonly 69 adopted solvents for organosolv pulping are ethanol, methanol, acetic acid, and formic 70

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acid, which are usually mixed with water and mineral acids as catalysts [7]. Taking into 71 account all of the above, organosolv pulping processes tend to be the most practical tech-72 nique used to recover lignin material from biomass feedstock. 73

Research on lignin and its derivatives is developing rapidly through the innovation 74 and improvement of raw materials, chemical synthesis, methods of preparation, and for-75 mulation. Nevertheless, to promote the sufficiency of lignin in potential applications, sev-76 eral researchers have investigated pathways to enhance lignin properties to meet physi-77 cochemical and biological requirements. Compared to the use of cellulose, which has been 78 commercialized for centuries, commercial applications of lignin use are rare. That is why 79 there is a great interest in developing future strategies for lignin utilization, especially for 80 softwood lignin. 81

This study, in the form of US9347177B2 patent evaluation, which is a technique for 82 studying the information present within and attached to patents, describes the state of the 83 art by introducing what has been patented in relation to methods used to recover deriva-84 tives of native lignin with a certain aliphatic hydroxyl content from softwoods as biomass 85 raw materials. Both regarding preparation methods and applications, this study suggests 86 a summary of the incorporation of recovered lignin derivatives in different polymer com-87 positions for the purpose of providing additional functionality, such as enhancing their 88 antioxidant property. Furthermore, samples of hybrid spruce trees, radiata pine, and lob-89 lolly pine were studied under various conditions to demonstrate the concept of the pre-90 sent invention. 91

2. Patent analysis

The studied patent was invented by Balakshin *et al.*, and it was filed by the company Fibria Innovations Inc. (Burnaby, BC, Canada) in 2015. The earliest priority date of this patent was May 28th, 2009, with 64 patent families [13].

The jurisdictions of the related 64 patent documents (i.e., patent applications and granted patents) correspond to the United States (27 patents), Canada (11 patents), Europe (8 patents), China (8 patents), Brazil (5 patents), and the global system for filing patent 98 applications (5 patents), known as the Patent Cooperation Treaty (PCT), and administered 99 by the World Intellectual Property Organization (WIPO) [14]. The evolution of patent documents, related to the US9347177B2 patent, as a function of application filing year, granted 101 year, and publication year is presented in Figure 3.



Figure 3. Filed, granted, and published patents concerning the related 64 patent families of the104US9347177B2 patent [15].105

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The International Patent Classification (IPC) is a hierarchical system in the form of 106 codes, which divides all technology areas into a range of sections, classes, subclasses, 107 groups, and subgroups. It is an international classification system that provides standard 108 information to categorize inventions and evaluate their technological uniqueness [16]. The 109 relevant IPC codes concerning the studied US9347177B2 patent are presented in Table 1. 110

Table 1. Relevant classifications of the US9347177B2 patent [16].

IPC codes	Description						
C07G1/00	Low-molecular-weight derivatives of lignin						
C08H7/00	Lignin; modified lignin; high-molecular-weight products derived therefrom						
C08J3/00	Processes of treating or compounding macromolecular substances						
C08K5/13	Use of organic ingredients and oxygen-containing compounds such as phenols and phenolates						
C08L23/02	2 Compositions of homopolymers or copolymers of unsaturated aliphatic hydrocarbons having only on						
	carbon-to-carbon double bond not modified by chemical after-treatment						
C08L57/00	Compositions of unspecified polymers obtained by reactions only involving carbon-to-carbon unsatu-						
	rated bonds						
C08L97/00	Compositions of lignin-containing materials						
C09K15/06	Anti-oxidant compositions containing organic compounds such as oxygen						
D21C11/00	Regeneration of pulp liquors						
D21H11/00	Pulp or paper, comprising cellulose or lignocellulose fibers of natural origin only						

3. Patent evaluation

3.1. Materials and methods

The properties of lignin materials differ greatly depending on the lignocellulosic 114 feedstock (e.g., hardwoods, softwoods, annual plants, etc.), and the method used to isolate 115 them from other biomass components, for which several processes have been suggested 116 (Figure 1). Through the US9347177B2 patent, the inventors described and claimed a 117 method for producing lignin materials from softwoods having a specific content of ali-118 phatic hydroxyl groups, resulting in a product with a predictable level of antioxidant ac-119 tivity. The claimed method includes pulping the fibrous biomass, heating the biomass, 120 isolating the lignin-rich material from the cellulosic pulp, and recovering lignin deriva-121 tives having an aliphatic hydroxyl content ranging from 2.5 to 7 mmol.g-1. 122

The invention was based on the principles of the organosolv pulping processes for 123 obtaining lignin derivatives with specific characteristics from softwoods. The inventors 124 have proposed, at first, different values of pulping conditions, including temperature and 125 pressure, which can vary between 100-300°C and 5-150 atm, respectively, under a reac-126 tion duration that varies between 1–300 min. Furthermore, the pH of the pulp liquor can 127 vary between 1.5–5.5. As well, the weight ratio of liquor-to-biomass may take different 128 values, such as 5:1-15:1, 5.5:1-10:1, and 6:1-8:1. Therefore, the adopted method at the end 129 allows the pulping of the biomass feedstock with an aqueous solution of an organic sol-130 vent (30% w/w or greater). The obtained liquor shows a pH value of 1–5.5. It was then 131 heated to about 100°C or greater. Simultaneously, the pressure of the reaction medium 132 was raised to about 5 atm or greater. These conditions were maintained for one minute or 133 longer, in order to separate the biomass feedstock constituents and recover the derivatives 134 of native lignin. 135

In order to prove the concept of the invention, recovery of the lignin derivatives has 136 been carried out from three softwood species including, hybrid spruce trees, radiata pine, 137 and loblolly pine grown in Canada, Chile, and the United States, respectively. Four sam-138 ples of each softwood biomass feedstock were treated using an acid catalyzed ethanol 139 organosolv pulping process under different conditions (Table 2). 140

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After pulping, the isolation of lignin derivatives was performed as follows: first, the 141 liquor was separated from the solids by pressing the pulped materials in a press to 142 squeeze out the liquid. The resulting liquor was then filtered through a coarse silk screen 143 to separate the chip residue from the liquor stream. Thereafter, the liquor stream was fil-144tered through fine filter paper to recover the lignin derivatives that are referred to as self-145 precipitated lignin derivatives (SPL). But, another part of these materials still remaining 146 in the filtered liquor is thus precipitated through dilution with cold water; this is referred 147 to as precipitated lignin (PL). The relative yield of each lignin derivative was determined 148 in reference to its original lignin (i.e., the sum of acid-insoluble lignin and acid-soluble 149 lignin) and the PL yield (% w/w) was calculated for each sample. 150

Table 2. Pulping conditions for the three softwood species at a 6:1 liquor-to-wood weight ratio.151

Softwood species	pН	Sulphuric acid (% w/w)	Time (min)	Temperature (°C)	Ethanol (% w/w)
Hybrid spruce trees	1.81-2.18	0.9–2.5	33-55	175-184	47-78
Radiata pine	1.92-2.63	0.23-1.9	33-110	179–194	44-61
Loblolly pine	1.83-3.2	0.13-2.1	39–79	170-199	42-73

3.2. Results and discussion

Benefiting from the chemical structure of the obtained lignin derivatives, including 153 phenolic and aliphatic hydroxyl content, they can be exploited in various polymer com-154 positions. They can comprise, in addition to the lignin derivatives, a polymer-forming 155 component and other ingredients such as adhesion promoters; dispersants; fillers and ex-156 tenders; fire and flame retardants; stabilizers; ultraviolet light absorbers, and viscosity 157 regulators. Based on the results, the inventors confirmed that the recovered lignin deriv-158 atives may have an aliphatic hydroxyl content of 2.5–7 mmol.g⁻¹, a phenolic hydroxyl con-159 tent of 2–8 mmol.g⁻¹, a molecular weight that varies in the range of 200–4000 g.mol⁻¹, and 160 any suitable polydispersity of 1–5. These features are correlated to the antioxidant capac-161 ity of lignin derivatives, which has been evaluated by the Radical Scavenging Index (RSI) 162 by using the 2,2-diphenyl-1-picrylhydrazyl (DPPH) assay. The inventors confirmed 163 through the invention that the normalized RSI (NRSI) value can range from 5 to 40.

3.2.1. Hybrid spruce trees

As results for the hybrid spruce wood chips, the yield of PL lignin derivatives varies 166 in the range of 40.42–71.84%; for which the content of the aliphatic hydroxyl groups was 167 assessed, ranging from 2.60 mmol.g⁻¹ (sample 1) to 3.04 mmol.g⁻¹ (sample 4). In addition, 168 the antioxidant activity of the analyzed PL lignin derivatives was evaluated by measuring 169 their radical scavenging capacity. For the hybrid spruce trees, the NRSI values vary in the 170 range of 26.23–30.64 (Table 3). 171

Table 3. Pulping conditions for different wood chip samples at 6:1 liquor-to-wood weight ratio and
characterization of the precipitated lignin.172173

Comulas	Hybrid spruce trees			Radiata pine			Loblolly pine					
Samples	1	2	3	4	1	2	3	4	1	2	3	4
pH	1.95	2.18	1.81	2.12	1.92	2.23	2.5	2.63	1.83	2.52	2.5	3.2
Sulphuric acid (% w/w)		0.9	2.5	1.2	1.9	0.9	0.47	0.23	2.1	0.35	0.38	0.13
Time (min)		55	36	41	33	55	57	110	39	79	73	53
Temperature (°C)		184	175	181	179	184	194	191	170	198	189	199
Ethanol (% w/w)		47	78	68	57	47	61	44	46	42	54	73
PL yield (% w/w)		40.42	71.84	70.59	48.76	38	48.55	27.01	29.8	16.4	39.3	23.3
Aliphatic hydroxyl contents (mmol.g ⁻¹)		2.66	2.75	3.04	2.80	3.31	3.66	3.78	2.65	2.90	3.42	3.81
NRSI	27.43	28.15	26.23	30.64	36.97	28.28	27.19	29.76	27.78	27.95	27.78	24.56

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As seen in Table 3, the yield of PL lignin derivatives of radiata pine wood chips is 176 lower than that of the previously mentioned hybrid spruce wood chip samples. It varies 177 between 27.01% and 48.76%, for which the aliphatic hydroxyl content of each sample was 178 determined. These contents ranged from 2.80 mmol.g⁻¹ (sample 1) to 3.78 mmol.g⁻¹ (sample 179 4). Antioxidant activity was also evaluated and showed NRSI values ranging from 27.19 180 to 36.97. 181

3.2.3. Loblolly pine wood chips

As results for this material, the yield of PL lignin derivatives varies between 16.4% 183 and 39.3%. For which the aliphatic hydroxyl content ranged from about 2.65 mmol.g⁻¹ 184 (sample 1) to 3.81 mmol.g⁻¹ (sample 4). In addition, the NRSI value showed low variety in 185 the range of 24.56 to 27.95 (Table 3). 186

4. Conclusions

Lignin has gained huge interest due to its attractive properties such as biodegrada-188 bility, non-toxicity, sustainability, antimicrobial and antioxidant properties. Through the 189 US9347177B2 patent, the inventors have successfully validated a method for the produc-190 tion of lignin derivatives from softwood species with a specific amount of aliphatic hy-191 droxyl groups, which is correlated to the antioxidant property. Based on patent classifica-192 tion, the invention covered by the patent concerns the processes of treating or compound-193 ing macromolecular substances and compositions of lignin-containing materials as well 194 as lignin and products derived therefrom. The proposed method, which is based on the 195 principles of the organosolv pulping processes, was used to obtain derivatives of native 196 lignin from hybrid spruce trees, radiata pine, and loblolly pine, resulting in a high yield 197 of lignin derivatives with significant antioxidant activity. The stable antioxidant activity 198 of these recovered lignin derivatives may be advantageous when formulating such com-199 positions, making these materials highly desired for wide applications, including, among 200 other things, nutritional supplements, inks, pigments, surfactants, oils, films, coatings, 201 and adhesives. 202

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Data Availability Statement: The data presented in this study is available within the US9347177B2 208 patent. 209

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Conflicts of Interest: The authors declare that the content of this article has no conflict of interest.213The authors have no relevant affiliations or financial involvement with any organization or entity214with a financial interest in or financial conflict with the subject matter or materials discussed in this215article.216

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