

Maderas-Cienc Tecnol 22(2):2020
Ahead of Print: Accepted Authors Version

DOI:10.4067/S0718-221X2020005XXXXXX

1
2 **EFFECT OF THERMAL TREATMENT VARIABLES ON THE**
3 **THERMOGRAVIMETRIC PROPERTIES OF EUCALYPT WOOD**

4
5 **Déborah Nava Soratto^{1, *}, Carlos Miguel Simões da Silva², Benedito Rocha Vital²,**
6 **Angélica de Cássia Oliveira Carneiro², Juliana Jerásio Bianche², Walter Torezani Neto**
7 **Boschetti², Thaís Pereira Freitas², Juliana Ceccato Ferreira³**

8 ¹Universidade Federal de Mato Grosso do Sul (UFMS), Campus CPCS, Chapadão do Sul,
9 MS, Brazil.

10 ²Universidade Federal de Viçosa (UFV), Departamento de Engenharia Florestal, Viçosa, MG,
11 Brazil.

12 ³Universidade Federal de Jataí (UFJ), Jataí, GO, Brazil.

13 **Corresponding author:** deborahsoratto@gmail.com

14 **Received:** August 24, 2018

15 **Accepted:** February 07, 2020

16 **Posted online:** February 08, 2019

17
18 **ABSTRACT**

19 Thermal treatments have the effect of reducing the hygroscopicity and improving the
20 resistance to microbiological attack of wood by the degradation of its chemical constituents.
21 During the treatments, the mass of the wood is reduced, a factor that can affect the quality of
22 the materials according to their use. The objective was to verify the effect of the thermal
23 treatment variables on the thermogravimetric properties and the chemical composition of
24 *Eucalyptus grandis*. The treatments were carried out in a vacuum oven with three atmosphere
25 conditions - vacuum; N₂; vacuum+N₂ at temperatures of 140, 180 and 220 °C for 6 hours. It
26 was observed that the mass loss during treatments differed only according to the temperatures
27 used. The extractive content, total lignin and holocellulose presented significant changes only
28 at 220°C in all three atmospheres. In the thermogravimetric analysis, the greatest value of
29 residual mass was found in the treatment that used nitrogen and 220 °C, thus demonstrating
30 that this treatment was more invasive, leading to the conclusion that the vacuum application
31 can help to reduce the degradation of the constituents of the eucalypti wood. wood, which can
32 lead to the production of thermally treated wood without great losses in the mechanical
33 properties.

34 **Keywords:** *Eucalyptus grandis*; chemical composition; mass loss; thermally treated wood
35 Thermogravimetric analysis.

36

INTRODUCTION

37

38

39 According to Xing & Li (2010), wood, for some applications, may require greater
40 biological durability and better dimensional stability and, according to Inari *et al.* (2007),
41 thermorectification or heat treatment is a way of modifying the chemical characteristics of the
42 wood. For instance, Ayata *et al.* (2017) cited significant improves on the resistance to fungi
43 decay for both softwood and hardwood species. Furthermore, Yildiz *et al.* (2006) add that the
44 heat treatment can modify the aesthetic appearance, hygroscopicity, and weather resistance, in
45 addition to improving dimensional stability and durability. This mode of treatment of wood
46 can be done in a number of ways, varying the temperatures used, the time the material
47 remains exposed to temperature, the atmosphere of the treatment site and the catalyst used.
48 Also, there is the influence of the specific characteristics of the treated wood species.

49 During the application of the treatments, the mass of the treated parts reduces, and the
50 authors Welzbacher *et al.* (2007) reported that this decrease due to the heat treatment could be
51 a reliable and accurate parameter to predict the resistance to decomposition of the thermally
52 treated wood.

53 Esteves & Pereira (2009) subjected wood samples to heat treatment in an inert
54 atmosphere, with temperatures between 200 and 230°C, and verified a reduction of the
55 hydrophilic characteristic of the wood due to the modification of the carbohydrates and lignin
56 present in it. Likewise, Martinka *et al.* (2014) report that using thermal treatment of spruce
57 wood increases the lignin content, reduces the proportion of hemicelluloses, and also slightly
58 reduces the degree of average polymerization of the cellulose. These authors further describe
59 that these modifications occur when increasing the maximum temperature and the treatment
60 time.

61 At temperatures below 150°C, the volatilization of some extractives and the loss of
62 free water while above this temperature break the chemical bonds of more stable products,
63 leading to the formation of acetic acid, formic acid, methanol, CO and CO₂ (Bourgois &
64 Guyonnet 1988). Dehydration of the wood also occurs through the loss of constituent water,
65 detected by the decrease in the concentration of OH-groups present in the wood. According to
66 Fengel and Wegener (1989), the heating of cellulose at high temperatures and in air
67 atmosphere leads to the production of carbonyl groups and hydroxyl groups due to the
68 presence of oxygen. However, hemicelluloses are more susceptible to chemical reactions of
69 degradation and less tolerant of the action of heat due to the presence of hydroxyl groupings

70 and their amorphous condition Watanabe *et al.* (1989).

71 Other studies may show that some treatment variables affect the final product more
72 than others, as found by Zanuncio *et al.* (2014), who verified that the temperature used was
73 more effective in altering the chemical composition of the wood than the duration of the
74 treatments. Moura *et al.* (2012), for example, by investigating wood of two different species,
75 *Eucalyptus grandis* and *Pinus caribaea* var. *hondurensis*, found that the increase in the
76 maximum treatment temperature resulted in higher mass losses, and they also mentioned that
77 the losses were higher in the former than in the wood of the latter species. Thus, also relevant
78 is the chemical composition of each species of wood, especially between hardwoods and
79 softwoods. The composition of the hardwood hemicelluloses has an influence on the greater
80 loss of mass. Softwoods contain mainly arabinoglucoronoxylan and galactoglucomannan,
81 while hardwoods contain a lower amount of glucomannan and mainly glucoxytan, which are
82 strongly acetylated, compared to softwood hemicelluloses (Fengel e Wegener 1989, Sjöström
83 1981). In addition, acetic acid released during the deacetylation of hemicellulose catalyzes the
84 depolymerization of less ordered carbohydrates such as hemicelluloses and amorphous
85 cellulose (Prins *et al.* 2006).

86 The atmosphere conditions used for the thermal treatment, such as vacuum, nitrogen,
87 steam or other one that limit the O₂ concentration, can influence directly on its efficiency and
88 final quality (Wentzel *et al.* 2019). Araújo *et al.* (2012), comparing wood treated in the press
89 and in the vacuum oven, concluded that the thermal treatment in the press can produce
90 undesirable effects on the mechanical properties, while the material treated in the vacuum
91 oven with a controlled atmosphere obtained lower losses in the same properties. In the same
92 way, Araújo *et al.* (2016) also verified that heat treatment in a vacuum resulted in a modified
93 wood that was less hygroscopic and significantly different when compared to wood treated
94 under a nitrogen atmosphere.

95 Candelier *et al.* (2013) used vacuum and nitrogen atmospheres with a temperature of
96 220°C in both, and concluded that the effect of the vacuum allows the removal of products
97 from the degradation of the volatiles produced by the wood. This removal has the
98 consequence of limiting the acid degradation of polysaccharides, a phenomenon occurring in
99 the treatment with nitrogen, for example, due to the formation of acetic acid and the
100 recondensation of products of degradation of the volatiles inside the structure of the wood as a
101 result of the non-activation of the vacuum.

102 In this way, knowing that the heat treatment and its variables are the cause of the

103 chemical degradation of wood and consequently generate a loss of mass in the treated
104 material, it is possible to say that control of the mass loss can be a way to control the quality
105 of the wood. Therefore, this study is based on the investigation of the effects of heat treatment
106 variables, temperature and treatment atmosphere on the loss of mass of the treated wood of
107 *Eucalyptus* sp., seeking to answer which temperature and atmosphere should be used
108 according to the purpose of the wood.

109

110

MATERIAL AND METHODS

111

112 *2.1 Heat treatment*

113

114 *Eucalyptus grandis* boards were obtained from a sawmill located in the municipality
115 of Viçosa/MG, Brazil. The planks were conditioned in a dry, airy, covered environment,
116 arranged in stacked cells, until they reached a moisture content close to 12% (dry basis). The
117 initial dry weight was calculated based on the 12% initial moisture content. Subsequently, the
118 material was sectioned in pieces with mean dimensions of 60 x 7.5 x 2 cm (length, width and
119 thickness) and subjected to selection, discarding visibly non-standard parts or defects such as
120 knots and cracks. The samples were then identified and their mass values checked for
121 subsequent calculations of the mass loss of the respective treatments to which they were
122 subjected.

123

124 The heat treatment of wood was carried out in a Marconi vacuum oven model MA-
125 027 (São Paulo, Brazil), equipped with temperature, pressure, and vacuum control, and gas
126 application. Treatments were applied in three atmosphere conditions - vacuum; nitrogen (N₂);
127 vacuum + nitrogen (N₂). For the first two conditions, the material was placed in a vacuum
128 oven already heated to the desired temperature, which was then applied for 6 hours. For the
129 third condition, the samples were subjected to three levels: one hour at room temperature in
130 the vacuum; one hour at 140 °C in nitrogen, and four hours at the desired final temperature in
131 the same atmosphere as above.

131

132 *2.2 Properties evaluated*

133

134 In addition to the values of mass loss, the structural chemical composition - contents
135 of extractives, total lignins and holocelluloses - and the thermogravimetric properties of the
136 samples submitted to each of the treatments were evaluated.

137 The extractive contents were determined according to the TAPPI 204 om-88 standard
138 (TAPPI, 2002). The total lignins contents were estimated according to the methodology
139 followed by Pereira *et al.* (2013). Holocelluloses contents were determined by difference,
140 subtracting from 100 the contents of extractives and total lignins.

141 The thermogravimetric properties were determined in a Shimadzu DTG-60H
142 apparatus in an atmosphere of nitrogen gas at a constant flow rate of 50 ml/min. The samples
143 were ground and selected between 200 and 270 mesh overlapping sieves and were subjected
144 to controlled heating at an average rate of 10 °C.min⁻¹ to the maximum temperature of 600°C.

145

146 **2.3 Experimental design**

147

148 The experiment was conducted in a completely randomized design (DIC), in a
149 factorial arrangement, consisting of three atmospheric conditions - vacuum; nitrogen; vacuum
150 + nitrogen - and three final temperatures - 140, 180 and 220 °C, in three replicates, each
151 having six sample units. The data were submitted to Analysis of Variance (ANOVA) and
152 when significant differences were observed at 5% of significance, the means were compared
153 between treatments by the Tukey test and with respect to the control by the Dunnett test.

154

155

RESULTS AND DISCUSSION

156

157 **3.1 Loss of mass during treatments**

158

159 Table 1 shows the values of mass loss during the heat treatment of eucalypt wood. It is
160 observed that the increase in the treatment temperature promoted a greater loss of mass, while
161 the change in the treatment atmosphere did not result in significant changes in these values.

162

163

164

165

166

167

168 **Table 1.** Mass loss, in average percentage, occurring during the treatment of *Eucalyptus*
 169 *grandis*. wood as a function of the treatment temperature and atmosphere.

Effect	Condition	Mass loss (%)
Temperature of treatment	140°C	6.71 c
	180°C	11.16 b
	220°C	17.34 a
Atmosphere of treatment	Nitrogen	12.55 a
	Vacuum	11.32 a
	Vacuum + nitrogen	11.33 a

170 Means followed by the same lowercase letters, for each effect along each column, did not differ significantly
 171 from each other by the Tukey test at 5% probability.

172
 173 The mass losses of all the treatments applied are in accordance with the values
 174 indicated by the "Thermowood" process for heat treatment of wood. According to Esteves *et*
 175 *al.* (2008), this Finnish patent recommends a minimum mass loss of 3% in order to increase
 176 dimensional stability and at least 5% to increase the natural durability of the wood. However,
 177 some values found in the present study are significantly higher than those found by other
 178 studies, such as Brito *et al.* (2006), who treated wood of *Eucalyptus grandis* and Olarescu *et*
 179 *al.* (2013), who treated lime wood. These authors verified mass losses of up to, respectively,
 180 9.7% and 9.3% for woods subjected to thermal treatments at 200°C.

181 Araújo *et al.* (2016) heat-treated wood of different species native to Brazil -
 182 *Aspidosperma populifolium*, *Dipteryx odorata* and *Mimosa scabrella* - using treatment,
 183 temperature and atmosphere conditions similar to those of the present study. However, by
 184 using a treatment time of only one hour, the authors found mass loss values of at most 2.21%
 185 for treatment at 220 °C. In addition, mean values in the vacuum condition, relative to the
 186 nitrogen atmosphere, were significantly higher for some species. Compared with the present
 187 study, performed for a longer treatment time (6 hours), it can be observed that, in addition to
 188 the temperature, the heating time has great relevance in the thermal degradation of the wood.

189

190 **3.2 Chemical composition**

191

192 Table 2 shows the structural chemical composition of thermally treated eucalypt wood
 193 samples. Changes in the proportions of the components are observed, with increases in
 194 extractive and lignin contents and decreases in holocelluloses contents due to the heat

195 treatment in relation to the control samples. However, there was no statistical difference
 196 between the treatments, only those with respect to the control.

197

198 **Table 2.** Structural chemical composition of *Eucalyptus grandis* wood subjected to thermal
 199 treatment under different temperature and atmosphere conditions.

Treatment		Holocelluloses	Lignin	Extractives
Untreated		67.62	28.94	3.44
	140°C	65.22	29.78	5.00
Vacuum	180°C	66.46	29.75	3.79
	220°C	61.21*	32.03*	6.77*
	140°C	66.67	29.90	3.43
Nitrogen	180°C	65.60	29.58	4.82
	220°C	57.44*	34.81*	7.75*
Vacuum	140°C	66.22	30.01	3.77
+	180°C	63.71*	31.08	5.21
Nitrogen	220°C	58.96*	33.24*	7.80*

200 Means followed by * differ significantly from the control at 5% significance by the Dunnett test.

201

202 The sum of the contents of hemicelluloses and cellulose is denominated holocelluloses
 203 and corresponds to the most expressive mass fraction of the wood. The reduction in its
 204 contents, mainly in the treatment at 220°C, occurs with the thermal degradation of the wood
 205 starting at the temperatures used in this study. The results are in agreement with Zanuncio *et al.*
 206 *(2014)*, who evaluated the heat treatment of *Eucalyptus* wood at different temperatures up
 207 to 230°C and times up to 7 hours. The authors observed the reduction of holocelluloses
 208 contents from 66.27% of the control to 54.12% in the most altered samples. According to
 209 Yang *et al.* (2007) and Bach *et al.* (2014), in this temperature range, mass loss occurs mainly
 210 in the fraction of hemicelluloses due to its structure with a low degree of polymerization,
 211 which makes it less stable to heating compared to cellulose, which presents the most compact
 212 and polymerized structure.

213 The lignin content of wood treated at 220°C was significantly higher than the control.
 214 Similar results were found by Moura *et al.* (2012), Zanuncio *et al.* (2014) and Da Silva *et al.*
 215 (2016), who observed increases in the lignins concentration when thermally treating
 216 *Eucalyptus* wood. The lignins contents increased proportionally in the wood because of the

217 degradation of other constituents, especially the hemicelluloses, as discussed by Figueiró *et*
218 *al.* (2019). Lignins are phenolic macromolecules with high thermal stability (Yang *et al.*
219 2007) and with very low mass loss rates during heat treatment, thus explaining the
220 concentration of their contents in thermally treated wood.

221 In general, the treatments caused an increase, in proportion, in the extractive content.
222 However, this increase was only significant at the temperature of 220°C, regardless of the
223 method used, when compared to the control. These results corroborate the results already
224 found by Zanuncio *et al.* (2014) and Da Silva *et al.* (2016), who observed increases in the
225 extractive content for the heat treatments performed at 230 and 220°C, respectively. The
226 increase in the extractive content can be explained by the formation and modification of other
227 compounds, such as the degradation of the hemicelluloses, for example, forming soluble
228 products in the reagents used for the removal of extractives Esteves *et al.* (2011), and also by
229 the formation of compounds during the heat treatment, especially at the higher temperatures.

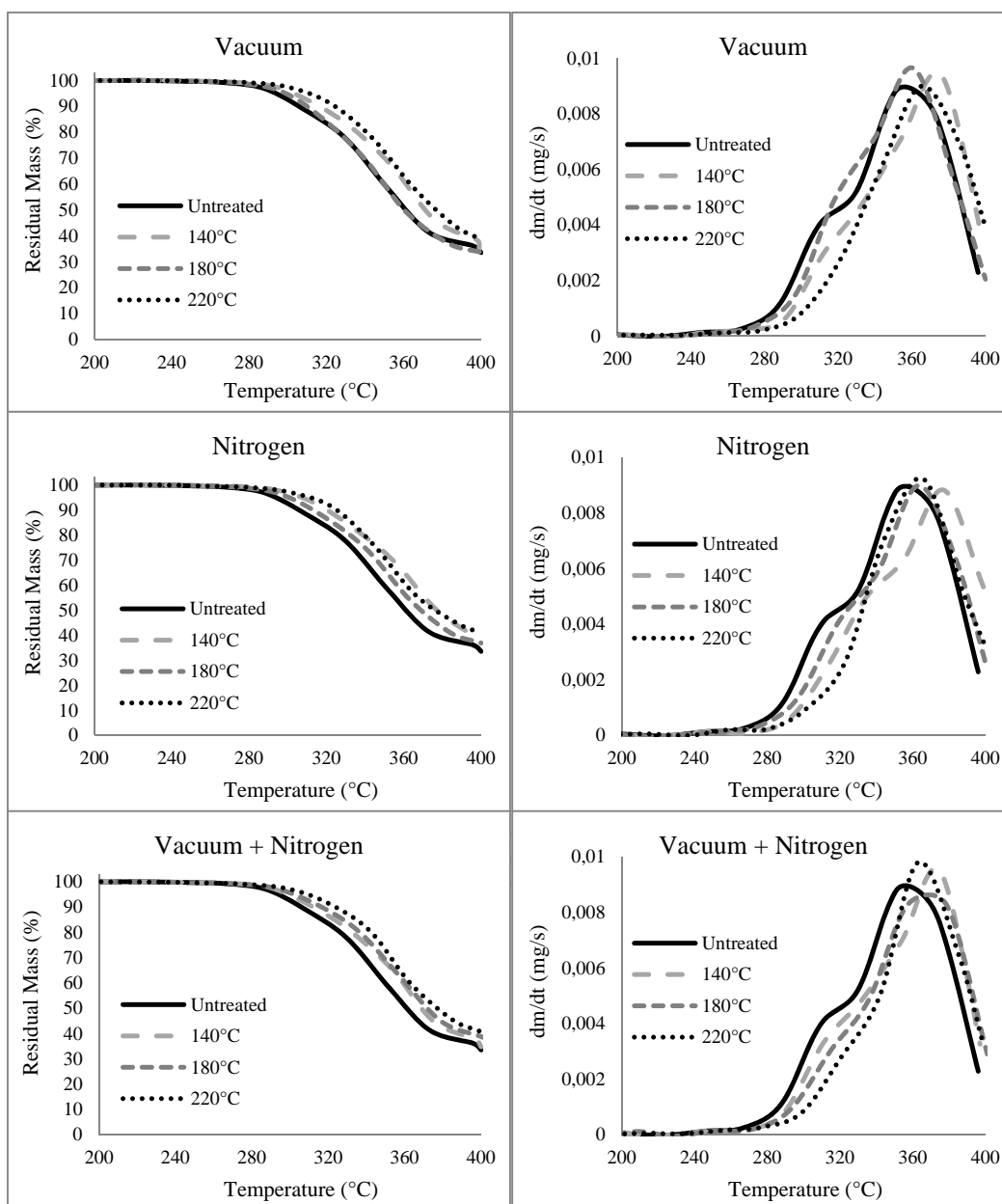
230

231 **3.3 Thermogravimetric analysis**

232

233 Figure 1 shows the curves obtained from the thermogravimetric analyses (TGA) and
234 their respective derivatives (DTG) of the thermally treated eucalypt wood under different
235 temperature and atmosphere conditions. The graphs were plotted between the temperatures of
236 200 and 400°C, because the thermal degradation was more relevant in this temperature range,
237 and they help to visualize the thermal degradation, allowing us to verify, with the derivatives,
238 the thermal decomposition peaks of the wood, which has already undergone heat treatment
239 previously.

240



241

242

243

244 **Figure 1.** Graphs obtained from the thermogravimetric analysis (TGA) and the respective
 245 thermogravimetric derivative (DTG) of thermally treated *Eucalyptus grandis* woods at
 246 different temperatures - 140, 180 and 220°C - as a function of the treatment atmosphere
 247 condition.

248

249

250

251

252

Figure 1 shows the increase in stability of the thermal degradation of the thermally treated wood in relation to the untreated wood (control). The samples submitted to the treatments at higher temperatures showed a mass loss with late onset and at lower rates, besides the increase of the residual mass in the thermogravimetric analysis. The effect of the

253 treatment atmosphere was less significant in the thermal degradation profiles, and its effect is
254 discussed later in Table 3.

255 The thermal degradation DTG profile of the control samples is divided into two
256 characteristic mass loss peaks. According to Bach *et al.* (2014), the first peak, which in the
257 present study was between 280 and 320°C, is attributed to thermal degradation of
258 hemicelluloses. The hemicelluloses are more reactive and susceptible to heat action due to
259 their amorphous structure, less compacted, with a low degree of polymerization and with high
260 amounts of hydroxyls Yang *et al.* (2007). The reduction and absence of this peak in the
261 thermally treated samples indicate, therefore, a lower content of these polysaccharides due to
262 the thermal degradation that occurred during the treatment.

263 The second mass loss peak, which in the present study was between 320 and 400°C, is
264 attributed to the thermal degradation of cellulose (Shen *et al.* 2010, Bach *et al.* 2014). The
265 peak mass loss for the cellulose is higher than the hemicelluloses, due to the greater amount
266 of energy required for the depolymerization of the cellulose chain and the breakdown of its
267 monomers. The increase of this peak in some thermally treated samples suggests a
268 concentration in the cellulose contents as a consequence of the degradation of hemicelluloses,
269 as discussed by Da Silva *et al.* (2016).

270 In the DTG graphs it was not possible to observe any mass loss peak that could be
271 related to lignin degradation. Lignins are macromolecules formed by aromatic units that are
272 characterised by high stability and thermal degradation, presenting low rates of mass loss
273 (Yang *et al.* 2007, Haykiri-Acma *et al.* 2010). According to Pereira *et al.* (2013), the final
274 residual mass presented by the TGA graphs consists mainly of residual lignin molecules. The
275 higher percentage values of residual mass above 400°C therefore indicate higher lignin
276 contents in the thermally treated samples than in the control sample.

277 Table 3 shows the mass loss percentage values obtained in each temperature range
278 during the thermogravimetric analysis. The highest variations were observed for the wood
279 samples submitted to the thermal treatment under nitrogen atmosphere. In this condition,
280 samples treated up to 180°C showed an increase in mass loss in the temperature range
281 between 200 and 300°C, followed by a reduction in the range between 300 and 400°C. For
282 the samples treated at 220°C, however, the trend showed a significant reversal, with reduction
283 of mass loss between 200 and 300°C and increase between 300 and 400°C.

284

285

286 **Table 3.** Mass loss (%), during the thermogravimetric analysis, as a function of the
 287 temperature ranges for the heat treatments and the control.

Treatment		Temperature range (°C)			Residual mass
		Up to 200	200–300	300–400	
Untreated		1.19	12.13	53.16	33.52
Vacuum	140°C	0.91	12.73	52.43	33.93
	180°C	1.33	17.53	46.37	34.77
	220°C	1.16	7.17	54.46	37.21
Nitrogen	140°C	1.04	15.60	48.08	35.28
	180°C	1.01	18.86	45.17	34.96
	220°C	1.06	5.91	54.58	38.44
Vacuum	140°C	0.84	13.33	52.15	33.68
+	180°C	1.18	15.01	47.61	36.19
Nitrogen	220°C	0.96	6.21	55.06	37.76

288

289 As discussed by Candelier *et al.* (2013), thermal treatments performed under vacuum
 290 and nitrogen conditions present significant differences. According to these authors, unlike the
 291 vacuum that facilitates the removal of the volatile organic compounds generated during the
 292 heat treatment, the nitrogen condition does not allow such an efficient elimination of these
 293 compounds, allowing them to remain for longer in contact with the wood. Some of these
 294 compounds have acidic pH, especially acetic acid, and they act as catalysts to accelerate the
 295 chemical modification reactions of polysaccharides, mainly hemicelluloses. With this, the
 296 nitrogen atmosphere usually presents greater chemical modifications of the wood compared to
 297 vacuum. Although there were no significant differences in our study, similar trends to that one
 298 were observed among the mean values.

299 Temperatures of 140 and 180°C are not sufficient to promote significant mass losses
 300 during heat treatment. However, they may initiate chemical modifications in the constituents
 301 of wood, especially hemicelluloses, due to thermal degradation. Thus, the fraction of
 302 hemicelluloses, with their altered chemical structure, tends to be more susceptible to heat
 303 exposure, increasing their mass loss between 200 and 300°C. As a consequence, the mass loss
 304 between 300 and 400°C tends to decrease. In addition, the milder heat treatments promote an
 305 increase in the crystallinity of the cellulose and its thermal stability (Esteves & Pereira 2009),
 306 also contributing to the lower mass loss in the mentioned temperature range.

307 The most intense heat treatment, applied at 220°C, was sufficient to degrade a
308 significant part of the hemicelluloses, thus concentrating the cellulose and lignin contents in
309 the final solid fraction, as discussed in Table 2 and Figure 1. With this, the wood samples
310 treated at that temperature showed lower mass loss between 200 and 300°C during the
311 thermogravimetric analysis. The loss of mass in the temperature range of 300 to 400°C and
312 the residual mass above 400°C presented higher values in relation to the control, due to the
313 higher concentrations of cellulose and lignin, respectively.

314

315 **4 CONCLUSIONS**

316

317 According to the results obtained in the present study, for thermally treated *Eucalyptus*
318 *grandis* wood up to 220°C under vacuum and / or nitrogen atmosphere, it can be concluded
319 that:

- 320 • The temperature has a direct effect on the heat treatment efficiency of this wood,
321 promoting gradual increases in mass loss, while the treatment atmosphere has a less
322 significant effect.
- 323 • The thermally treated wood presents significant changes in the proportions of the
324 chemical constituents, mainly due to the increase in temperature, such as lower
325 holocelluloses contents and higher lignin and extractive contents in relation to the
326 control samples.
- 327 • There was greater thermal stability, with a delay at the beginning of mass loss and an
328 increase in residual mass values during thermogravimetric analysis.
- 329 • For the eucalypt wood, the nitrogen atmosphere tends to present larger changes in the
330 chemical components when compared to the vacuum atmosphere as a function of the
331 temperature increase.

332 The results show that the heat treatment is capable of promoting significant changes in
333 the structure and chemical composition of *Eucalyptus grandis* wood and allows us to observe
334 the best temperature and atmosphere according to the level of chemical decomposition and
335 mass loss. It is interesting to seek methods that treat the wood causing the least damage
336 possible for its use. These changes are reflected in the other indices of material quality, so it is
337 necessary to know the treatment condition capable of promoting the most advantageous
338 increments to the purpose for which each type of treated wood is destined.

339

340 **References**

341

342 **Araújo, S.O.; Vital, B.R.; Mendoza, Z.M.S.H.; Vieira, T.A.; Carneiro, A.C.O. 2012.**
343 Properties of thermorectified wood of *Eucalyptus grandis* and *Eucalyptus*. *Scientia*
344 *Forestalis* 40 (95): 327-336.

345

346 **Araújo, S.O.; Vital, B.R.; Oliveira, B.; Carneiro, A.C.O.; Lourenço, A.; Pereira, H.**
347 **2016.** Physical and mechanical properties of heat treated wood from *Aspidosperma*
348 *populifolium*, *Dipteryx odorata* and *Mimosa scabrella*. *Maderas-Cienc Tecnol* 18 (1): 143-156.
349 DOI: <https://doi.org/10.4067/S0718-221X2016005000015>.

350

351 **Ayata, U.; Akcay, C.; Esteves, B. 2017.** Determination of decay resistance against *Pleurotus*
352 *ostreatus* and *Coniophora puteana* fungus of heat-treated scotch pine, oak and beech wood
353 species. *Maderas-Cienc Tecnol* 19(3):309-316. DOI: [https://doi.org/10.4067/S0718-](https://doi.org/10.4067/S0718-221X2017005000026)
354 [221X2017005000026](https://doi.org/10.4067/S0718-221X2017005000026).

355

356 **Bach, Q.; Tran, K.; Skreiberg, O.; Khalil, R.A.; Phan, A.N. 2014.** Effects of wet
357 torrefaction on reactivity and kinetics of wood under air combustion conditions. *Fuel* 137:
358 375-383. DOI: <https://doi.org/10.1016/j.fuel.2014.08.011>.

359

360 **Bourgois, J.; Guyonnet, R. 1988.** Characterization and analysis of torrefied wood. *Wood*
361 *Science Technology* 22 (2): 143-155. DOI: <https://doi.org/10.1007/BF00355850>.

362

363 **Brito, J.O.; Garcia, J.N.; Bortoletto Junior, G.; Pessoa, A.M.C.; Silva, P.H.M. 2006.** The
364 density and shrinkage behavior of *Eucalyptus grandis* wood submitted to different
365 temperatures of thermorectification. *Cerne* 12 (2): 182-188. URL:
366 <http://cerne.ufla.br/site/index.php/CERNE/article/view/413>.

367

368 **Candelier, K.; Dumarçay, S.; Pétrissans, A.; Desharnais, L.; Gérardin, P.; Pétrissans,**
369 **M. 2013.** Comparison of chemical composition and decay durability of heat treated wood
370 cured under different inert atmospheres: nitrogen or vacuum. *Polymer Degradation and*
371 *Stability* 98: 677-681. DOI: <https://doi.org/10.1016/j.polymdegradstab.2012.10.022>.

372

373 **Da Silva, C.M.S.; Carneiro, A.C.O.; Pereira, B.L.C.; Vital, B.R.; Alves, I.C.N.;**
374 **Magalhães, M.A. 2016.** Stability to thermal degradation and chemical composition of woody
375 biomass subjected to the torrefaction process. *European Journal of Wood and Wood Products*
376 74 (6): 845-850. DOI: <https://doi.org/10.1007/s00107-016-1060-z>.

377

378 **Esteves, B.M.; Domingos, I.L.; Pereira, H.M. 2008.** Pine wood modification by heat
379 treatment in air. *BioResources* 3 (1): 142-154.

380

381 **Esteves, B.M.; Pereira, H.M. 2009.** Wood modification by heat treatment: a review.
382 *BioResources* 4 (1): 370-404.

383

384 **Esteves, B.; Videira, R.; Pereira, H. 2011.** Chemistry and ecotoxicity of heat treated pine
385 wood extractives. *Wood Science Technology* 45 (6): 661-676. DOI:
386 <https://doi.org/10.1007/s00226-010-0356-0>.

387

- 388 **Fengel, D.; Wegener, G. 1989.** Wood: Chemistry, Ultrastructure, Reactions. Walter de
389 Gruyter, Berlin.
390
- 391 **Figueiró, C.G.; Vital, B.R.; Carneiro, A.C.O.; Silva, C.M.S.; Magalhães, M.A.; Fialho,**
392 **L.F. 2019.** Energy valorization of woody biomass by torrefaction treatment: a brazilian
393 experimental study. *Maderas-Cienc Tecnol* 21(3): 297-304. DOI: [https://](https://doi.org/10.4067/S0718-221X2019005000302)
394 doi.org/10.4067/S0718-221X2019005000302.
395
- 396 **Haykiri-Acma, H.; Yaman, S.; Kucukbayrak, S. 2010.** Comparison of the thermal
397 reactivities of isolated lignin and holocellulose during pyrolysis. *Fuel Processing Technology*
398 91: 759-764. DOI: <https://doi.org/10.1016/J.FUPROC.2010.02.009>.
399
- 400 **Inari, G.N.; Petrissans, M.; Gerardin, P. 2007.** Chemical reactivity of heat-treated wood.
401 *Wood Science Technology* 41: 157-168. DOI: <https://doi.org/10.1007/s00226-006-0092-7>.
402
- 403 **Martinka, J.; Hroncová, E.; Chrebet, T.; Balog, K. 2014.** The influence of spruce wood
404 heat treatment on its thermal stability and burning process. *European Journal of Wood and*
405 *Wood Products* 72: 477-486. DOI: <https://doi.org/10.1007/s00107-014-0805-9>.
406
- 407 **Moura, L.F.; Brito, J.O.; Silva Júnior, F.G. 2012.** Effect of thermal treatment on the
408 chemical characteristics of wood from *Eucalyptus grandis* W. Hill ex Maiden under different
409 atmospheric conditions. *Cerne* 18 (3): 449-455. DOI: [https://doi.org/10.1590/S0104-](https://doi.org/10.1590/S0104-77602012000300012)
410 [77602012000300012](https://doi.org/10.1590/S0104-77602012000300012).
411
- 412 **Olarescu, M.C.; Campean, M.; Ispas, M.; Cosereanu, C. 2013.** Effect of thermal treatment
413 on some properties of lime wood. *European Journal of Wood and Wood Products* 72: 559–
414 562. DOI: <https://doi.org/10.1007/s00107-014-0809-5>.
415
- 416 **Pereira, B.L.C.; Carneiro, A.C.O.; Carvalho, A.M.M.L.; Colodette, J.L.; Oliveira, A.C.;**
417 **Fontes, M.P.F. 2013.** Influence of chemical composition of Eucalyptus wood on gravimetric
418 yield and charcoal properties. *BioResources* 8 (3): 4574-4592.
419
- 420 **Prins, M.J.; Ptasinski, K.J.; Jansen, F.J.J.G. 2006.** Torrefaction of wood, part 2. Analysis
421 of products. *Journal of Analytical Applied Pyrolysis* 77: 35–40. DOI:
422 <https://doi.org/10.1016/j.jaap.2006.01.001>.
423
- 424 **Shen, D.K.; Gu, S.; Bridgwater, A.V. 2010.** The thermal performance of the
425 polysaccharides extracted from hardwood: Cellulose and hemicelluloses. *Carbohydrate*
426 *Polymers* 82: 39-45. DOI: <https://doi.org/10.1016/j.carbpol.2010.04.018>.
427
- 428 **Sjöström, E. 1981.** Wood polysaccharides, in Wood chemistry. Fundamentals and
429 applications. *Academic Press* 3: 49–67.
430
- 431 **TAPPI. 2002.** Technical Association of the Pulp and Paper Industry. TAPPI test methods T
432 204 om-88: solvent extractives of wood and pulp. In: TAPPI Standard Method. Atlanta, GA.
433
- 434 **Xing, D.; Li, J. 2010.** Effects of heat treatment on thermal decomposition and combustion
435 performance of *Larix* spp. wood. *BioResources* 9 (3): 4274-4287.
436

- 437 **Yang, H.; Yan, R.; Chen, H.; Lee, D.H.; Zheng, C. 2007.** Characteristics of hemicellulose,
438 cellulose and lignin pyrolysis. *Fuel* 86: 1781-1788. DOI:
439 <https://doi.org/10.1016/j.fuel.2006.12.013>.
440
- 441 **Yildiz, S.; Gezer, E.D.; Yildiz, U.C. 2006.** Mechanical and chemical behavior of spruce
442 wood modified by heat. *Building and Environment* 41 (12): 1762-1766. DOI: DOI:
443 <https://doi.org/10.1016/j.buildenv.2005.07.017>.
444
- 445 **Watanabe, T.; Ohnishi, J.; Yamasaki, Y.; Kaizu, S.; Koshijima, T. 1989.** Binding-site
446 analysis of the ether linkages between lignin and hemicelluloses in lignin-carbohydrate
447 complexes by DDQ-oxidation. *Agricultural and Biological Chemistry* 53: 2233–2252. DOI:
448 <https://doi.org/10.1080/00021369.1989.10869603>.
449
- 450 **Welzbacher, C.R.; Brischke, C.; Rapp, A.O. 2007.** Influence of treatment temperature and
451 duration on selected biological, mechanical, physical and optical properties of thermally
452 modified timber. *Wood Mater Science and Engineering* 2: 66-76. DOI:
453 <https://doi.org/10.1080/17480270701770606>.
454
- 455 **Wentzel, M.; Brischke, C.; Militz, H. 2019.** Dynamic and static mechanical properties of
456 *Eucalyptus nitens* thermally modified in an open and closed reactor system. *Maderas-Cienc*
457 *Tecnol* 21(2):141-152. DOI: <http://dx.doi.org/10.4067/S0718-221X2019005000201>.
458
- 459 **Zanuncio, A.J.V.; Nobre, J.R.C.; Motta, J.P.; Trugilho, P.F. 2014.** Chemistry and
460 colorimetry of thermorectified wood from *Eucalyptus grandis* W. Mill ex Maiden. *Revista*
461 *Árvore* 38 (4): 765-770.
462
463
464
465
466
467