Castor oil based polyurethane adhesive content on OSSB produced with soybean straw

Efeito do teor de adesivo poliuretano à base de óleo de mamona em painéis OSSB de palha de soja

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Abstract

riented Structural Straw Board (OSSB) panels are composites made from straw originally from agricultural residues and nonformaldehyde based adhesive, whose main application is for construction as a closing and ceiling material. The objective of this study was to evaluate the effect of the polyurethane adhesive content on physical and mechanical properties of OSSB produced with locally available SbS (soybean straw). Four castor oil based polyurethane ratios (6%, 9%, 12% and 15%, by mass of the SbS) were tested. OSSB was evaluated according to European standards for wood-based composites; also, a physical and chemical characterization of the SbS was performed. SbS has an irregular shape, which reduced the OSSB mechanical performance due to the low densification of the composite and the development of transversal cracks on the SbS after the OSSB manufacturing process. The static bending and thickness swelling performance of SbS OSSB was not improved by the increase of polyurethane adhesive content. SbS low ash content and neutral pH extractives indicates that probably its chemical properties should not affect the setting of the adhesive.

Keywords: Glycine max L. Agricultural residue. Structural composite.

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Resumo

Painéis OSSB (Oriented Structural Straw Board) são compósitos produzidos com palha proveniente de resíduos agrícolas e utilizando de adesivo sem formaldeído, cuja principal aplicação se volta para a construção civil como material de fechamento e forros. Este trabalho teve como objetivo avaliar o efeito do teor de poliuretano no adesivo sobre propriedades físicas e mecânicas do OSSB produzido com SbS (palha de soja) localmente disponível. Quatro composições de poliuretano à base de óleo de mamona (6%, 9%, 12% e 15%, em relação à massa de palha) foram testadas. Painéis OSSB foram avaliados de acordo com os padrões europeus para compósitos à base de madeira; também, uma caracterização física e química da SbS foi realizada. A SbS possui formato irregular, o que reduziu o desempenho mecânico do OSSB devido à baixa densificação do compósito e ao desenvolvimento de trincas transversais na SbS após o processo de fabricação do compósito. O desempenho em testes de flexão estática e inchamento de espessura do OSSB não foi melhorado pelo aumento do teor de adesivo de poliuretano. A SbS apresentou baixo teor de cinzas (menos que 4.3%) e pH neutro, isto indica que estas propriedades químicas provavelmente não afetaram na cura do adesivo.

Palavras-chave: Glycine max L. Resíduo agrícola. Compósito estrutural.

Introduction

According to the USDA (UNITED..., 2019), more than 360 million metric tons of soybean oilseeds were harvested worldwide in 2018/2019, and Brazil is the second main soybean producer in the world, after the United States. Lal (2005) and Krisnawati and Adie (2015) determined values ranging from 0.8 to 2.6 of soybean dry straw to grain ratio, demonstrating the large availability of this residue.

Agricultural residues are often considered a problem for farmers due to their low monetary return, but they can be viewed as an economic opportunity, especially for developing countries where they can have a positive impact (ADEL *et al.*, 2016). Some residues (*e.g.* wheat straw, rice straw and rice husk) receive interest due to their low cost, great availability, reduction of environmental pollution and large number of natural polymers in their composition (ARGENTO *et al.*, 2019; LIMA *et al.*, 2019).

Barbu, Reh and Çavdar (2017) indicate that the production of composites with agricultural residues is considered attractive, especially due to the increase in the price of wood as a raw material. According to Väisänen *et al.* (2016) it is required that these products do not present inferior performance in comparison to currently available wood-based composites, otherwise the selection of the agricultural residue as a raw material will not happen.

Studies applying SbS (soybean straw) residues in the production of composites have been developed (REINPRECHT *et al.*, 2015; SITZ; BAJWA, 2015; SHAO; YANG; JIANG, 2016; MARTELLI-TOSI *et al.*, 2017) and showed positive results in comparison to traditional materials (*e.g.* timber, plywood and fiberboards), although these composites are classified as non-structural according to EN 300 standard (EUROPEAN..., 2006).

It is also important to note that there have not been studies performed to evaluate the thermal conductivity of SbS based composites, even though this property is used to determinate heat transfer and insulation of building constructions, such as cladding, roughing and flooring (TRIPATHI; RICE, 2017; ÇAVUŞ *et al.*, 2019).

This paper aimed to analyze the feasibility of producing a structural composite applying locally available SbS, in order to increase its economic value.

OSSB is a commercially available composite with structural performance that uses rice or wheat straw and had its first manufacturing plant starting operations in China in 2009 (HAN *et al.*, 2012). The straw is split longitudinally prior to the production of the board due to the action of two grooved rollers with different tangential speed, so the binder is allowed to have contact with the inside of the split straw (BACH; DOMIER; HOLOWACH, 1999).

OSSB is composed of three layers of straw that are oriented perpendicular to each other, offering better dimensional stability and mechanical performance in both directions of the plane (MARRA, 1992); this same principle is applied to wood-based composites, such as plywood, CLT (Cross Laminated Timber) and OSB (Oriented Strand Board). Image analysis is used in order to evaluate the straw orientation and layers thickness, due to the need of better characterizing and understanding this composite structure.

Another feature of OSSB is the use of formaldehyde-free adhesive (*i.e.* p-MDI resin) in its composition, due to a better adhesion to straw in general (BACH; DOMIER; HOLOWACH, 1999; HAN *et al.*, 2012; CHENG; HAN; FANG, 2013). Also, the limits of formaldehyde release of adhesives applied to lignocellulosic composites are under focus due to society's demand for more ecological processes and because its contribution to poor indoor air quality (STEFANOWSKI; CURLING; ORMONDROYD, 2017; WANG *et al.*, 2018).

Polyurethane is another type of formaldehyde-free adhesive (polyurethane based) that has a wide range of applications in industry (TANG *et al.*, 2008). This binder has been substantially investigated as a replacement of conventional adhesives in wood-based and other lignocellulosic composites (MERLINI; SOLDI; BARRA, 2011; FIORELLI *et al.*, 2012, 2013; BERTOLINI *et al.*, 2013; VALARELLI *et al.*, 2014; CHEN; TAI, 2018; NASCIMENTO *et al.*, 2018; VARANDA *et al.*, 2018; FERRO *et al.*, 2019; MACEDO *et al.*, 2019; SHIROSAKI *et al.*, 2019). The use of castor oil polyurethane adhesive in the composition SbS OSSB in this paper is justified due the abundance of the *Ricinus communis* L. plant in Brazil, whose seeds are used for the production of the resin (CANGEMI *et al.*, 2006).

The goal of this study was to evaluate the potential of using SbS and polyurethane adhesive for producing composites with structural performance.

Methods

SbS was collected two days after the soybean harvest in the region of Itapeva-SP, in Brazil. The soybean clone used was M5917IPRO in a clayey soil with 14 plants per linear meter. SbS was physically and chemically characterized prior to the production of the strawboard. A commercially available polyurethane adhesive (KEHL®) with 100% solid content was used, which was obtained by reaction of polymeric isocyanate with castor oil as a polyol in the mass ratio of 50:50.

SbS physical characterization

Straw density was determined according to the method described by Lam *et al.* (2008). SbS was considered a hollow cylinder and had its mass measured with a laboratory scale and dimensions – diameter, wall thickness and length – measured with a caliper in order to determine the straw volume and density, as shown on Equation 1:

$$S. D. = \frac{Mi}{(d \cdot t - t^2) \cdot \pi \cdot l} \cdot 10^6$$
 Eq. 1

Where:

 $S.D. = \text{straw bulk-density, in kg·m}^{-3}$;

Mi =mass of the internode section, in g;

d = internode section diameter, in mm;

t = wall thickness of the internode section, in mm; and

l =length of the internode, in mm.

SbS had its mass determined in a laboratory scale before and after being dried in an air-forced oven at 103 °C for 24 hrs. Straw moisture content (S.M.C.) was determined using the initial mass (M_1) and the dried mass (M_2), as described at Equation 2:

$$S.M.C. = \frac{(M_1 - M_2)}{M_2} \cdot 100$$
 Eq. 2

Where:

S.M.C. = straw moisture content, in %;

 M_I = initial mass, in g; and

 M_2 = dried mass, in g.

SbS is composed of the plant's stem and petioles, each of them presenting different dimensions. A comparison of number of internodes, average diameter, average wall thickness, average internode length and total length for both parts of the soybean plant was carried out.

SbS chemical characterization

SbS was initially reduced to particles using a hammer mill (TE-650, TECNAL. Brazil) until reaching dimensions of 40- to 60- mesh. Total Extractive Content (*T.E.C.*) and Hot Water Extractive Content (*H.W.E.C.*) were determined according to TAPPI 204 method. *T.E.C.* was determined as the difference between masses of the particles before (*mb*) and after (*ma*) a three stage extraction process, as shown in Equation 3:

$$T.E.C. = \frac{(mb - ma)}{ma} \cdot 100$$
 Eq. 3

Where:

T.E.C. = total extractive content, in %;

mb = mass before extraction, in g; and

ma = mass after extraction, in g.

The first stage of extraction used 1/3 ethanol and 2/3 benzene solution for 5 h, the second stage used ethanol solution for 4 h, and in the last stage the extraction used boiling water for 1 h. H.W.E.C. followed the same

principle, except that there was only one stage of extraction in boiling water for 3 h. The resulting solution of the *H.W.E.C.* test had its pH measured.

Ash Content (A.C.) of SbS was determined according to EN 1755-01 (AMERICAN..., 2015) method. Two grams samples were heated in a laboratory furnace at 575 °C for 3 h. A.C. was considered as the ratio between the sample mass after three hours at the furnace (mf) and initial mass (mi), as described in Equation 4·

$$A. C. = \frac{mf}{mi} \cdot 100$$
 Eq. 4

Where:

A.C. = ash content, in %;

mf = final mass, in g; and

mi = initial mass, in g.

OSSB production process

OSSB was produced in laboratory scale according to the method described by Bach, Domier and Holowach (1999) and Shirosaki *et al.* (2019). SbS was manually split along its main axis to expose its inner regions, and then was dried in a laboratory oven at 103 °C for 24 h before the production of the boards. Dried split SbS was mixed with castor oil based polyurethane adhesive and the mattress was formed inside a retaining square box with 0.42 m side.

The mattress was pressed in room temperature for 10 min with pressure of 0.4 MPa. Then the board was pressed in a heated hydraulic press (PHH 80T, Hidralmac. Brazil) at 100 °C in three cycles of 180 s (there was a pressure release of 3 s between cycles) with pressure of 4 MPa, following the Ferreira, Silva and Campos (2017) method. The board had a nominal thickness of 15 mm and target density of 650 kg·m⁻³ and was composed of three perpendicular layers in the proportion of 30 : 40 : 30 by mass. Each layer had the straw oriented longitudinally in a preferential direction. Four adhesive contents (ranging from 6% to 15% in relation to the dried straw mass) were tested, in order to determine its optimal amount in the composition of OSSB.

OSSB physical characterization

The determination of density of the boards was based on EN 323 standard (EUROPEAN ...1993a). Specimens (with nominal dimensions of $0.05 \text{ m} \times 0.05 \text{ m} \times 0.015 \text{ m}$) had their length, width and thickness measured with a caliper and mass measured with a laboratory scale. Density was calculated according to Equation 5:

$$D = \frac{m}{l \cdot w \cdot t} \cdot 10^6$$
 Eq. 5

Where:

 $D = \text{density}, \text{ in kg·m}^{-3};$

m =mass of the specimens, in g;

l = length of the specimens, in mm;

w =width of the specimens, in mm; and

t = thickness of the specimens, in mm.

Moisture content (M.C.) determination method was based on EN 322 standard (EUROPEAN..., 1993b). Specimens with at least 0.02 kg had their initial mass (mi) and final mass (mf) measured with a precision scale. The final mass was defined as the mass of the specimen dried in a laboratory oven at 103 °C until reaching a constant value – less than 0.1% variation between consecutive measurements taken every six hours. The moisture content of the boards was defined according to Equation 6:

$$M.C. = \frac{(mi - mf)}{mf} \cdot 100$$
 Eq. 6

Where:

M.C. = moisture content, in %;

mi = initial mass, in g; and

mf = dried mass, in g.

Thickness Swelling (*T.S.*) after immersion in water for 2 and 24 h were determined according to EN 317 standard (EUROPEAN..., 1993c). Square specimens (0.05 m x 0.05 m) had their thickness measured with a caliper before (*ti*) and after (*tf*) immersion in water at room temperature and neutral pH for 2 h and 24 h, as shown in Equation 7:

$$T.S. = \frac{(tf - ti)}{ti} \cdot 100$$
 Eq. 7

Where:

T.S. = straw moisture content, in %;

ti = initial thickness, in mm; and

tf = final thickness, in mm.

The optical micrographs of the specimens were taken by a Lupe (M80, Leica. Brazil) with a coupled camera (DFC295, Leica. Brazil) connected to a computer using Leica QWin Standard V 3.5.1 software (2008) for image processing. Conductivity tests were performed with a Thermal Properties Analyzer (KD2 Pro, Decagon Devises, Inc.) in an acclimatized chamber (SM-3.5S, THERMOTRON) at 20 °C.

OSSB mechanical characterization

Mechanical tests were performed using a universal testing machine (DL 30000, EMIC. Brazil). The three-point bending test was based on EN 310 standard (EUROPEAN..., 1993d), where specimens (350 mm x 50 mm) were tested using a span of 300 mm with a crosshead speed of 7 mm/min. *MOE* and *MOR* were determined according to Equations 8 and 9, respectively.

$$MOE = \frac{l^3 \cdot (P_2 - P_1)}{4 \cdot w \cdot t^3 \cdot (a_2 - a_1)}$$
 Eq. 8

Where:

MOE = Modulus of elasticity, in MPa;

l = length of span, in mm;

 $(P_2 - P_I)$ = is the increment of load on the load-deflection curve, where P_2 was approximately 40% and P_1 10% of the maximum load, in N;

w = width of the specimens, in mm;

t = thickness of the specimens, in mm; and

 $(a_2 - a_I)$ = the increment in deflection due to (P2 - PI) in the load-deflection curve.

$$MOR = \frac{3 \cdot P_{max} \cdot l}{2 \cdot w \cdot t^2}$$
 Eq. 9

Where:

MOR =modulus of rupture, in MPa;

 $P_{max} = \text{maximum load, in N};$

l = length of the span, in mm;

w = width of the specimens, in mm; and

t = thickness of the specimens, in mm.

Internal Bonding (*IB*) test was performed according to EN 319 standard (EUROPEAN..., 1993e). Square specimens (50 mm x 50 mm) were bonded to wood braces using fast bonding contact adhesive. Crosshead speed of 0.5 mm/min was applied until failure of the samples, and *IB* was determined according to Equation 10.

$$I.B. = \frac{P_{max}}{l \cdot w}$$
 Eq. 10

Where:

IB = Internal bond, in MPa;

 $P_{max} = \text{maximum load, in N};$

w = width of the specimens, in mm; and

l =length of the specimens, in mm.

Resistance to screw withdrawal was performed according to EN 320 standard (EUROPEAN..., 2011), as seen in Figure 1. Specimens (75 mm x 75 mm) had their edge and face screw withdrawal strength measured by the maximum tension applied before the failure of the sample.

Statistical analysis

The average interval of confidence for the physical and chemical evaluated properties of the SbS was determined, tested with p < 0.05. The factor studied was the polyurethane adhesive content in OSSB production, totaling four treatments with six replications for each test. The Tukey test was performed by ANOVA analysis, tested with p < 0.05. The software used to interpret the results was R version 3.2.3.

Results and discussions

SbS physical and chemical properties

The physical and chemical properties determined for SbS are shown in Table 1. SbS had intermediate values of density in comparison to wheat straw, which range from 395 to 781 kg·m⁻³ (LAM *et al.*, 2008; TUMULURU *et al.*, 2014; SILVA *et al.*, 2019), and higher values when compared to rice straw, which range from 162 to 194 kg·m⁻³ (ZHANG; GHALY; LI, 2012). The SbS density was measured with average moisture content of 21.49% (as seen on Table 1), considered high enough to start its degradation (GOODHEW; GRIFFITHS; WOOLLEY, 2004; BOUASKER *et al.*, 2014). Although SbS degradation analysis was not the focus of this paper, mold formation was observed after a couple of weeks when the residue was collected on the day of the soybean harvest, justifying the need of a preliminary drying.

It was determined that more than 90% of SbS extractives were soluble in hot water and did not alter their pH. The *T.E.C.* amount determined for Sb was higher than those reported for rice straw (10%) and wheat straw (7%) (WORASUWANNARAK; SONOBE; TANTHAPANICHAKOON, 2007; SHMULSKY; JONES, 2019). The SbS *A.C.* shown in Table 1 is in the range of values (2% - 5%) determined by Reddy and Yang (2009); in contrast to *T.E.C.*, wheat straw and rice straw had higher *A.C.* than SbS, 6.49% and 9.2%, respectively (MOUSAVI *et al.*, 2013).

Figure 1 - Edge and face screw withdrawal resistance test

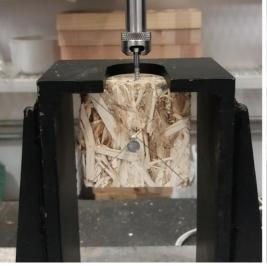




Table 1 - Physical and chemical properties of SbS

Properties	Average	Standard Deviation	Coefficient of Variation	Average Confidence Interval*
S.D.** (kg·m ⁻³)	529	65	12.37	505 - 553
S.M.C. (%)	21.49	3.94	18.35	20.01 - 22.96
T.E.C. (%)	12.59	0.39	3.12	11.61 - 13.57
H.W.E.C. (%)	11.36	0.22	1.92	10.82 - 11.90
pН	6.96	0.07	1.04	6.78 - 7.14
A.C. (%)	3.73	0.32	8.51	3.34 - 4.13

Note: *P < 0.05.

**S.D.: Straw Density;

S.M.C.: Straw Moisture Content; T.E.C.: Total Extractive Content;

H.W.E.C.: Hot Water Extractive Content; and

A.C.: Ash Content.

High A.C. usually changes the pH of the raw material and accelerates the setting of the adhesive, impairing the composite mechanical performance due the reduction of adhesion between the reinforcement and the matrix (IWAKIRI, 2005). This was not likely to happen since SbS had low A.C., compared to rice and wheat straw, and neutral pH as shown in Table 1, indicating that its chemical properties probably had no negative effect on the OSSB mechanical performance.

The average geometrical dimensions of the soybean stem and petioles are shown in Table 2, as well as the coefficient of variation and the minimum and maximum measured values. It was observed that the soybean stem and petioles differed in almost every parameter measured, except for the total available length. The coefficients of variation were high, indicating a low performance of the adopted procedure.

The number of internodes represents the amount of divisions on SbS that are expected to break during the straw processing for OSSB production. Bach, Domier and Holowach (1999) and Han *et al.* (2012) describe that a raw material with the least number of internodes (*i.e.* longer internode length) and a controlled moisture content are desired in order to reduce the generation of fines during the manufacturing process. The effect of these internode divisions is shown in Figure 2 as an irregularity on the OSSB surface, which could difficult or even prevent the successful application of coatings on the board.

Nevertheless, the wide range of results indicates the necessity of a different industrial process than the one applied to wheat straw for the production of OSSB. This can be inferred, as the minimum value obtained of the external diameter is smaller than the maximum value of the wall thickness, due to the variation of the dimensions of the plant along its height. In addition, some plants do not show petioles on harvest season, which are represented by the "zeros" on the number of internodes and on the total available length shown in Table 2.

SbS OSSB physical properties

SbS OSSB physical properties (*i.e.* density, moisture content and thickness swelling) are shown in Table 3. OSSB density reached 631 kg·m⁻³ in composites produced with 15% resin content, close to the results reported by Cheng, Han and Fang (2013) and Sitz and Bajwa (2015) for OSSB and MDF (Medium Density Fiberboard) produced with wheat straw and pMDI resin, respectively. The OSSB density increased with higher amounts of polyurethane adhesive content, which can also be observed in Figure 3, where the composition of 15% shows SbS to be more compacted.

Bertolini *et al.* (2013) and Varanda *et al.* (2018) studied wood composites produced with polyurethane obtaining higher densities (from 800 to 1100 kg·m⁻³). On the other hand, Valarelli *et al.* (2014) achieved close values to the 15% polyurethane composition SbS OSSB density in the bamboo particleboard while Chen and Tai (2018) obtained inferior values in bamboo charcoal composites produced with PU adhesive.

The density of the lignocellulosic material used in PU adhesive matrix affects the final density of the composite, so it was expected that SbS OSSBs densities were at least the same of the SbS, but only the 12% and 15% polyurethane adhesive content boards were. This was due to the empty spaces caused by the geometry of SbS and the lower compaction of the 6% and 9% polyurethane adhesive content treatments, as seen on Figure 2 and Figure 3, respectively.

Table 2 - Dimensions of soybean stem and petioles

Properties		Stem	Petioles
Number of Internodes (units)	Average	15a*	5b
	C.V.**	14.67	93.51
	Minimum	10	0
	Maximum	19	17
Average Diameter (10 ⁻³ m)	Average	4.47a	2.26b
	C.V.	45.01	37.87
	Minimum	0.34	0.52
	Maximum	10.37	5.59
Average Wall Thickness (10 ⁻³ m)	Average	1.14a	0.62b
	C.V.	58.92	45.79
	Minimum	0.22	0.18
	Maximum	4.08	1.66
Average Internode Length (10 ⁻³ m)	Average	4.8a	12.7b
	C.V.	36.31	65.92
	Minimum	1.0	1.0
	Maximum	11.5	32.0
Total Available Length (10 ⁻² m)	Average	76.6a	69.8a
-	C.V.	11.31	78.26
	Minimum	56.0	0.0
	Maximum	91.0	202.0

Note: *P < 0.05.

**C.V.: Coefficient of Variation (%).

Figure 2 - Surface imperfection in OSSB due to SbS internode division (magnification of 100X)



Table 3 - Physical properties of SbS OSSB according to polyurethane adhesive content

Properties	T6*	Т9	T12	T15
D (kg·m ⁻³)	459a**	494b	559c	631d
<i>M.C.</i> (%)	9.6a	9.5a	8.8b	8.4c
T.S. 2 h (%)	72.8a	52.0b	40.6bc	27.6c
T.S. 24 h (%)	92.7a	63.0b	62.7b	51.0b

Nota: *T6: 6% adhesive content; T9: 9% adhesive content;

T12: 12% adhesive content;

T15: 15% adhesive content;

D: Density;

M.C.: Moisture Content; and T.S.: Thickness Swelling.

**The same letter in the same line signifies that there is not a significant difference among the average values at 5% of probability.

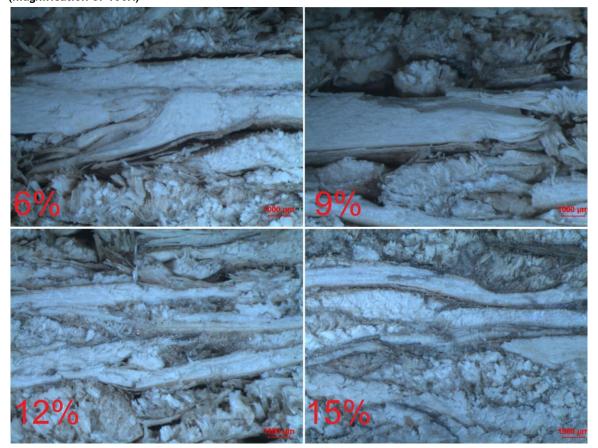


Figure 3 - SbS OSSB edge optical microscopies for the four percentages of polyurethane tested (magnification of 100X)

The *M.C.* determined for all tested OSSB fullfiled the requirements of EN 300 standard (EUROPEAN..., 2006), which must range from 2% to 12%. The values of *M.C.* were slightly reduced by the increase of PU adhesive content showing a decrease in water affinity, also observed in the results of *T.S.* for 2 h (see Table 3). Nevertheless, over 50% of *T.S.* 24h was achieved in the first 2 hrs of water immersion and more than 9% of PU content did not reduce *T.S.* 24 hrs significantly.

The maximum value allowed for *T.S.* 24 hrs according to EN 300 standard (EUROPEAN..., 2006) is 20%, for load bearing boards on dry conditions, less than half the average values obtained for SbS OSSB. *T.S.* 24h for wheat straw composites – OSSB and MDF – with pMDI binder ranged from 9.2% to 20.6% (CHENG; HAN; FAN, 2013; SITZ; BAJWA, 2015); and lignocellulosic particleboards with PU adhesive *T.S.* 24h ranged from 8.17% to 22.69% (FIORELLI *et al.*, 2012, 2013; VALARELLI *et al.*, 2014). The low performance of SbS OSSB is justified due the low adhesion of SbS with PU adhesive, as shown on Table 4 for *I.B.* tests results.

Specimens showed a thermal conductivity ranging from 0.193 to 0.202 W \cdot m⁻¹ · K⁻¹, which characterizes this material as a thermal insulator (TRIPATHI; RICE, 2017). Çavuş *et al.* (2019) determined values between 0.090 and 0.197 W \cdot m⁻¹ · K⁻¹ in their study with 31 different wood species and Binici, Aksogan and Demirhan (2016) found values between 0.075 and 0.1588 W \cdot m⁻¹ · K⁻¹ for corn straw boards. The conductivity of SbS OSSB is slightly higher due to its high content of SbS extractives, which probably increased the thermal conductivity despite the low density of the SbS and the composite.

SbS OSSB mechanical properties

The static bending *MOE* and *MOR* in both directions of the board, internal bonding and screw withdraw tests results are shown on Table 4. SbS OSSB *MOE* performance did not increase for PU content higher than 9% of binder was used in its composition and *MOR* reached its maximum value with 12% PU, similar to *T.S.* 24 hr results. EN 300 standard (EUROPEAN..., 2006) determines *MOE* values of at least 3500 MPa

and 1400 MPa and MOR values of 18 MPa and 9 MPa in the major and minor direction of load bearing boards in dry conditions, respectively.

OSSB produced with wheat straw and pMDI resin showed *MOE* of 4000 MPa and *MOR* of 20 MPa on the major axis, while SbS MDF reached *MOE* values close to 1420 MPa and *MOR* of 7.84 MPa as seen on SbS OSSB (CHENG; HAN; FANG, 2013; SITZ; BAJWA, 2015). The low static bending performance of OSSB is due the presence of transversal cracks on SbS (see Figure 4) which reduced the *MOE* and *MOR* values. These fragile cracks were developed because SbS was compressed to form the OSSB, and due to its irregular shape (as showed on the SbS physical and chemical properties subsection) the SbS was bent and twisted (see Figure 3) in the manufacturing process.

The effect of adhesive content of SbS OSSB was clearly seen on *I.B.* test results, as the 15% PU content composition performance was significantly higher than the other compositions tested, showing values close to those required for the EN 300 standard (EUROPEAN..., 2006) for load bearing boards in dry conditions. The *I.B.* performance obtained is also close to other straw based composites produced with 3% to 4% pMDI binder and 14% urea formaldehyde binder (CHENG; HAN; FAN, 2013; BEKHTA; KORKUT; HIZIROGLU, 2013; SITZ; BAJWA, 2015), but inferior to PU bonded composites (FIORELLI *et al.*, 2012 and 2013). The rupture of *I.B.* specimens occurred on the composite interface (see Figure 5), which created an irregular separation plane due to the heterogeneous dimensions of SbS (described in the SbS physical and chemical properties subsection).

Table 4 - Mechanical properties of SbS OSSB according to polyurethane adhesive content

Properties	T6*	Т9	T12	T15
MOE major (MPa)	683a**	1522b	1972b	1943b
MOE minor (MPa)	291a	537ab	498ab	626b
MOR major (MPa)	3.68a	6.95a	12.73b	14.48b
MOR minor (MPa)	2.37a	4.86b	5.12b	3.58ab
<i>I.B.</i> (MPa)	0.01a	0.04a	0.07a	0.30b
S.W. on edge (N)	200a	340ab	580bc	850c
S.W. on face (N)	490a	500a	730a	1190b

Nota: *T6: 6% adhesive content;

T9: 9% adhesive content;

T12: 12% adhesive content; T15: 15% adhesive content;

MOE: Modulus of Elasticity;

MOR: Modulus of Rupture;

I.B.: Internal Bonding; and

S.W.: Screw Withdraw.

**The same letter in the same line signifies that there is not a significant difference among the average values at 5% of probability.

Figure 4 - Transversal crack on SbS showed after the OSSB production (magnification of 100X)



Figure 5 - Internal bond specimen separated after its failure



OSSB edge and face *S.W.* strength improved with the increase of PU in its composition reaching values close to the commercially available OSSB (NOVOFIBRE, 2019), although there are no minimum standard requirements for this evaluated property. Sitz and Bajwa (2015) determined the face *S.W.* strength of 164 N for SbS MDF, which is inferior to the values obtained in this paper.

Conclusions

SbS has heterogeneous dimensions according to its measured parameters, which were: internode length, external diameter and wall thickness. 6% and 9% castor oil based polyurethane adhesive content OSSB had lower density than the SbS itself due to the raw material geometry. On the other hand, higher binder content boards (12% and 15%) showed a better densification, but developed transversal cracks on SbS due to its irregular shape and the OSSB manufacturing process impairing the boards' mechanical performance.

Some properties of SbS OSSB were clearly affected by the PU adhesive content, such as density, moisture content, internal bonding and screw withdrawal strength, but static bending *MOE* and *MOR* and thickness swelling did not improve with higher adhesive content. High SbS moisture content indicates the necessity of its drying before the manufacturing process. Due to the neutral extractives' pH and the lower ash content it can be concluded that the chemical properties of SbS did not affect the setting of the PU adhesive.

It was verified that the boards exhibited good performance as an insulating material, but not as a structural element. Therefore, they can be applied in construction as vertical or horizontal closing elements.

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