

EFFECT OF STYRENE MALEIC ANHYDRIDE ON PHYSICAL AND MECHANICAL PROPERTIES OF RECYCLED POLYSTYRENE WOOD FLOUR COMPOSITES

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ABSTRACT

In this work, the influence of three types of styrene maleic anhydride (SMA) oligomers on the adhesion of polystyrene wood plastic composites (WPC) was investigated. The composites were processed on a twin-screw co-rotating extruder below 200°C using 20 wt% of wood flour. The styrene maleic anhydride with different content of maleic anhydride groups, 30%, 25% and 20% (w/w) and levels of 1, 2 and 4% of coupling agents (styrene maleic anhydride 2000, styrene maleic anhydride 3000 and styrene maleic anhydride EF40) in the composite formulations were tested. Mechanical, physical and morphologic properties were evaluated. The styrene maleic anhydride improves the compatibility of hydrophilic wood flour with hydrophobic polystyrene matrix. It has been observed that the addition of styrene maleic anhydride increased the wood plastic composites mechanical properties with the incorporation of 2% wt of styrene maleic anhydride 2000. The mechanical properties showed to be dependent on content of maleic anhydride in the coupling agent. Treated and non-treated wood plastic composites showed similar density values, but the void content was reduced for treated composites. Scanning electron microscopy reveal the better adhesion between polymer and matrix when coupling agent were used.

Keywords: Coupling agent, injection moulding, mechanical properties, *Pinus elliottii*, polystyrene, wood plastic composites.

INTRODUCTION

The utilization of biomass for processing novel composites has attracted growing interest because of this eco-friendly and renewable nature (Sliwa *et al.* 2012, Nafchi *et al.* 2015). Materials reinforced with wood flour and others natural fibres increased considerably in recent years (Ornaghi Jr. *et al.* 2014). Wood flour is easily available, light and cheap, and it can be added to commodity matrices in large quantities thus offering economically advantageous solutions (Poletto *et al.* 2014, Khonsari *et al.* 2015). The main application areas of wood plastic composites (WPC) are the automotive and building industries (Nafchi *et al.* 2015) in which they are used in structural applications as door panels, decking, furniture, window parts, etc. However, in wood industry a large amount of wood waste is generated at different stages of the wood processing and such waste is mainly destined for landfill. Industrial region of Caxias do Sul, situated in northeast region of Rio Grande do Sul/Brazil, generates higher quantities of industrial and urban wastes. The furniture industry generates approximately 5500 m³/month of *Pinus elliottii* sawdust. On the other hand, plastic waste is one the major components of global municipal solid waste and, as such, it presents a promising raw materials source for WPC, especially because of large volume (Cui *et al.* 2008). Expanded polystyrene (EPS) is commonly used

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for packing materials, and after use EPS usually ends up in landfills or is incinerated. The estimate for generation of waste EPS in Caxias do Sul is about 1240 m³/month. The EPS wastes are deposited in the city landfill reducing drastically the area of the landfill. Thus, increased use of recycled plastics offers the prospect of lessening waste disposals and reducing the products costs (Adhikary *et al.* 2008). Hence, the development of new value added WPC products using wood waste and recycled plastic is assuming greater importance in environmental conservation.

One of the main disadvantages of WPC is the low compatibility between the hydrophilic wood filler and hydrophobic polymer matrix (El-Sabbagh 2014, Kim *et al.* 2007). Thus, two ways of improvement of the interface properties are generally used surface treatment of the reinforced fibres in order to increase their compatibility with the surrounding polymer, or modification of the matrix (El-Sabbagh 2014). The introduction of functional groups inside the polymer chains of the matrix aims to create chemical or physical interactions with the reinforcing fibres, and is a solution which leads to excellent results in practice. The maleic anhydride introduced by radical copolymerization in styrene constitutes one of the most efficient modifications agents used to functionalize the styrenic polymers (Devaux *et al.* 2002). Then, high molecular weight copolymers based on polystyrene and maleic anhydride may act as coupling agents in polystyrene composites reinforced by natural fibers. Sommerhuber *et al.* (2016) developed WPC based on recycled polystyrene reinforced with wood flour and evaluated the effect of 3% of SMA on the composite mechanical properties. The authors observed a significantly improve in tensile and flexural strength properties attributed to the better adhesion between wood and polymer matrix caused by SMA. Poletto *et al.* (2015) used 2% of SMA as coupling agent in polystyrene cellulose fiber composites. The addition of coupling agent substantially improves the mechanical and dynamic mechanical properties evaluated.

Therefore, the main goal of this study was to investigate the effect of three different styrene-co-maleic anhydride (SMA) oligomers in the adhesion of polystyrene composites reinforced with wood flour wastes. The influence of the maleic anhydride concentration in the copolymers on the mechanical, physical and morphologic properties of the WPC was evaluated.

MATERIALS AND METHODS

Materials

EPS wastes were obtained from a sorting unit called Associação de Recicladores Serrano, Caxias do Sul, Brazil, and it had a melt flow index (MFI) of 20g/10min (200°C/5 kg). Wood flour of *Pinus elliottii* (PIE) was obtained from Madarco Co., Caxias do Sul, Brazil. The three types of SMA used as a coupling agent were supplied by Sartomer Company, Exton/USA. SMA 2000 contains 30% w/w of maleic anhydride groups and weight average molecular weight (M_w) of 7500 g/mol, SMA 3000 with 25% w/w of maleic anhydride groups and M_w of 9500 g/mol and SMA EF40 contains 20% w/w of maleic anhydride groups and M_w of 10500 g/mol. The chemical structures of SMA used were showed in Figure 1. SMA was incorporated at 1, 2 and 4% in weight of coupling agents (SMA 2000, SMA 3000 and SMA EF40) in the WPC.

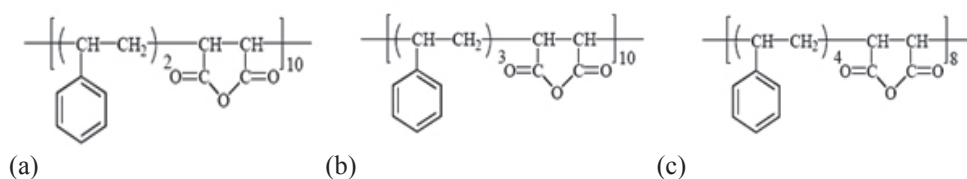


Figure 1. Chemical structure of SMA used as coupling agents, SMA 2000 (a), SMA 3000 (b), and SMA EF40 (c).

Composite preparation

The wood flour was dried in an oven at 105°C for 24 h. WPC samples with 20 w/w% of wood flour and with and without 1, 2, and 4 wt% of poly(styrene-co-maleic anhydride) oligomer were processed in a co-rotating twin-screw extruder COR-20-32-LAB MH Equipment-Brazil at 200 rpm, with L/D ratio of 32 into pellets. The nine barrel temperature zones were controlled at between 160°C and 190°C. Specimens for mechanical and physical tests were injection molded in a HIMACO LH 150-80 equipment at a barrel temperature of 180°C and mold temperature of $40 \pm 2^\circ\text{C}$, with screw speed of 100 rpm and pressure of 650 bar.

Wood flour particle size distribution

The wood flour particle size distribution was determined based on ASTM D1921 - Method A. A sample of 50g of wood flour previously dried was placed on a mechanical sieving device with selected sieves (28, 35, 48, 65, 100, 150, 200, 325 mesh). The measurements were carried out in triplicate.

Mechanical properties

Flexural tests were performed according to ASTM D790 at a cross-head speed of $1,5 \text{ mm}\cdot\text{min}^{-1}$ using EMIC DL 3000 testing machine. Izod impact strength was measured with a CEAST Resil 25 pendulum using unnotched specimens according to ASTM D256. Each test value was obtained as the average of at least five independent measurements.

Physical properties

Density for polystyrene and WPC were determined according ASTM D792. Each density value was obtained as the average of at least three independent measurements.

Water absorption test

For water absorption, rectangular specimens having dimensions of 100 mm x 13 mm x 3,5 mm were used. The specimens were dried in an oven at 80°C for 24 h, cooled in a desiccators using silica gel and immediately weighed. The water absorption test were carried out in triplicate immersing the specimens for 2 h, 24 h and 48 h in distilled water at 23°C. After immersion, the excess water on the surface of the specimens was wiped up using a piece of soft cloth and the final weights of the specimens were then taken. From the difference of weights of the specimens percentage of water absorption was calculated.

Morphological study

Studies on the morphology of wood flour and composites samples were carried out using a SHIMADZU Superscan SS-550, scanning electron microscope (SEM). The cryo fracture surface specimens were sputter-coated with gold before the analysis in order to eliminate electron charging.

Statistical analysis

The statistical analysis of variance for composite mechanical properties has been carried out using commercial software (EXCEL). A one-way analysis of variance and t-tests were used to evaluate the statistical difference among groups. Values of $p < 0,05$ were considered significant.

RESULTS AND DISCUSSIONS

Wood flour characterization

Fiber size and shape are among the most important factors related to composite materials (Bledzki *et al.* 2010). The effective surface area, which influence on mechanical properties inversely depends on particle size and shape (Bledzki *et al.* 2010, Poletto *et al.* 2012). Figure 2 shows the particle size and shape of the wood flour used in the WPC formulations. The particle size distribution is showed in the inset of Figure 2(a). It was observed that approximately 80% of all fibers were distributed in the range of 40-150 μm , although the distribution varied. Short and tiny fibers are preferable for the development of WPC formulations, since these provide a higher specific surface area and greater surface area of contact with the polymer matrix which usually produces better fiber-matrix adhesion when the coupling agent is used and an increase in the mechanical properties are normally observed (Gallagher *et al.* 2013).

The fiber surface morphology plays a vital role in the case of composite materials. The external surface features of the wood flour can be observed in Figure 2(b). The wood fiber surface is rough, which can provide mechanical anchoring to the polymer matrix and thus lead to improved mechanical properties compared to particles with smooth surfaces.

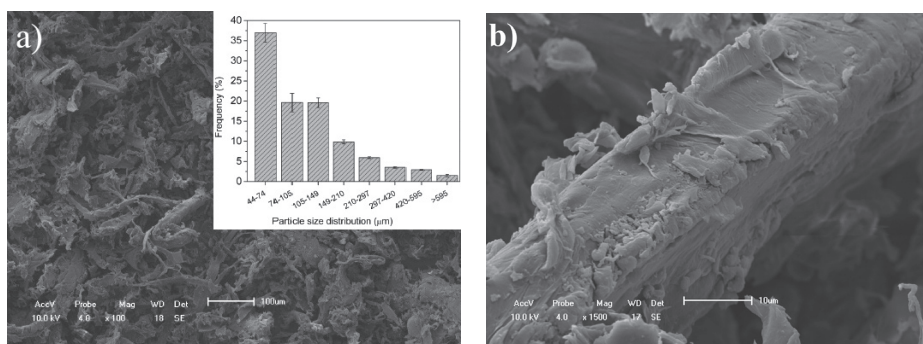


Figure 2. (a) SEM micrograph with magnification of 100x of wood flour and a (b) SEM micrograph detail (magnification 1500x) of wood fiber surface morphology.

Mechanical Properties

Table 1 showed the flexural strength and flexural modulus for the WPC with different SMA oligomers studied. The quantities of styrene-co-maleic anhydride necessary to obtain a maximum resistance in flexural strength are about 1-2%. The maximum value was obtained with 2% of the SMA 2000, as can be observed in Table 1. This coupling agent contains 30% of maleic anhydride groups, more content than others, and weight average molecular weight of 7500 g/mol, the lowest weight average molecular weight. The fact that the strong maleic anhydride concentration in oligomers tends to increase the probability of these functional groups being presents near to the wood flour. In addition, the low molecular weight of these SMA oligomers gives them sufficient fluidity in the molten state and thus supports their migration towards the fibre surface (Devaux *et al.* 2002, Bledzki *et al.* 2005). This phenomenon causes a preferential localization of SMA to the fibre-matrix interface, and thus increases the possibilities of interaction with wood flour (Gallagher *et al.* 2013). The statistical analysis reveal that no significant differences between flexural strength values were obtained using 1% or 2% of coupling agent for the three coupling agents tested. However, SMA 2000 presented better results than the others two oligomers used. It can be also concluded from the statistical analysis that uncompatibilized WPC had lower flexural strength than composites treated with SMA, which may be associated with the better compatibility between matrix and filler promoted by the coupling agent. The usage of SMA 3000 and

SMA EF40 decreased the composites flexural strength when compared with composites treated with SMA 2000. This might be explained based on the lower maleic anhydride content in SMA 3000 and SMA EF40 when compared with SMA 2000, resulting in lower adhesion between matrix and filler. The composites treated with SMA 2000 presented higher flexural modulus when compared with the others oligomers used. In addition, the flexural modulus followed the same behavior that the flexural strength and composites with 2 wt% of SMA 2000 showed the highest modulus values. Adding 1% of SMA 2000 did not affect the flexural modulus when compared with the composite without coupling agent and further increase in the coupling agent content (2wt% SMA 2000) did not alter the flexural modulus values appreciably. SMA 3000 promotes a significantly decrease in flexural modulus when compared with untreated composite and treated with SMA 2000. This behavior is probably associated with the lower maleic anhydride content in SMA 3000.

Impact strength decrease with the addition of 2 wt% of SMA 2000, as can be seen in Table 1, probably because occur an increase in the formation of entanglement between the fiber and matrix (Nygård *et al.* 2008). Furthermore, the sufficient number of maleic anhydride groups in SMA 2000 attached onto the PS chains causes strong interfacial interaction, probably due the formation of chemical bonds between these groups and hydroxyl groups of wood flour (Naghmouchi *et al.* 2015). Statistical analysis reveal that no significant differences between impact strength values were observed with addition of different levels of SMA 2000 or SMA 3000. Composites treated with SMA EF 40 showed the highest impact strength values. This result may be attribute to the lower content of maleic anhydride groups in this coupling agent, with reduces the formation of entanglement between the fiber and matrix, which can promote a better energy absorption by the polymer matrix during the impact test.

Table 1. Effect of coupling agent type and percent on the mechanical properties.

Mechanical properties evaluated	Coupling agent									
	0	SMA 2000			SMA 3000			SMA EF40		
		1	2	4	1	2	4	1	2	4
Flexural strength (MPa)	54,94 ^d (2,73)	64,81 ^a (1,73)	66,69 ^a (2,91)	44,18 ^b (2,85)	56,51 ^{d,f,g} (2,26)	59,22 ^{d,f} (1,63)	42,52 ^e (1,92)	58,13 ^{d,g,h} (3,35)	59,06 ^{d,h} (0,96)	44,98 ^c (3,91)
Flexural modulus (MPa)	4564 ^a (72)	4427 ^{a,d} (140)	4701 ^{a,b} (143)	4850 ^b (122)	4213 ^{c,d,e} (188)	4209 ^c (88)	4380 ^c (138)	4194 ^c (73)	4661 ^{a,f} (284)	4756 ^f (148)
Impact Strength (J/m)	95,98 ^a (1,62)	97,20 ^a (3,92)	93,23 ^a (5,00)	92,82 ^a (4,58)	92,62 ^a (4,75)	97,24 ^a (2,39)	91,22 ^b (2,74)	102,0 ^c (1,04)	107,43 ^d (3,30)	98,51 ^a (2,51)

Values for standard deviation are given in parentheses.

Results with the same superscript letter in their respective line are significantly the same.

The addition of SMA 2000 increased the flexural strength and flexural modulus, but drastically reduced the impact strength. This behavior is more evident when 1-2wt% of SMA 2000 was used. The interactions formed between polymer matrix and wood flour when SMA 2000 was used probably reduces the molecular mobility of matrix which reduces the energy absorption on impact, causing a reduction in composite impact strength. However, based on the flexural tests composites with 2 wt% of SMA 2000 showed the better mechanical performance when compared with the others composites studied.

Density

Mechanical properties of polymer composites are well known to be affected strongly by internal defects such as voids (Takagi *et al.* 2008, Poletto *et al.* 2011). Consequently, the density and void content usually serve as good indicators for composite strength (Borja *et al.* 2006, Takagi *et al.* 2008).

The measured density values were compared to the calculated ones on the basis of the following relationship, based on the rule of mixtures:

$$\rho_c = \rho_{EPSr} W_{EPSr} + \rho_{wf} W_{wf} \quad (\text{Equation 1})$$

where ρ_c is the calculated density of the composite material (g/cm^3), ρ_{EPSr} is the density of EPS-r ($1,07 \text{ g}/\text{cm}^3$), ρ_{wf} is the density of the wood flour ($1,41 \text{ g}/\text{cm}^3$), W_{EPSr} is the weight fraction of EPS-r (wt%), and W_{wf} is the weight fraction of the wood flour (wt%).

Void content was obtained according to:

$$V = 100 \left(\frac{\rho_c - \rho_m}{\rho_c} \right) \quad (\text{Equation 2})$$

where V is the void content (%), and ρ_m is the measured density of the composite material (g/cm^3).

The experimental results for polymer matrix, composites with and without 2 wt% of SMA 2000 and calculated density value were shown in Figure 3(a). The composite without coupling agent showed slightly lower density than composite with SMA. However, the experimental results were lower than the calculated values.

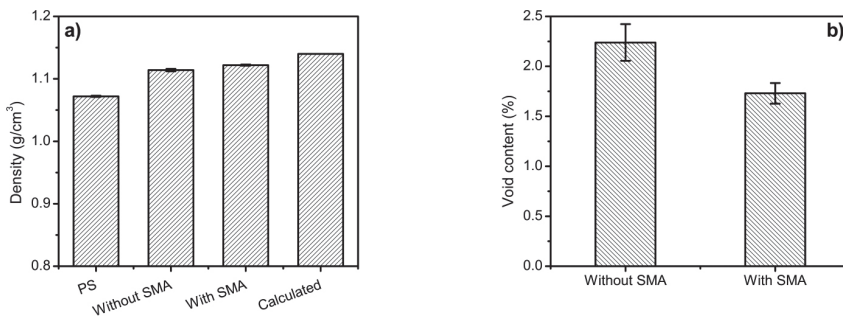


Figure 3. Experimental and calculated density values (a) and void content (b) for matrix and composites with 2wt% of SMA 2000.

The void content for WPC with coupling agent was lower than the non-treated composites, as can be seen in Figure 3(b). Composites with good mechanical properties the void content values should be less than 3% (Padma Priya *et al.* 2006). WPC with coupling agent had void content value less than 2%, while the non-treated composite value were close to 3%. Thus, the treated composite showed lower void content and better mechanical properties than non-treated one. The decrease in void content for treated composite may indicate the existence of a good bonding between wood flour and matrix in the composite as a result of using SMA 2000.

Water absorption

One of the drawbacks related to the use of lignocellulosic filler in composite materials is their high sensitivity to water, which adversely affects their mechanical properties as well as their long term durability in outdoor applications (Naghmouchi *et al.* 2015). Table 2 shows the water absorption behavior for composites with and without 2 wt% of SMA 2000. The water absorption for polystyrene was negligible during the time test, and the results were not shown.

The composites treated with SMA 2000 presented higher water absorption values when compared with composites without coupling agent. The addition of SMA 2000 introduces higher quantities of maleic anhydride groups into the composite. These maleic anhydride groups are hydrophilic and may promote an increasing in water absorption for treated composites.

Table 2. Water absorption values for composites with and without coupling agent.

Samples	Water absorption		
	2 h (%)	24 h (%)	48 h (%)
Without SMA 2000	0,138 ± 0,015	0,402 ± 0,001	0,519 ± 0,055
With SMA 2000	0,174 ± 0,014	0,467 ± 0,009	0,662 ± 0,027

Morphology characteristics

The distribution and compatibility between the fiber and the polymer matrix could be observed in SEM study. The SEM micrographs of composites without and with 2 wt% of coupling agent are shown in Figure 4(a) and (b), respectively. In Figure 4(a), examination of the cryo fracture surface of composite without coupling agent indicated the presence some voids where the fibres have been pulled-out and bigger gaps between the wood flour and matrix, which is evidence of weak interfacial adhesion at the interface (Poletto *et al.* 2011, George *et al.* 2013). The SEM micrograph of treated composite was presented in Figure 4(b). This SEM micrograph shows the strong bonding and reduced of pulled-out traces. This result demonstrated that addition of 2 wt% of SMA 2000 provides strong interfacial adhesion and good wetting, as evidenced by the almost complete absence of voids in polymer matrix and gaps between the fibre and matrix (Kim *et al.* 2007, Csikós *et al.* 2015).

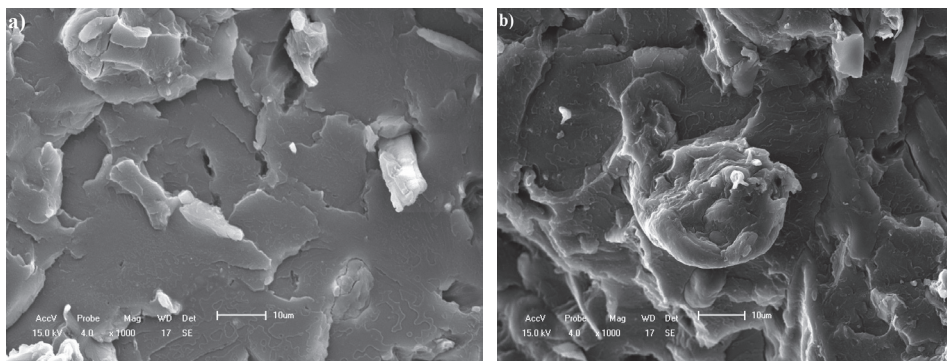


Figure 4. SEM micrographs with magnification of 1000x for (a) composite fracture surfaces without coupling agent and (b) composite fracture surfaces with coupling agent.

CONCLUSIONS

The improvement in composite mechanical properties using SMA as coupling agent was strongly dependent on the amount of maleic anhydride into the oligomer and the molecular weight of SMA. Low molecular weight gives sufficient fluidity for SMA in the molten state and thus supports the migration towards the fibre surface improving the adhesion between matrix and wood flour. The highest amount of maleic anhydride groups offers sufficient interactions between SMA and wood flour. The flexural and impact strengths of SMA-treated composites were increased compared to those of SMA non-treated composites. Satisfactory results in mechanical properties mainly in flexural strength were obtained with 2% in weight of SMA 2000. The density are almost the same for composites without and with coupling agent, however the void content was reduced in treated composites. Composites treated with SMA 2000 presented higher water absorption than untreated ones, probably because SMA 2000 introduces higher quantities of hydrophilic maleic anhydride groups into the composite. The SEM micrographs confirm better adhesion between wood flour and polystyrene matrix when coupling agent was used.

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