



Thermoplastic starch/wood composites: Interfacial interactions and functional properties



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ABSTRACT

Thermoplastic starch (TPS)/wood composites were prepared from starch plasticized with 36 wt% glycerol. The components were homogenized by dry-blending, extruded and injection molded to tensile bars. Tensile properties, structure, deformation, water adsorption and shrinkage were determined as a function of wood content, which changed between 0 and 40 vol% in 7 steps. The modification of TPS with wood particles improves several properties considerably. Stiffness and strength increases, and the effect is stronger for fibers with larger aspect ratio. Wood fibers reinforce TPS considerably due to poor matrix properties and strong interfacial interactions, the latter resulting in the decreased mobility of starch molecules and in the fracture of large wood particles during deformation. Strong interfacial adhesion leads to smaller water absorption than predicted from additivity, but water uptake remains relatively large even in the presence of wood particles. The shrinkage of injection molded TPS parts is very large, around 10%, and dimensional changes occur on a very long timescale of several hundred hours. Shrinkage decreases to a low level already at 15–20 vol% wood content rendering the composites good dimensional stability.

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1. Introduction

The constant threat of depleting fossil fuel sources and the increasing environmental awareness of the public resulted in increasing interest in natural polymers in all fields of life (Markarian, 2008). Cellulose, lignin and starch are renewable materials produced in very large quantities each year. The use of natural-based raw materials or additives in plastics has become industrial practice recently and as a consequence, the interest of the scientific community increased as well. Cellulose and wood are used as reinforcement, lignin to produce various chemicals, while under some conditions starch is applied as a thermoplastic polymer. Starch itself cannot be processed with traditional thermoplastic processing technologies because of the large size and stiffness of the molecules, and the resulting high glass transition temperature. Because of strong hydrogen bonds developing among the molecules, neat starch cannot be melted and processed without further treatment (Jang & Pyun, 1996; Swanson, Shogren, Fanta,

& Imam, 1993). Chemical modification and plasticization partially resolves these problems and plasticized starch can be processed by extrusion, injection molding or other processing methods. Besides the advantages (processability, flexibility, biodegradability, etc.) of plasticized starch, it also has disadvantages like poor mechanical properties, water sensitivity, poor dimensional instability, etc. In order to overcome these drawbacks thermoplastic starch (TPS) is often modified by blending with other polymers (Cercle, Sarazin, & Favis, 2013; Landreau, Tighzert, Bliard, Berzin, & Lacoste, 2009), the addition of fillers or reinforcements (Castillo et al., 2013; de Carvalho, Curvelo, & Agnelli, 2001; Kuciel, Kúznar, Mikula, & Liber-Knéc, 2012; Ma, Yu, & Kennedy, 2005) or the preparation of nanocomposites (Bagdi, Müller, & Pukánszky, 2006; Castillo et al., 2013; Huang, Yu, & Ma, 2004). In order to maintain one of the most important advantages of starch, i.e. biodegradability, it is modified mostly with aliphatic polyesters and natural fibers (Avérous, Fringant, & Moro, 2001; Benezet, Stanojlovic-Davidovic, Bergeret, Ferry, & Crespy, 2012; Chakraborty, Sain, Kortschot, & Cutler, 2007; Huang et al., 2004; Sreekumar, Gopalakrishnan, Leblanc, & Saiter, 2010; Sreekumar, Leblanc, & Saiter, 2010; Torres, Arroyo, & Gomez, 2007).

A considerable number of papers deal with the effect of natural fibers on the properties of thermoplastic starch. Most of them study the influence of fiber type and amount usually determining

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Table 1
Particle characteristics of the studied lignocellulosic fibers.

Fiber	Abbreviation	$D[4,3]^a$ (μm)	Length ^b (μm)	Diameter ^b (μm)	Aspect ratio ^b
Arbocel CW 630	W35	39.6	93.5	33.3	3.5
Filtracel EFC 1000	W68	213.1	363.4	63.9	6.8
Arbocel FT 400	W126	171.2	235.2	21.8	12.6

^a Volume average particle size.

^b Average values determined from scanning electron micrographs.

properties at one or two fiber contents. All kinds of fibers have been used as reinforcement in TPS including various forms of cellulose (Avérous et al., 2001 (natural cellulose fibers from leafwood); Ayadi & Dole, 2011 (natural cellulose fibers from leafwood); Benezet et al., 2012 (wheat straw fibers, hemp fibers, cotton linter fibers); Curvelo, de Carvalho, & Agnelli, 2001 (bleach pulp fiber); Ma et al., 2005 (micro winceyette fibers); Martins et al., 2009 (bacterial cellulose and vegetable cellulose fibers); Muller, Laurindo, & Yamashita, 2009; Soykeabkaew, Laosat, Ngaokla, Yodsuan, & Tunkasiri, 2012 (bacterial cellulose); Soykeabkaew, Supaphol, & Rujiravanit, 2004 (jute and flax fibers); Sreekala, Goda, & Devi, 2008), jute (Soykeabkaew et al., 2012; Torres et al., 2007; Wollerdorfer & Bader, 1998), sisal (Girones et al., 2012; Sreekumar, Gopalakrishnan, et al., 2010; Sreekumar, Leblanc, et al., 2010; Torres et al., 2007; Wang, Thompson, & Liu, 2012), wheat straw (Benezet et al., 2012), hemp (Benezet et al., 2012; Girones et al., 2012; Kunanopparat, Menut, Morel, & Guilbert, 2008; Ochi, 2006), cotton (Benezet et al., 2012; Moriana, Karlsson, & Ribes-Greus, 2010; Prachayawarakorn, Sangnitidej, & Boonpasith, 2010), flax (Saiah et al., 2009; Soykeabkaew et al., 2004), ramie (Lu, Weng, & Cao, 2006; Sreekala et al., 2008), etc. Somewhat less papers deal with TPS/wood composites (Abbott, Palazuela Conde, Davis, & Wise, 2012; Agnantopoulou, Tserki, Marras, Philippou, & Panayiotou, 2012; Avérous & Boquillon, 2004; Chakraborty et al., 2007; Kuciel et al., 2012; Kuciel & Liber-Knec, 2009), although wood is cheaper and simpler to handle during processing. Some of the papers cited investigates the effect of fiber characteristics on mechanical properties and conclude that stiffness and strength increase both with increasing fiber length and content. (Avérous et al., 2001). Others, but much fewer in number, also mention interfacial interactions and based mostly on SEM micrographs (Avérous et al., 2001; Lu et al., 2006; Prachayawarakorn, Ruttanabus, & Boonsom, 2011), the shift of absorption bands in FTIR (Prachayawarakorn et al., 2011), and changing glass transition temperature deduce that interfacial adhesion is strong between natural fibers and starch (Avérous & Boquillon, 2004; Avérous et al., 2001; Curvelo et al., 2001; Martins et al., 2009; Soykeabkaew et al., 2012). Limited number of attempts exist which estimate interfacial adhesion quantitatively (Avérous & Boquillon, 2004).

Since water sensitivity is a drawback of starch composites, its change with fiber modification is studied relatively often and the results generally indicate that water absorption decreases with increasing fiber content (Ayadi & Dole, 2011; Curvelo et al., 2001; Lu et al., 2006; Ma et al., 2005; Soykeabkaew et al., 2012). This change is explained with the smaller water uptake of the fibers compared to that of starch. Attempts to determine the dimensional stability of starch/fiber composites are much less frequent, although shrinkage might be considerable hindering application in some areas. Existing papers indicate that shrinkage decreases with increasing fiber content, i.e. modification is beneficial from this aspect as well (Avérous et al., 2001).

The study of existing literature indicated that considering their practical relevance relatively small number of papers have been published on TPS/wood composites (Abbott et al., 2012; Agnantopoulou et al., 2012; Avérous & Boquillon, 2004; Chakraborty et al., 2007; Kuciel et al., 2012; Kuciel & Liber-Knec,

2009). Even less report systematic experiments carried out as a function of fiber content in a wide composition range (Abbott et al., 2012; Agnantopoulou et al., 2012; Avérous & Boquillon, 2004). As a consequence, the goal of our work was to prepare TPS/wood composites with wood fillers of different particle characteristics, determine their reinforcing effect and influence on properties in a wide composition range. Considerable attention is paid to interfacial interactions, and reinforcement as well as interfacial adhesion are estimated quantitatively in comparison with commodity polymer/wood composites. The effect of wood characteristics and content on water absorption and shrinkage were studied and are discussed in detail.

2. Experimental

The corn starch used in the experiments was supplied by Hungrana Ltd. Hungary and its water content was 12%. Glycerol with 0.5% water content was obtained from Molar Chemicals Ltd. Hungary and it was used without further purification or drying. Three different wood fibers were used as reinforcement, all three obtained from Rettenmaier and Söhne GmbH, Germany. The particle characteristics of the fibers are listed in Table 1. Since in several of our previous projects aspect ratio turned out to be one of the most important characteristics of wood fibers (Renner, Kenyó, Móczó, & Pukánszky, 2010; Renner, Móczó, & Pukánszky, 2009), we included it into the abbreviation of the samples used. Accordingly, WP126 indicates a wood fiber with an aspect ratio of 12.6. Composites were prepared with 5, 10, 15, 20, 30 and 40 vol% wood content.

Both corn starch and the fibers were dried in an oven before composite preparation (100 °C for 24 h and 105 °C for 4 h, respectively). Thermoplastic starch containing 36 wt% glycerol was prepared by dry-blending in a Henschel FM/A10 high speed mixer at 2000 rpm. The wood fibers were introduced into the mixer at the end of the plasticization process. TPS was produced by processing the dry-blend on a Rheomex 3/4" single screw extruder attached to a Haake Rheocord EU 10V driving unit at 140–150–160 °C barrel and 170 °C die temperatures, and 60 rpm screw speed. TPS was injection molded into standard ISO 527-1A tensile bars using a Battenfeld BA 200 CD hydraulic machine at 140–160–170 °C barrel, 180 °C nozzle and 40 °C mold temperatures, 140 bar injection pressure, 54 mm/s injection rate, and 35 s cooling time.

The mechanical properties of the specimens were characterized by tensile testing using an Instron 5566 universal testing machine at 5 mm/min cross-head speed and 115 mm gauge length at 23 °C and 50% RH. Micromechanical deformations during tensile testing were followed by recording acoustic emission signals with a Sensophone AED 40/4 apparatus with 20 db threshold. The particle characteristics of the fibers and composite structure were characterized by polarized optical (POM) and scanning electron microscopy (SEM). POM micrographs were recorded on cryo-cut slices of 10 μm thickness with cross polarizers using a Leitz EM FC7 microscope. SEM micrographs (Jeol JSM-6380) were taken from the fracture surface of composite samples created during tensile testing. The crystalline structure of the TPS matrix, the wood fibers and the composites was characterized by X-ray diffraction using a Phillips PW 1830/PW 1050 equipment with $\text{CuK}\alpha$ radiation at 40 kV and 35 mA anode

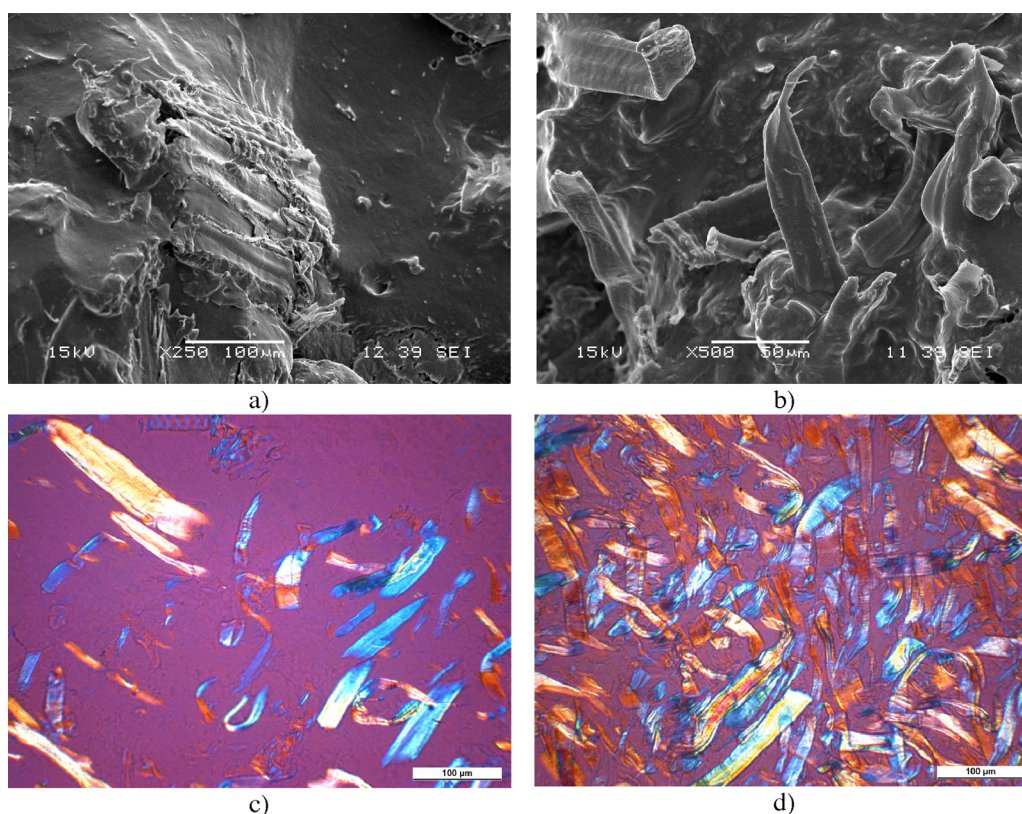


Fig. 1. Fiber distribution and the homogeneity of TPS/wood composites studied by microscopy; (a) W68, 10 vol%, SEM, (b) W126, 10 vol%, SEM, (c) W126, 5 vol%, POM, (d) W126, 40 vol%, POM.

excitation. Glass transition temperature and other thermal transitions were determined with dynamic mechanical analysis using a Perkin Elmer Pyris Diamond DMA apparatus in single cantilever mode with constant amplitude (10 μm) and frequency (1 Hz) in the temperature range between -100 and 80°C with a heating rate of $2^\circ\text{C}/\text{min}$. Water absorption was determined at 23°C and 52% relative humidity by the measurement of the weight increase of the specimens. The desired relative humidity was achieved with saturated solution of $\text{Mg}(\text{NO}_3)_2$. Shrinkage was followed by the measurement of the length of the specimens as a function of time. The specimens were kept in a desiccator over silica gel to avoid the complicating influence of water absorption.

3. Results and discussion

The results are presented in several sections. Mechanical properties and structure are described first and then the reinforcing effect of the fiber is discussed in the next section. Special attention is paid to interfacial interactions, and the effect of wood reinforcement on functional properties is discussed in the final section of the paper.

3.1. Structure and mechanical properties

Aggregation, attrition and orientation of anisotropic particles are the most important structure related phenomena in particulate filled and fiber reinforced composites. Previous studies on wood reinforced composites have shown that aggregation rarely occurs, but fibers might touch each other purely from geometric reasons (Dányádi, Móczó, & Pukánszky, 2010; Dányádi, Renner, Móczó, & Pukánszky, 2007; Renner, Kenyó, et al., 2010; Renner et al., 2009). Long fibers may even form a loose network (Faludi, Dora, Renner, Móczó, & Pukánszky, 2013; Faludi, Hári, Renner, Móczó, &

Pukánszky, 2013). SEM micrographs taken from the fracture surface of two composites is shown in Fig. 1a and b. A single large particle embedded into the matrix is seen in Fig. 1a presenting the micrograph recorded on a TPS/W68 composite containing 10 vol% wood. In Fig. 1b several broken fibers appear on the surface of a TPS/W126 composite of the same composition and the micrograph clearly indicates the good dispersion of the particles. One must be aware of the fact that SEM micrographs can always be debated and they can be used only as supplementary evidence to support conclusions from other results. Nevertheless, they can demonstrate the fracture of the fibers quite well as shown by Fig. 1b. They are completely inadequate to demonstrate homogeneity and the quality of dispersion cannot be judged reliably based on them. On the other hand, POM micrographs taken at smaller magnification offer a clear picture of structure (Fig. 1c and d). Such a micrograph recorded on a TPS/W126 composite at 5 vol% wood content shows the homogeneous distribution of the fibers in the TPS matrix (Fig. 1c). At 40 vol% wood content the fibers touch each other (Fig. 1d) as shown earlier (Faludi, Dora, et al., 2013), but real aggregates do not form even at this fiber loading. Nevertheless, the physical contact of the fibers probably influences mechanical properties to some extent. Microscopy clearly proves that the distribution of wood particles is homogeneous at small, but physical contacts form among them at large fiber contents.

Stiffness is used the most frequently to estimate the effect of fillers and fibers on the properties of composites. The modulus of our TPS/wood composites is plotted against fiber content in Fig. 2a. According to the figure stiffness increases from around 0.35 GPa up to approximately 4.5 GPa, i.e. modification with wood results in very strong reinforcement. The initial steep increase slows down at larger fiber contents indicating some structural effect, probably the mere physical contact of the particles discussed above (Dányádi et al., 2007; Renner, Kenyó, et al., 2010; Renner et al., 2009) or the

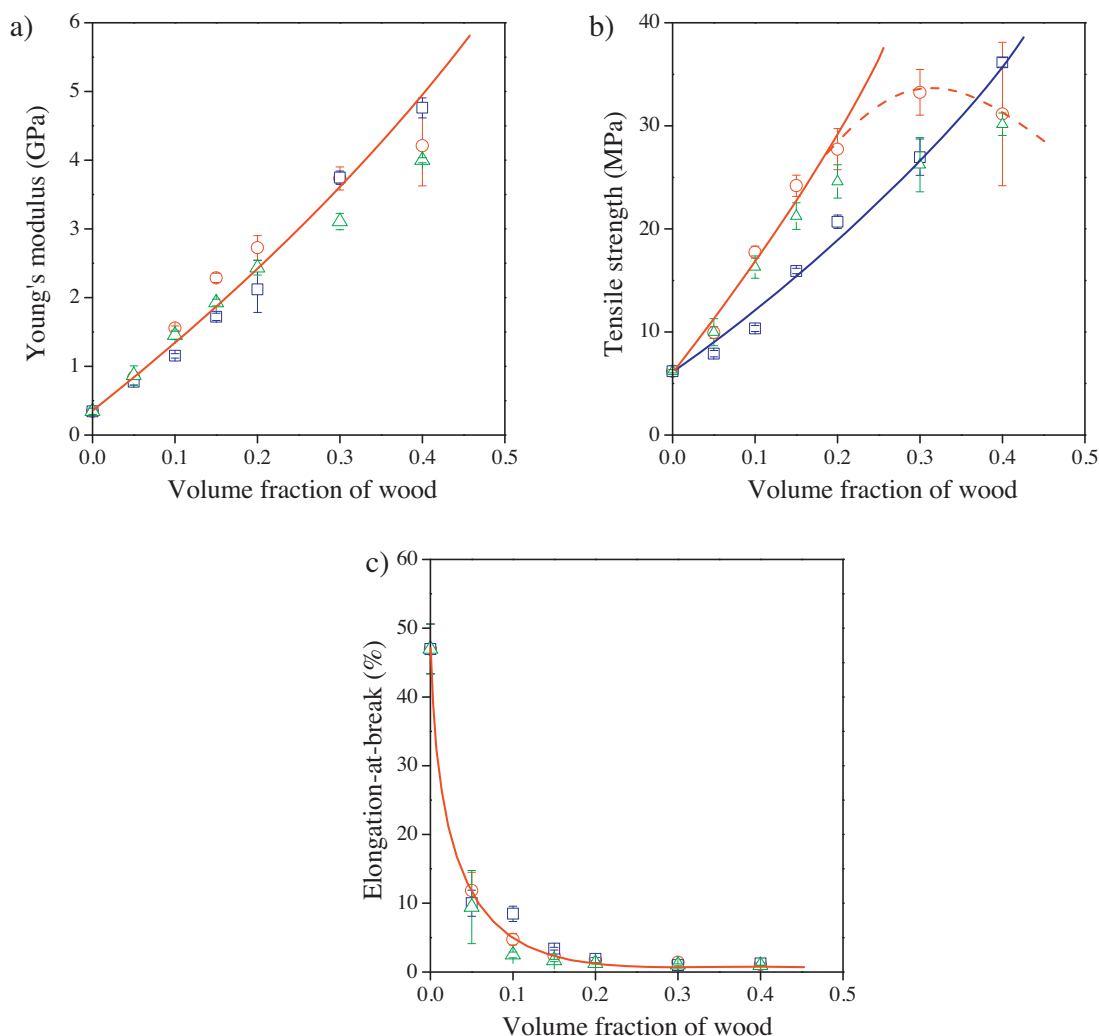


Fig. 2. Mechanical properties: effect of wood fiber type and content on the stiffness (a), strength (b) and elongation-at-break (c) of TPS/wood composites; symbols: (○) W126, (□) W68 and (△) W35.

formation of a network in the case of fibers with large aspect ratio (Faludi, Hári, et al., 2013). The composition dependence of mechanical properties strongly supports this explanation since the largest deviation from the tendency observed at large fiber content occurs for the small (W35) and the long (W126) fibers (see Table 1). This latter has larger reinforcing effect at small fiber contents than the other two wood flours (W35, W68).

The tensile strength of composites containing the three wood fibers is plotted against wood content in Fig. 2b. The same conclusions can be drawn here as from the composition dependence of modulus. Strength increases at least as strongly as stiffness and the different fibers influence also this property dissimilarly. Small (W35) and long particles (W126) reinforce TPS more than the third wood flour (W68), but structural effects are also stronger in the former two cases. Deformability, i.e. the elongation-at-break of the specimens decreases drastically with increasing wood content (Fig. 2c), but such an effect is usually expected both in particulate filled and fiber reinforced composites. The effect of the three fibers is very similar; specific surface area seems to influence deformability, but differences are quite small. We can conclude from these results that wood reinforcement is beneficial if stiffness and strength are the targeted properties, considerable reinforcement occurs, but its extent is difficult to judge from the qualitative evaluation of the results.

3.2. Reinforcement

Reinforcement, i.e. the load-bearing capacity of the second component, can be estimated quantitatively by the use of the simple model developed earlier to describe the composition dependence of tensile yield stress (Turcsányi, Pukánszky, & Tüdös, 1988), tensile strength (Pukánszky, 1990) or fracture resistance (Pukánszky & Maurer, 1995) of particulate filled polymers. The model takes the following form for tensile strength

$$\sigma_T = \sigma_{T0} \lambda^n \frac{1 - \varphi}{1 + 2.5\varphi} \exp(B\varphi) \quad (1)$$

where σ_T and σ_{T0} are the true tensile strength ($\sigma_T = \sigma \lambda$ and $\lambda = L/L_0$) of the composite and the matrix, respectively, n is a parameter taking into account strain hardening, φ is the volume fraction of the fiber and B is related to its relative load-bearing capacity, i.e. to the extent of reinforcement, which, among other factors, depends also on interfacial interaction. If we transform Eq. (1) into a linear form and plot the natural logarithm of reduced tensile strength against fiber content, we obtain a linear correlation (Fig. 3a), the slope of which is proportional to the load-bearing capacity of the reinforcement and under certain conditions to the strength of interaction. The correlations are linear at small fiber loadings indeed, but deviations are observed at large wood contents. Such deviations usually

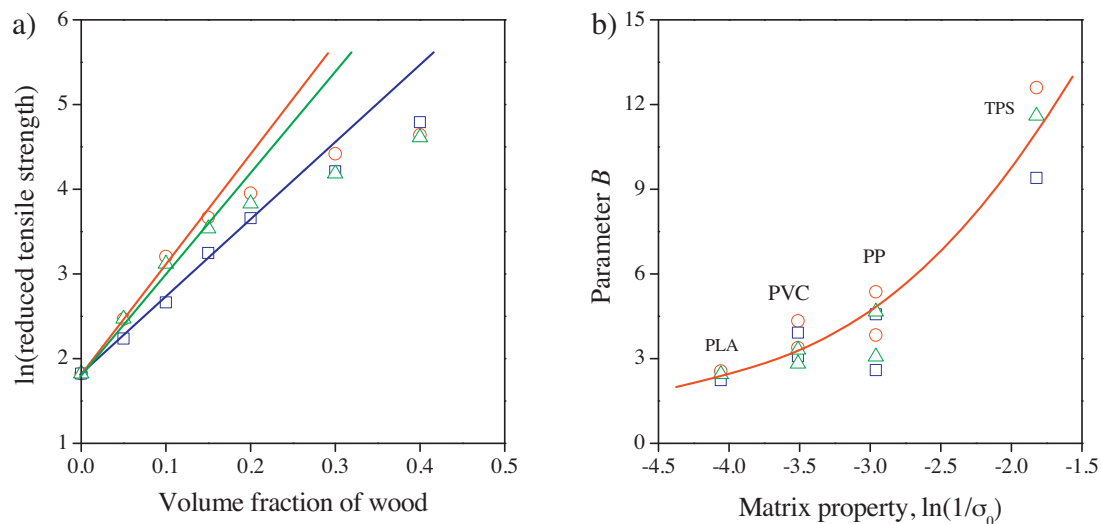


Fig. 3. Reinforcement: (a) tensile strength of TPS/wood composites plotted against wood content in the linear representation of Eq. (1). Symbols: (○) W126, (□) W68, (△) W35; (b) influence of matrix strength on the reinforcing effect of wood fibers.

indicate structural effects, i.e. they confirm our tentative explanation presented in the previous section. The slope of the straight lines is parameter B , which expresses the load-bearing capacity of the second component, i.e. reinforcement. The value of B ranges from 9.4 to 12.6 for the three wood fibers used, and they are very large compared to usual particulate filled commodity polymers; B is often smaller than 1 for PP/CaCO₃ composites, for example (Kiss, Fekete, & Pukánszky, 2007; Pukánszky, 1990).

The reinforcing effect of a filler or fiber depends on interfacial interactions, but also on the properties of the matrix (σ_0), which plays an important role in the actual value of B . This role can be seen even better if we plot B values as a function of the natural logarithm of reciprocal matrix strength, i.e. $\ln(1/\sigma_0)$ (see Pukánszky, 1990). The correlation is presented for various thermoplastic wood composites in Fig. 3b. We can see that the inherent property of the matrix dominates in the determination of the extent of reinforcement. However, reinforcement depends also on interfacial adhesion, thus we need further study and analysis to define its strength.

3.3. Interfacial adhesion

As described in the introductory part, several approaches are used to estimate the strength of interfacial interactions in TPS/natural fiber composites. FTIR and the determination of the glass transition temperature were mentioned as specific examples. The temperature dependence of the loss modulus of TPS and that of the composites containing the three wood reinforcements at 30 vol% is plotted in Fig. 4. Both TPS and the composites exhibit two transitions, one at low temperature which can be assigned to the relaxation of smaller structural units and which is related to the amount of glycerol (Avérous & Pollet, 2012; Forsell, Mikkilä, Moates, & Parker, 1997), while the second above room temperature which is identified as the glass transition temperature of TPS. Reinforcement with wood results in considerable increase of this latter indicating large decrease in the mobility of starch molecules. This is rather surprising, since very small changes can be seen in the T_g of particulate filled polymers and only at large contact surfaces (Bleach, Nazhat, Tanner, Kellomäki, & Törmälä, 2002; Mortezaei, Famili, & Kokabi, 2010). The probable reasons for the large effect are strong hydrogen bonds and the stiffness of starch molecules; a few secondary bonds are sufficient to decrease mobility. Even the presence and competitive interaction of the small glycerol molecules

do not influence this effect. Wood properties do not seem to play a role here, but we must call the attention here to the fact that the specific surface area of the three fillers is very similar. We can conclude that strong interactions develop between TPS and wood at the segmental and molecular level, but we do not know much about macroscopic interfaces.

Various deformation processes may take place around the inclusions in composites. Besides debonding, i.e. the separation of the interface between the matrix and the reinforcement, several other processes including matrix yielding, the pull-out or fracture of the fibers, cavitation, etc. may also take place during the deformation of the composites (Dányádi et al., 2007; Faludi, Dora, Renner, Móczó, & Pukánszky, 2013; Imre et al., 2012; Renner et al., 2007; Renner, Móczó, Suba, & Pukánszky, 2010). Some of these are burst-like processes emitting elastic waves, which can be picked up by appropriate sensors. The result of such an acoustic emission measurement is presented in Fig. 5a. The small circles are individual acoustic events (signals); one of the continuous lines shows the cumulative number of events (right axis), while the other is the stress vs. elongation trace (left axis) shown as reference. We can see that no acoustic activity can be detected below certain deformation or stress, and a characteristic stress (σ_{AE}) can be determined in the way indicated in Fig. 5a. This stress is related to the initiation

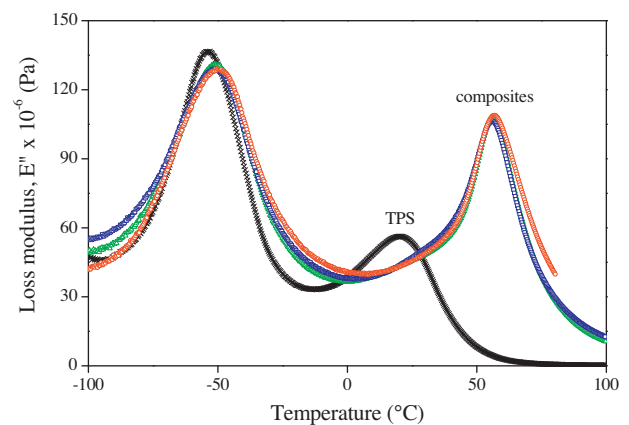


Fig. 4. Temperature dependence of the loss modulus of TPS and its wood composites at 30 vol% wood content; decreased mobility in the presence of wood; symbols: (○) W126, (□) W68 and (△) W35.

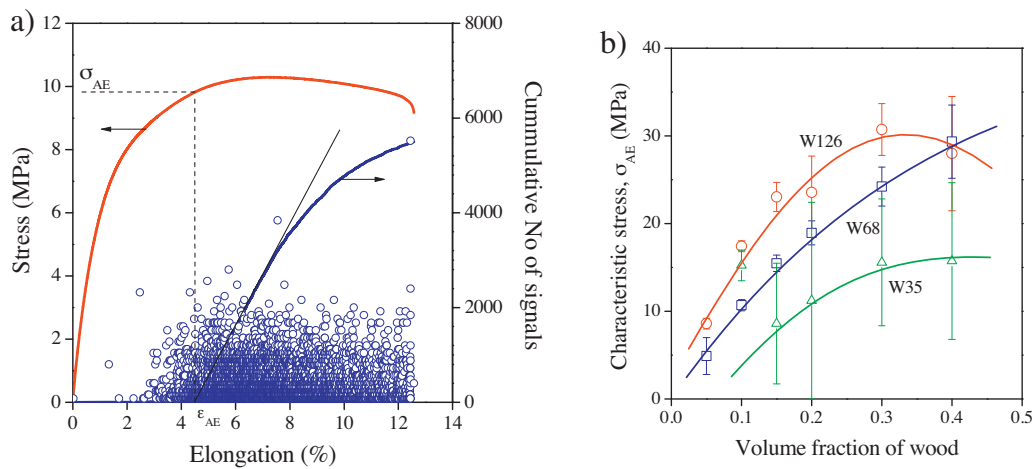


Fig. 5. Acoustic emission testing of TPS/wood composites. (a) TPS/W126 composite, wood content: 5 vol%. Symbols: (○) individual acoustic events. Stress (left axis) and cumulative number of events (right axis) vs. elongation traces are plotted as reference. Determination of characteristic stress (σ_{AE}); (b) characteristic stress of the dominating micromechanical deformation mechanism plotted against fiber content; symbols: (○) W126, (□) W68 and (△) W35.

of the dominating micromechanical deformation process taking place around wood particles upon deformation. The composition dependence of this characteristic stress is plotted against wood content in Fig. 5b. We can see differences in the effect of the three fibers; the longest and largest fibers initiate the process at larger, while the small particles at smaller stress. We cannot identify the dominating deformation process either from the value of the characteristic stress or the shape of the cumulative number of signal trace; it can be debonding or the fracture of the wood particles, as shown earlier (Dányádi et al., 2007; Renner et al., 2009; Renner, Kenyó, et al., 2010).

If debonding is the dominating process, interfacial adhesion can be estimated quantitatively with the help of a model developed earlier (Pukánszky & Vörös, 1993). According to the model debonding stress can be expressed as

$$\sigma^D = -C_1 \sigma^T + C_2 \left(\frac{E F_a}{R} \right)^{1/2} \quad (2)$$

where σ^D and σ^T are debonding and thermal stresses, respectively, E the Young's modulus of the matrix, F_a the strength of interfacial adhesion, which is equal to the reversible work of adhesion, if adhesion is created by secondary forces, R the radius of the particles, while C_1 and C_2 are geometric constants related to the debonding process. The parameters of the equation were calculated from measurements done on polymer/filler pairs with known characteristics (E , R , F_a). If we know the stiffness of the matrix and the size of the particles, which we usually do, we can calculate the strength of adhesion (F_a) (Renner, Mócsó, Vörös, & Pukánszky, 2010). In our case we can assume that $\sigma^D = \sigma_{AE}$ and calculate F_a from Eq. (2). The results are listed in Table 2 for the three set of composites. Polypropylene (PP) reinforced with one of the fillers (W35) is used as reference. PP is a commodity polymer used in large quantities also as the matrix of wood composites. Moreover, interfacial adhesion can be changed considerably by coupling, thus the adhesion

values determined at poor and good adhesion can serve well as reference. In the absence of coupling interaction is poor in PP, while covalent bonds are assumed to form when a functionalized coupling agent (MAPP) is used. We can see one order of magnitude difference in the strength of adhesion in the two cases. The adhesion calculated in the TPS composite containing the small particles (W35) is similar to that determined in PP at good adhesion, but considerably larger values are obtained for the other two fibers indicating very good adhesion. Since parameter B appearing in Eq. (1) is also related to interfacial adhesion, we can use it also for the estimation of the strength of interaction, but we must compensate for the effect of changing matrix property. We did that by multiplying B by $\ln(1/\sigma_0)$ and the results are also listed in Table 2. They convey practically the same message as F_a , i.e. interfacial adhesion is very strong in TPS/wood composites. Even if not debonding, but some other mechanism dominates in TPS composites, it is certain that strong interfacial interaction develops between the components considerably influencing properties.

We could not define the dominating micromechanical deformation process taking place during the deformation of TPS/wood composites from the composition dependence of properties or from acoustic emission measurements. We assumed that SEM micrographs taken from fracture surfaces created during tensile testing would offer further information about interfacial adhesion and processes occurring around the particles. Two micrographs taken from composites containing 10 vol% fibers were presented in Fig. 1a and b in a previous section. The fracture surface of the TPS/W68 composite presented in Fig. 1a shows a very large broken particle which is firmly embedded in the matrix. A larger number of broken fibers can be observed on the surface of the TPS/W126 composite. Most of the fibers fracture perpendicularly to their axis, which occurs only at very strong interactions. In spite of all uncertainties of the method, the SEM study obviously supports our conclusion that interaction is strong in the TPS/wood composites studied and not debonding, but fiber fracture is the dominating process. Apart from the firm embedding of the particles into the TPS matrix, very few characteristic features can be seen on the fracture surface of TPS/W35 composites (not shown).

3.4. Functional properties

In order to fulfill their function, products prepared from TPS composites must satisfy certain conditions. The adsorption of excessive water modifies properties, stiffness and strength

Table 2
Comparison of estimated interfacial adhesion in TPS/wood and PP/wood composites.

Matrix	Fiber	Coupling	Adhesion, F_a (mJ/m ²)	Parameter B	$B \ln(1/\sigma_0)$
PP	W35	No	150	2.3	−6.8
PP	W35	MAPP	1190	4.7	−13.8
TPS	W35	No	1200	11.6	−21.1
TPS	W68	No	3800	9.4	−17.1
TPS	W126	No	3900	12.6	−22.9

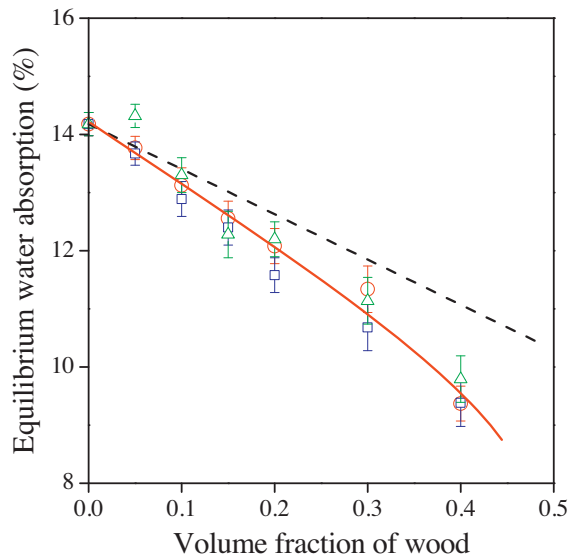


Fig. 6. Effect of wood content and fiber type on the equilibrium water absorption (M_{∞}) of TPS/wood composites. The broken line indicates additivity; symbols: (○) W126, (□) W68 and (△) W35.

decrease or the part can fail completely. Similarly, shrinkage may result in unacceptable dimensional changes. Earlier studies showed that the reinforcement of TPS with natural fibers decreases its water uptake, i.e. results in the improvement of this characteristic (Ayadi & Dole, 2011; Curvelo et al., 2001; Lu et al., 2006; Ma et al., 2005; Soykeabkaew et al., 2012). Equilibrium water absorption was determined from adsorption isotherms by fitting the following form of Fick's law to the experimental results

$$M_t = M_{\infty} \left(1 - \frac{8}{\pi^2} \left(\exp(-at) + \frac{1}{9} \exp(-9at) + \frac{1}{25} \exp(-25at) \right) \right) \quad (3)$$

where M_t is time dependent weight increase, M_{∞} the final (equilibrium) water uptake reached after infinite time, t the time of adsorption and a a constant characterizing the overall rate of water adsorption. The equilibrium water uptake of the composites is plotted in Fig. 6 showing that water absorption decreases with wood content, as reported in the literature, and the effect of the three wood fibers is very similar. The broken line in Fig. 6 indicates

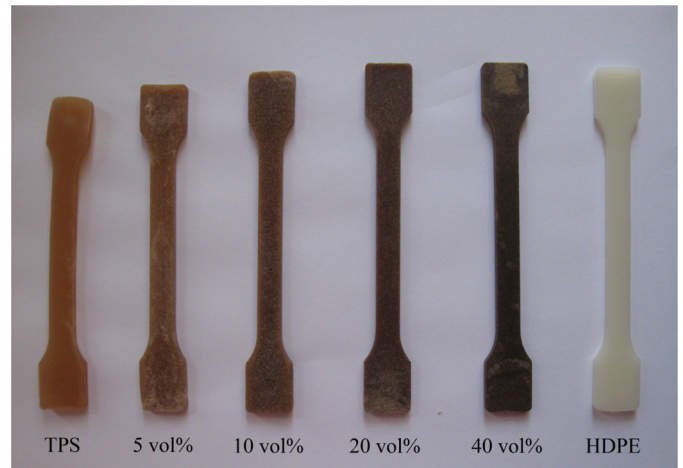


Fig. 7. Shrinkage of injection molded bars produced from TPS/W68 composites of various wood contents. Shrinkage was measured 345 days after production.

additivity, i.e. the value of water uptake calculated from that of TPS and wood for each composition. We can see that the actual water absorption of the composites is smaller than the additive value. A probable explanation might be the strong interaction between the components discussed in the previous section. We must call the attention here to the fact though, that water absorption is rather large even at 40 vol% wood content, i.e. it might hinder the application of these materials in certain fields.

The shrinkage of TPS is another important factor which may limit its application. Fig. 7 shows the photo of injection molded TPS/W68 composite tensile bars with different fiber contents taken 345 days after production. A HDPE bar is used as reference. HDPE is a crystalline polymer with a relatively large shrinkage in engineering practice. Compared to commodity and especially to engineering thermoplastics the shrinkage of TPS is extremely strong and some warpage also occurs. Such shrinkage obviously cannot be tolerated in certain applications. We recorded XRD traces on the bars as a function of time to check if the slow crystallization of TPS results in the considerable shrinkage. We did not find any changes in crystalline structure and concluded that the reason of these large dimensional changes is the limited flexibility of the starch molecules and their long relaxation times. Injection molding results in non-equilibrium structure and the presence of the

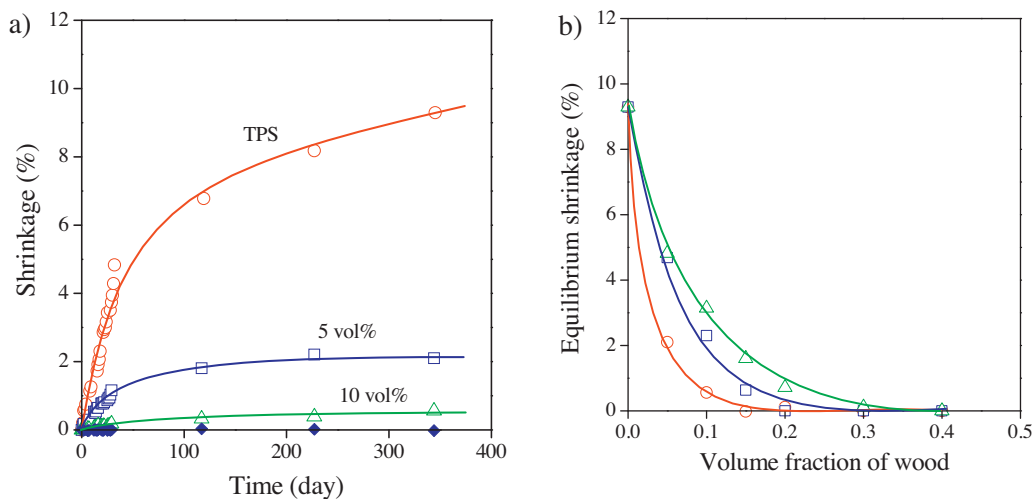


Fig. 8. Shrinkage of TPS/wood composites: (a) time dependence of the shrinkage of TPS/W126 composites at different wood contents; symbols: (○) 0, (□) 5, (△) 10 vol%; (b) Dependence of equilibrium shrinkage on wood content and fiber type; symbols: (○) W126, (□) W68, (△) W35.

plasticizer allows relaxation, but only on a very long time scale. The time dependence of shrinkage is shown in Fig. 8a for the composites containing the large aspect ratio fibers (W126). Shrinkage increases with time according to a saturation function on a very long timescale. Shrinkage is very large for TPS, but the presence of already 5 vol% fiber decreases shrinkage considerably, while above 10 vol% shrinkage can be practically neglected, at least compared to that of neat TPS. We should like to call the attention here to the long time scale of the experiment. The effect of wood content on the final, equilibrium shrinkage can be seen much better in Fig. 8b together with the effect of fiber characteristics. Shrinkage decreases very fast with increasing wood content and reaches very small values already at 15 vol% wood content. In accordance with literature data, fibers with larger aspect ratio hinder shrinkage in a larger extent than short particles. Obviously, wood reinforcement has very advantageous effect on the shrinkage of TPS and it may extend the application area for these materials considerably.

4. Conclusions

The modification of TPS with wood particles improves several properties considerably. Stiffness and strength increases, and the effect is stronger for fibers with larger aspect ratio. Wood fibers reinforce TPS considerably due to poor matrix properties and strong interfacial interactions, the latter resulting in the decreased mobility of starch molecules and in the fracture of large wood particles during deformation. Strong interfacial adhesion leads to smaller water absorption than predicted from additivity, but water uptake remains relatively large even in the presence of wood particles. The shrinkage of injection molded TPS parts is very large, around 10%, and dimensional changes occur on a very long timescale of several hundred hours. Shrinkage decreases considerably to a low level already at 15–20 vol% wood content rendering the composites good dimensional stability. Wood reinforcement of TPS is generally advantageous for most application areas.

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