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IMPROVEMENTS IN WOOD THERMOPLASTIC MATRIX COMPOSITE MATERIALS PROPERTIES BY PHYSICAL AND CHEMICAL TREATMENTS

Abstract: This paper presents a short overview of the developments made in the field of wood thermoplastic composites in terms of surface treatment, flammability, matrix/reinforcement model, properties and application of recycled polymer matrices. The usage of lignocellulosic fibers as reinforcement in composite materials demands well formed interface between the fiber and the matrix. Because of the different nature of reinforcement and matrix components some physical and chemical treatment methods which improve the fiber matrix adhesion were introduced, as well as the improvements of lignocellulosic fibers and thermoplastic polymer matrix based composites flammability characteristics. These physical and chemical treatments influence the hydrophilic character of the lignocellulosic fibers, and therefore change their physical and mechanical properties.

Keywords: bio-composites, wood thermoplastic composites, recycled polymer matrices

1. Introduction

Wood, as a natural polymer matrix composite, is the oldest material of this type. Constitutive components of wood are lignin with hemicelluloses as a soft polymer matrix, and rigid cellulosic microfibrils as reinforcement, so called lignocellulosic fibers (Rong et al., 2011). The properties of wood reinforcing fibers depends on the type of plants, as well as the parts of a plant, because each vegetal creation has its own crystalline organization (Bledzki and Gassan. 1999). The application of lignocellulosic fibers as a reinforcement in polymer matrix composites has received

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increased attention. From the all lignocellulosic fibres, wood fiber is the most broadly used as composite materials reinforcement. Generally known lignocellulosic composites are laminated lumber, plywood, fiberboard, particleboard and wood plastic composite materials (WPCs) (Rowell, 1990). A large amount of wood waste in different form such as wood pulp, fibers or flour has been generated in wood industry. This wood waste forms are very suitable as a reinforcement for thermoplastic polymer matrices (Woodhams et al.. 1984). On the other hand. incompatibility between non-polar hydrophobic thermoplastic matrices and polar hydrophilic wood reinforcement creates problems during the production of composites and results with their law mechanical properties (Yeh and Gupta,

2008). Due to poor adhesion between the polar wood and the non-polar polymer, increasing amount of wood reinforcement causes reduction in composite strength (Klyosov, 2007). In order to overcome this problem, a lot of research has been done (Bledzki and Gassan, 1999; Satyanarayana et al., 2009; Adekomaya et al., 2016). Exposure to UV radiation and humidity also cause the reduction of WPC mechanical properties due to reduced fiber/matrix adhesion properties, lignin degradation and polymer chain scission, so changes occurring during accelerated weathering also were researched (Beg and Pickering, 2008; Stark and Matuana, 2003; Matuana and Kamdem, 2002; Matuana et al., 2001). As a matrix materials in WPC virgin but also recycled polymers have been used (Yeh et al., 2009; Adhikary et al., 2008; Augier et al., 2007). The most used thermoplastic polymers as a matrix in wood plastic composite materials are polyethylene, polypropylene, poly vinyl chloride, polystyrene and acrylonitrilebutadiene-styrene (Bledzki and Gassan, 1999; Yeh et al., 2009).

Because of superior strength/weight and stiffness/weight ratios of WPCs, their use in automotive industries is increasing. Automotive components, such as dashboards, door panels, seats, and cabin linings were made from WPCs (Ashori, 2008). Currently, aspirations of using WPCs for car parts includes replacements glass fiber reinforced composites and steels (Bismarck *et al.*, 2006).

2. Surface treatment

The surface structure and energy of the wood reinforcement can be changed by using different methods of physical and chemical surface treatment. These changes influence the mechanical bondings fiber/matrix, leading to different interface in composite material and different mechanical properties. The cellulose fibers treatment with oxygen plasma increased the free surface energy with increasing treatment time or level of

corona treatment, which leads to an improved wettability of composite materials (Belgacem *et al.*, 1994; Sakata *et al.*, 1993). Physical treatments also include thermotreatment, stretching and calandering processes. Chemical modification includes delignification pretreatment, fiber surface modification and coupling agents' application (Bledzki and Gassan, 1999).

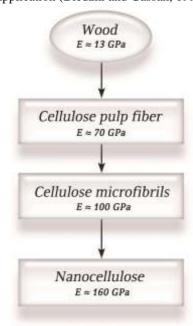


Figure 1. Levels of wood structure

Due to complex wood composition, modifications can be applied on the each level of structure observation, Figure 1.

2.1. Delignification

Removal of lignin from wood fibers increases fiber/matrix adhesion due to better physical bonding between cellulose fiber and thermoplastic matrix, as it has been shown in Figure 2 (Mai *et al.*, 1983; Beg and Pickering, 2008).



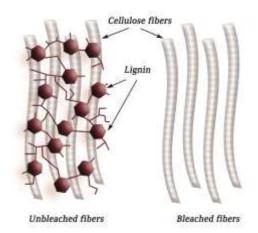


Figure 2. Schematic representation of the removal of lignin from wood fibers

Lignin and hemicelluloses are the most responsible component for ultraviolet degradation of wood (Nevell and Zeronian, 1985). and Pickering Beg (2008),investigated moisture/ultraviolet (UV) weathering performance of unbleached and bleached 40 wt% wood fiber reinforced polypropylene (PP) composites, with 3 wt% of a maleated polypropylene (MAPP) coupling agent. Initially, bleached fiber composites showed better mechanical properties, but after accelerated weathering, extent of the mechanical properties reduction was almost the same for both bleached and unbleached wood fiber composites. The changes occurred during accelerated weathering were registrated with differential X-ray diffraction, thermogravimetric analysis, scanning calorimetry and scanning electron microscopy. For long-term application of PP composites authors recommended usage of PP UV stabilizer.

2.2. Coupling agents and fiber surface modification

Interfacial adhesion in WPC can be improved by the chemical modification with the introduction of a coupling agent. Often this coupling agent is a functionalized polymer. As functionalized polyolefin

composite matrix for improving interfacial adhesion, prevent debonding and the formation of voids into the composite materials, maleated polypropylene (MAPP) or polyethylene (MAPE) has been used (Keener et al., 2004; Li and Matuana, 2003; Myers et al., 1990; Kazayawoko et al., 1999; Oiu et al., 2003; Kazayawoko et al., 1997; Ndiaye et al., 2011; Hristov et al., 2004; Lu et al., 2005; Bledzki et al., 2002; Beg and Pickering, 2008). The surface of wood reinforcement contains the-OH groups which form chemical bond with maleic anhydride groups. Result of this formation is improved interfacial bonding. Schematic representation of the modification of interface by maleated polypropylene in wood/PP composite material is shown in Figure 3. For modification of wood reinforcement the other anhydrides, such as succinic or phthalic, also were used (Matsuda, 1987; Zhang et al., 1994; Gellerstedt and Gatenholm, 1999; Kokta et al., 1990), but maleic shows the best improvement in mechanical properties.

Beside anhydrides, in order to enhance interfacial adhesion in wood composites, poly[methylene(polyphenyl isocyanate)] (Maldas *et al.*, 1989), copper amine (Jiang *et al.*, 2003; Kamdem *et al.*, 2004), chromated copper arsenate (Stilwell *et al.*, 2003) and silane (Kokta *et al.*, 1990; Bengtsson and Oksman, 2006), were also used as chemical coupling agents.

Improved flammability of WPC materials is of the great importance for thermal insulators and automotive constitutive elements (Ashori, 2008). Commonly the composites based on polypropylene matrix and wood fiber reinforcement were used for purposes and many studies of flammability of iPP composites with various content of long cellulose fibers was conducted (Borysiak et al., 2006; Helwig and Paukszta, 2000; Lv et al., 2005; Bras et al., 2005). There are two methods for improvements of WPC flammability with retardant agents: during the manufacturing process and during the finishing process

(Kim and Pal, 2010; Kozlowski and Helwig, 1998). Typically additive flame retardants minerals (aluminum hydroxide, are magnesium hydroxide, red phosphorus or borates), but modification of organohalogens and organophosphorus compounds also results of significant improving flammability (Sain et al., 2004; Kozlowski and Wladyka-Przybylak, 2000; Anna et al., 2003). Baysal et al. (2007) used boric acid and borax mixture for impregnation of pine wood reinforcement/vinyl copolymers. Tests show improvements in antiswelling reduced absorption, efficiency, water modulus of elasticity, resistance against decay caused by fungi and decrease in flammability.

Dányádi et al. (2010), researched influence of surface modification on interfacial interactions in PP/wood composites by addition of a MAPP, benzylation as chemical modification of the wood surface and application of surfactants. The addition of a maleated polymer improve interfacial adhesion of PP/wood composites, but does not influence homogeneity, viscosity and water absorption regardless of coupling agent amount. Homogeneity was improved surfactants. and water absorption decreasing was obtained by chemical modification of wood by benzylation.

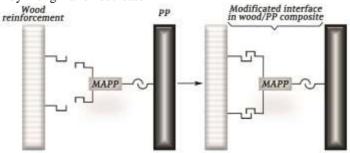


Figure 3. Schematic representation of the interface modification in wood/PP composite material with functionalized polymer coupling agents (MAPP)

Previous research of Dányádi et al. (2007) reported influence of MAPP different molecular weights and functionality on interfacial adhesion of PP/wood composites with wood content from 0 to 70 wt%. They proved that MAPP with the larger molecular weight and smaller functionality provide better improvement of composite strength. On the other hand, MAPP with the smaller molecular weight reduces viscosity and improves processability of WPC. upgrading of WPC processability is of the great importance because of increased production of different components for the civil and automotive industry, packaging or furniture (Bledzki and Gassan, 1999; Adekomaya et al., 2016; Bledzki et al., 2005; Jacob, 2006; Bledzki et al., 2006).

Resent research (Xu et al., 2014) includes application of chitosan as a coupling agent in

wood/PVC composites. Beside its biological activities. Chitosan owns excellent thermal and anti-bacterial mechanical, properties (Amri et al., 2013; Lu et al., 2010; Martinez-Camacho et al., 2010; Cheung et al., 2007; Vasile et al., 2013). It has similar function as the chemical coupling agents with amino-groups because of consisting amino-polysaccharide, Figure 4. The big advantages of chitosan are low price and environmentally friendly characteristics which makes it suitable for the production of biocomposite materials.

Xu et al, (2014) investigated effects of chitosan on interfacial adhesion and on the thermal and rheological properties of wood flour/polyvinyl chloride composites. They reveal optimal amount and particle size of chitosan for better heat resistance capacity, glass transition temperature and thermal



stability of analyzed wood composites. It was proved that low or high content of chitosan resulting in the low vicat softening temperature which value is the highest when the amount of chitosan achieves an optimal value of 30 phr. The vicat softening temperature of wood/PVC composites with chitosan as coupling agents were compared

with the same composite with silane as a popular chemical coupling agent. The results proved the better heat resistance capability of wood/PVC composites with chitosan coupling agent. Xu *et al.* (2014) researched effects of particle sizes of chitosan and prove that smaller particle size provide the better interfacial compatibility.

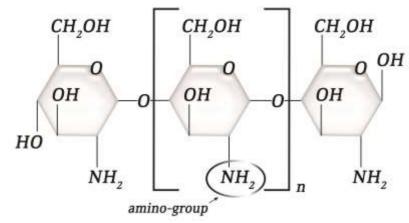


Figure 4. Chemical structure of chitosan

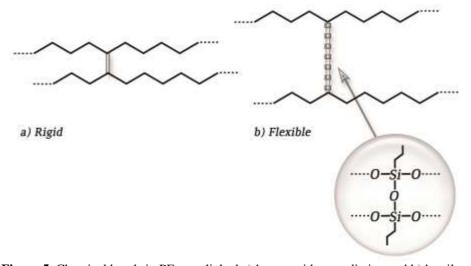


Figure 5. Chemical bonds in PE crosslinked a) by peroxide or radiation and b) by silane technology

Beside the most common used coupling agents functionalized polymers MAPP and MAPE, the silanes are also widely used

(Oksman and Clemons, 1998; Kokta *et al.*, 1989; Raj *et al.*, 1989). Silanes are monomeric silicon compounds with four



substituent groups attached to the silicon atom. These substituent groups can be very different in composition and reactivity (nonreactive, inorganically or organically reactive). Vinyl silanes can be grafted onto the polyethylene chains giving crosslinked polyethylene. While the C-C bond obtained in PE crosslinked by peroxide or radiation is rigid, structure of PE crosslinked by silane technology contains flexible bonds, Figure 5. In this manner used polyethylene as matrix materials significantly improves properties of composite with wood reinforcement (Kokta et al., 1989; Kokta et al., 1989). Content of wood reinforcement in WPC with silane grafted HDPE matrix could be high up to the point when there are no free silanol groups which could further react with hydroxyl groups in cellulose fibers (Bengtsson et al., 2005; Park and Balatinecz, 1998). Bengtsson and Oksman (2006) investigated the use of silane technology in crosslinking PE/wood flour composites. Softwood from spruce and pine was used as wood reinforcement. The results show significantly higher flexural strength and elongation at break in the silane crosslinked composites compared with non-crosslinked ones. Improved adhesion between the wood reinforcement and crosslinked PE and strengthening of the matrix upon crosslinking cause toughness and impact strength improvements of the composite materials. Disadvantage of such prepared WPC is the lower flexural modulus in the crosslinked samples noncrosslinked ones. The authors explain the lower flexural modulus as a consequence of a lower crystallinity in the crosslinked composites and plastification of non-reacted silane. The proved lower creep in the crosslinked composites was interpreted with reduced viscous flow due to crosslinking and increased adhesion between wood reinforcement and thermoplastic PE matrix.

In order to increase adhesion and interface properties of wood fibers and non-polar thermoplastics such as, HDPE, PP and PVC, a pretreatment of fibers was often applied. For this purpose a lot of methods and the different treatments have been researched (Shukla *et al.*, 1993; Liao *et al.*, 1997). The organic carboxylic acids (e.g. stearic acid, acetic anhydride) also have been applied for surface modification (Mahlberg *et al.*, 2001; Zafeiropoulos *et al.*, 2002).

Zheng et al. (2007) investigated the effect of benzoic acid surface modification on the mechanical properties of the PVC matrix composites reinforced with lignocellulosic fibers. They proved that treatment with benzoic acid provides 37% the better tensile strength of the composite compared to the tensile strength of untreated composite. The lignocellulosic fiber treatment with benzoic acid significantly improved dispersion of the fiber in PVC thermoplastic matrix which is supported by SEM micrographs of impact fracture testing composites materials.

An another example of dual modification in WPC has been given by Wang et al. (2015). They improved the interfacial adhesion through modifying both composite components, thermoplastic matrix (PP) and bamboo wood fiber. In order to establish strong interfacial interaction between polar reinforcement and non-polar matrix, MAPP was used as a coupling agent and bamboo cellulose fiber was oxidized with 2,2,6,6tetramethylpiperidine-1-oxy radical (TEMPO). It was proved that reactive hydroxyl groups on the fiber surface created by oxidation with TEMPO successfully bonded with MAPP modified PP. It was supported by FTIR, XRD and TGA analysis. Tensile strength of dual modified WPC cellulose fiber/PP based on 50% of fiber content is significantly improved.

2.2. Water absorption

Water has a adverse influence on the mechanical properties of WPC (Stark and Matuana, 2004). In contact with water and moisture, reaction of oxidation take place-on the wood fiber surfaces and causes swelling of the WPC. Degradation of lignin or thermoplastic matrix in WPC during



weathering make cellulosic fibers unprotected and cause moisture absorption inside the composite material (Stark, 2006).

Chemical treatments can increase interface adhesion between the polar fiber and non-polar matrix, but also can decrease the water absorption of hydrophilic wood reinforcement. As mentioned bleached wood fiber reinforced composites containing MAPP, shows better mechanical properties after accelerated weathering than the unbleached one (Beg and Pickering, 2008). Generally, the main disadvantage of wood fibre/thermoplastic polymer composite materials is their hydrophilic nature. Due to using of wood fibre reinforced composites in outdoor applications demands their greater environmental and dimensional stability, study of their mechanical properties under the influence of weathering attracted great attention (Beg and Pickering, 2008; Cantero et al., 2003).

Mechanical properties before and after accelerated weathering of wood reinforcement/thermoplastic LLDPE-g-MAPP matrix was investigated by Kallakas et al. (2015). Three different wood flour fractions sizes from birch and bleachedchemi-thermo-mechanical aspen reinforcement surface were treated with two coupling agents, aminopropyltriethoxysilane (APTES) and polyvinyalcohol (PVA). The best tests results were obtained with composites made of APTES modified wood flour and LLDPEg-MAPP. Increasing the wood flour fractions size in the composites increases the amount of absorbed water as well as swelling. Consequently, flexural modulus and strength, and deflection of tested WPC decreased. Damages on the cross-section of WPC after weathering tests were supported by SEM analysis.

Butylina *et al.* (2012), studied softwood/PP composites behavior exposed a long-term weathering to Finnish climatic conditions. For analyzing the weathering caused degradation of the composites a combination

of SEM, FTIR spectroscopy, colorimetric measurements and DSC were used. It is proved that the composite with a higher polypropylene content and lubricant was less damaged by weathering. Impact strength of analyzed WPC was determinated by Charpy test method. The results reveal a decrease of impact strength for composites with higher water absorption after weathering.

Effect of wood species and particle size on water absorption in WPC with HDPE matrix and MAPP as coupling agent, were analyze by (Dorostkar *et al.*, 2014). Results showed decreased water absorption percent with increasing reinforcement particle size, but significant differences of water absorption property between different species was not observed.

3. Recycled polymer matrices

Products obtained from recycled thermoplastic polymers include also matrices in WPC. The most common recycled are polypropylene, polyvinyl polymers chloride, acrylonitrile butadiene styrene (ABS), high-density polyethylene (HDPE), etc. Pattanakul et al. (1991) investigate properties of the recycled HDPE from used milk bottles and proved that the properties of the recycled and virgin polymer does not have difference of great scale. On the other hand, wood fibers and flour from industry waste are very suitable as reinforcement for thermoplastic polymer matrices, giving composite materials for automotive furniture. components. enterior design. packaging and many other commercial products. Adhikary et al. (2008) used sawdust of pine softwood and recycled or virgin HDPE with MAPP coupling agent for manufacturing composite samples. Results show that the content of wood reinforcement significantly influences water absorption and swelling of WPC. Samples prepared with coupling agents MAPP showed improved characteristics. Only 3-5wt% of MAPP increase amount of optimal boow reinforcement from 30wt% to 50wt% for

achievement equivalent stability properties. Improved interfacial bonding of composite with modified HDPE matrix is observed by SEM images of the fracture surfaces of WPC. Authors concluded that dimensional stability and mechanical properties of composites can be achieved by increasing the polymer content or by addition of coupling agents.

Ashori and Nourbakhsh (2009) compared properties of WPCs with recycled HDPE and recycled PP. In both case reinforcement was obtained from old newspaper and it was used in different percentage. For improvement of reinforcement/matrices interface MAPP was used as coupling agent. Results report that WPC with recycled HDPE achieved moderately superior mechanical properties compared with composite with recycled PP. Also, composites made with low content of reinforcement without coupling agent show lower mechanical properties. Less water absorption, dimensional stability and good mechanical properties can be obtained using MAPP as coupling agents in 2-4 wt%. The similar approach was done with recycled and virgin ABS making composites with maple wood flour as reinforcement (Yeh et al., 2009). The coupling agents were not used.

Currently, recycling of WPC waste attracts attention due to economy and environment protecting reasons. According to European Guideline 2000/53/EG administered by the European Commission, means of transport have to be produced with 95% recyclable materials consisting reproduced WPCs (Ashori et al., 2008). Petchwattana et al. (2012), studied changes in the mechanical and physical properties upon closed-loop recycling and reprocessing reinforcement/PVC of wood matrix composite materials. The composite were produced by mixing of WPC scrap and WPC Based on the mechanical properties appropriate mixing ratio of the WPC scrap: WPC freshfeed at 30:70 was established by maximizing the use of WPC scrap. Due to chain scission of PVC melted during each re-processing pass a drop in the

flexural strength and flexural modulus was observed. Implementation of recycled WPC materials in the production of new constitutive parts and improvement of their properties is still developing.

5. Conclusions

The main disadvantage of lignocellulosic fibers is their strong polar character. Contrary, thermoplastic polymers has nonpolar character which creates incompatibility with lignocellulosic fibers. To overcome this problem many physical and chemical surface treatments were developed. Application of coupling agent can significantly increase the interface adhesion of polar lignocellulosic fiber with thermoplastic matrices, and also decrease the water absorption of composite materials. The most commonly used coupling agents with the high improvements WPC properties are maleated polypropylene (MAPP), chitosan and silane. Environmentally friendly characteristic is the big advantage of chitosan which makes it suitable for the production of bio-composite Lignocellulosic materials. pretreatment is preferable to improve interfacial adhesion and decrease water absorption. For this purpose different surface modification were studied, including the oldest one - delignification. Organic carboxylic acids, benzoic acid, and 2,2,6,6tetramethylpiperidine-1-oxy radical show very favorable results. For cellulosic fiber lignin and thermoplastic matrix in WPC have protecting role, so their degradation cause serious damage in WPC. Due to requirements for outdoor application of WPC many researches focused on their accelerated weathering. Bleached wood fiber reinforced composites containing MAPP, show good mechanical properties after accelerated weathering. The good results were obtained bv using aminopropyltriethoxysilane (APTES) coupling agents.

Recycling of thermoplastic polymers and their composites attract attention due to



enviroment protection requirements. As the most common recycled matrices are polypropylene, polyvinyl chloride, acrylonitrile butadiene styrene (ABS) and high-density polyethylene (HDPE). It is proved that the properties of the recycled and virgin HDPE does not have difference of great scale. Mechanical properties of composites with recycled polymer matrices can be improved by using coupling agents.

WPC waste can be included in re-processing with maximum of 30%. Application possibility of recycled WPC materials and improvement of their properties is still developing.

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References:

- Adekomaya, O., Jamiru, T., Sadiku R., & Huan Z. (2016). A review on the sustainability of natural fiber in matrix reinforcement A practical perspective, *Journal of Reinforced Plastics and Composites*, 35(1), 3–7.
- Adhikary, K.B., Pang, S., & Staiger, M.P. (2008). Dimensional stability and mechanical behaviour of wood–plastic composites based on recycled and virgin high-density polyethylene (HDPE). *Composites Part B*, *39*, 807–815.
- Amri, F., Husseinsyah, S., & Hussin, K. (2013). Mechanical, morphological and thermal properties of chitosan filled polypropylene composites: the effect of binary modifying agents. *Composites Part A*, 46, 89–95.
- Anna, P., Zimonyi, E., Mariton, A., Matko, S., Keszi, S., Bertalan, G., & Marosi, G. (2003). Surface treated cellulose fibres in flame retarded PP composites. *Macromolecular Symposia*, 202, 246–264.
- Ashori, A. (2008). Wood–plastic composites as promising green-composites for automotive industries! *Bioresource Technology*, 99, 4661–4667.
- Ashori, A., & Nourbakhsh, A. (2009). Characteristics of wood–fiber plastic composites made of recycled materials. *Waste Management*, 29, 1291–1295.
- Augier, L., Sperone, G., Vaca-Garcia, C., & Borredon, M.E. (2007). Influence of the wood fibre filler on the internal recycling of poly (vinyl chloride)-based composites. *Polymer Degradation and Stability*, *92*, 1169-1176.
- Baysal, E., Yalinkilic, M.K., Altinok, M., Sonmez, A., Peker, H., & Colak, M. (2007). Some physical, biological, mechanical, and fire properties of wood polymer composite (WPC) pretreated with boric acid and borax mixture. *Construction and Building Materials*, 21, 1879–1885.
- Beg, M.D.H., & Pickering K.L. (2008). Accelerated weathering of unbleached and bleached Kraft wood fibre reinforced polypropylene composites. *Polymer Degradation and Stability*, *93*, 1939–1946.
- Beg, M.D.H., & Pickering, K.L. (2008). Reprocessing of wood fibre reinforced polypropylene composites. Part II: Hygrothermal ageing and its effects. *Composites Part A*, 39, 1565–1571.
- Belgacem, M., Bataille, P., & Sapieha, S. (1994). Effect of corona modification on the mechanical properties of polypropylene/cellulose composites. *Journal Of Applied Polymer Science*, 53(4), 379-385. http://dx.doi.org/10.1002/app.1994.070530401

- Bengtsson, M., & Oksman, K. (2006). The use of silane technology in crosslinking polyethylene/wood flour composites. *Composites Part A*, 37, 752–765.
- Bengtsson, M., Gatenholm, P., & Oksman, K. (2005). The effect of crosslinking on the properties of polyethylene/wood flour composites. *Composites Science and Technology*, 65(10), 1468–1479.
- Bismarck, A., Baltazar-Y-Jimenez, A., & Sarlkakis, K. (2006). Green composites as Panacea? Socio-economic aspects of green materials. *Environment, Development and Sustainability*, 8(3), 445–463.
- Bledzki, A.K., & Gassan, J. (1999). Composites reinforced with cellulose based fibers. *Progress in Polymer Science*, 24, 221–274.
- Bledzki, A.K., Faruk, O., & Huque, M. (2002). Physico-mechanical studies of wood fiber reinforced composites. *Polymer-Plastics Technology* and *Engineering*, 41(3), 435–451.
- Bledzki, A.K., Faruk, O., & Sperber, V.E. (2006). Cars from bio-fibres. *Macromolecular Materials and Engineering*, 291(5), 449–457.
- Bledzki, A.K., Letman, M., Viksne, A., & Rence, L. (2005). A comparison of compounding processes and wood type for wood fibre-PP composites. *Composites Part A*, 36(6), 789–797.
- Borysiak, S., Paukszta, D., & Helwig, M. (2006). Flammability of woodepolypropylene composites. *Polymer Degradation and Stability*, *91*, 3339–3343.
- Bras, M.L., Duquesne, S., Fois, M., Grisel, M., & Poutch, F. (2005). Intumescent polypropylene/flax blends: a preliminary study. *Polymer Degradation and Stability*, 88, 80–84.
- Butylina, S., Hyvärinen, M., & Kärki, T. (2012). A study of surface changes of wood-polypropylene composites as the result of exterior weathering. *Polymer Degradation and Stability*, 97, 337-345.
- Cantero, G., Arbelaiz, A., Mugika, F., Valea, A. (2003). Mondragon I. Mechanical behaviour of wood/polypropylene composites: effects of fibre treatments and ageing processes. *Journal of Reinforced Plastics Composites*, 22, 37–50.
- Cheung, H.Y., Lau, K.T., Lu, T.P., & Hui, D. (2007). A critical review on polymer-based bioengineered materials for scaffold development. *Composites Part B*, 38(3), 291–300.
- Dányádi, L., Janecska, T., Szabó, Z., Nagy, G., Móczó, J., & Pukánszky, B. (2007). Wood flour filled PP composites: Compatibilization and adhesion. *Composites Science and Technology*, 67, 2838–2846.
- Dányádi, L., Móczó, J. & Pukánszky B. (2010). Effect of various surface modifications of wood flour on the properties of PP/wood composites. *Composites Part A*, 41, 199–206.
- Dorostkar, A., Rafighi, A., & Madhoushi, M. (2014). Investigation on Water Absorption Property of Wood Plastic Composite, *International Journal of Plant, Animal and Environmental Sciences*, 4(2), 633-638.
- Gellerstedt, F., & Gatenholm, P. (1999). Surface properties of lignocellulosic fibers bearing carboxylic groups. *Cellulose*, *6*, 103–121.
- Helwig, M., & Paukszta, D. (2000). Flammability of composites based on polypropylene and flax fibers. *Molecular Crystals* and *Liquid Crystals*, 354, 373-380.
- Hristov, V.N., Krumova, M., Vasileva, S., & Michler, G.H. (2004). Modified polypropylene wood flour composites. II Fracture deformation and mechanical properties. *Journal of Applied Polymer Science*, 92, 1286–1292.

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- Jacob, A. (2006). WPC industry focuses on performance and cost. *Reinforced Plastics*, 50, 32–33
- Jiang, H.H., Kamdem, D.P., Bezubic, B., & Ruede, P. (2003). Mechanical properties of poly(vinyl chloride)/wood flour/glass fiber hybrid composites. *Journal of Vinyl and Additive Technology*, 9(3), 138–145.
- Kallakas, H., Poltimäe, T., Süld, T.M., Kers, J. & Krumme, A. (2015). The influence of accelerated weathering on the mechanical and physical properties of wood-plastic composites. *Proceedings of the Estonian Academy of Sciences*, 64, 94–104.
- Kamdem, D.P., Jiang, H.H., Cui, W.N., Freed, J., & Matuana, L.M. (2004). Properties of wood plastic composites made of recycled HDPE and wood flour from CCA-treated wood removed from service. *Composites Part A*, 35(3), 347–355.
- Kazayawoko, M., Balatinecz, J.J., & Matuana, L.M. (1999). Surface modification and adhesion mechanism in woodfiber–polypropylene composites. *Journal of Materials Science*, *34*(24), 6189–6199.
- Kazayawoko, M., Balatinecz, J.J., & Woodhams, R.T. (1997). Diffuse reflectance Fourier transform infrared spectra of wood fibers treated with maleated polypropylenes. *Journal of Applied Polymer Science*, 66(6), 1163–1173.
- Keener, T.J., Stuart, R.K., & Brown, T.K. (2004). Maleated coupling agents for natural fibre composites. *Composites Part A*, 35(3), 357–362.
- Kim, J.K., & Pal, K. (2010). Recent Advances in the Processing of Wood-Plastic Composites, Engineering Materials, Heidelberg, Germany: Springer-Verlag Berlin Heidelberg.
- Klyosov, A.A. (2007). Wood-plastic composites. Hoboken, NJ: Wiley.
- Kokta, B.V., Maldas, D., Daneault, C., & Beland, P. (1990). Composites of polyvinyl chloride–wood fibers. I: effect of isocyanate as a bonding agent. *Polymer-Plastics Technology and Engineering*, 29(1–2), 87–118.
- Kokta, B.V., Raj, R.G., & Daneault, C. (1989). Use of wood flour as filler in polypropylene: studies on mechanical properties. *Polymer-Plastics Technology* and *Engineering*, 28(3), 247–259.
- Kozlowski, R., & Helwig, M. (1998) Progress in fire retardants for lignocellulosic materials. In: *Proceedings of the 6th Arab international conference on materials science, materials & fire*. Alexandria. Egypt: Springer Berlin Heidelberg. 1-11
- Kozlowski, R., & Wladyka-Przybylak, M. (2000). *Natural polymers: wood and lignocellulosics*. In: Horrocks, R. (ed.) Fire Retardant Materials. Cambridge, UK: Woodhead Publishing Limited,
- Li, Q., & Matuana, L.M. (2003). Effectiveness of maleated and acrylic acid-functionalized polyolefin coupling agents for HDPE–wood-flour composites. *Journal of Thermoplastic Composite Materials*, *16*, 551–564.
- Liao, B., Huang, Y.H., & Cong, G.M. (1997). Influence of modified wood fibers on the mechanical properties of wood fiber-reinforced polyethylene. *Journal of Applied Polymer Science*, 66, 1561–1568.
- Lu, J.Z., Wu, Q., & Negulescu, I.I. (2005). Wood–fiber/high density-polyethylene composites: coupling agent performance. *Journal of Applied Polymer Science*, *96*(1), 93–102.
- Lu, X.L., Qiu, Z.Y., Wan, Y., Hu, Z.L., & Zhao, Y.X. (2010). Preparation and characterization of conducting polycaprolactone/chitosan/polypyrrole composites. *Composites Part A*, *41*(10), 1516–1523.

- Lv, P., Wang, Z., Hu, K., & Fan, W. (2005). Flammability and thermal degradation of flame retarded polypropylene composites containing melamine phosphate and pentaerythritol derivatives. *Polymer Degradation and Stability*, 90, 523-534.
- Mahlberg, R., Paajanen, L., & Nurmi, A. (2001). Effect of chemical modification of wood on the mechanical and adhesion properties of wood fiber/polypropylene fiber and polypropylene/veneer composites. *Holz als Roh–und Werkstoff*, 59, 319–326.
- Mai, Y.W., Hakeem, M.I., & Cotterell, B. (1983). Effects of water and bleaching on themechanical properties of cellulose fibre cements. *Journal of Materials Science*, *18*, 2156–2162.
- Maldas, D., Kokta, B.V., & Daneault, C. (1989). Composites of polyvinyl chloride—wood fibers: IV. Effect of the nature of fibers. *Journal of Vinyl and Additive Technology*, 11(2), 90–99.
- Martinez-Camacho, A.P., Cortez-Rocha, M.O., Ezquera-Brauer, J.M., Graciano-Verdugo, A.Z., Rodriguez-Felix, F., & Castillo-Ortega, M.M. (2010). Chitosan composite films: thermal, structural, mechanical and antifungal properties. *Carbohydrate Polymers*, 82(2), 305–315.
- Matsuda, H. (1987). Preparing and utilisation of esterified woods bearing carboxylic groups. *Wood Science and Technology*, 21, 75–88.
- Matuana L.M., & Kamdem D.P. (2002). Accelerated ultraviolet weathering of PVC/wood fibre composites. *Polymer Engineering Science*, 42(8), 1657–1666.
- Matuana, L.M., Kamdem, D.P., & Zhang, J. (2001). Photoaging and stabilization of rigid PVC/wood-fibre composites. *Journal of Applied Polymer Science*, 80(11), 1943–1950.
- Myers, G.E., Kolosick, P.C., Chahyadi, I.S., Coberly, C.A., Koutsky, J.A., & Ermer, D.S. (1999) Extruded wood-flour polypropylene composites: effect of a maleated polypropylene coupling agent on filler–matrix bonding and properties. *MRS Proceedings*, 197, 67–77.
- Ndiaye, D., Matuana L.M., Morlat-Therias, S., Vidal, L., Tidjani, A., & Gardette, J.L. (2011). Thermal and Mechanical Properties of Polypropylene/Wood-Flour Composites. *Journal of Applied Polymer Science*, 119, 3321–3328.
- Nevell, T.P., & Zeronian, H.S., (1985). *Cellulose chemistry and its applications*. New York: Wiley.
- Oksman, K., & Clemons, C. (1998). Mechanical properties and morphology of impact modified polypropylene-wood flour composites. *Journal of Applied Polymer Science*, 67(9), 1503–1513.
- Park, B.D., & Balatinecz, J.J. (1998). Short term flexural creep behavior of wood-fiber/polypropylene composites. *Polymer Composites*, 19(4), 377–382.
- Pattanakul, C., Selke, S., Lai, J.M. (1991). Properties of recycled high-density polyethylene from milk bottles. *Journal of Applied Polymer Science*, 43(11), 2147–2150.
- Petchwattana, N., Covavisaruch, S., & Sanetuntikul, J. (2012). Recycling of wood–plastic composites prepared from poly(vinyl chloride) and wood flour. *Construction and Building Materials*, 28, 557–560.
- Qiu, W., Zhang, F., Endo, T., & Hirotsu, T. (2003). Preparation and characteristics of composites of high-crystalline cellulose with polypropylene: effects of maleated polypropylene and cellulose content. *Journal of Applied Polymer Science*, 87(2), 337–345.

International Journal for Quality Research



- Raj, R.G., Kokta, B.V., Maldas, D., & Daneault, C. (1989). Use of wood fibers in thermoplastics. VII. The effect of coupling agents in polyethylene-wood fiber composites. *Journal of Applied Polymer Science*, *37*(4), 1089–1103.
- Rong, M.Z, Zhang, M.Q., Liu, Y., Yang, G.C. & Zeng, H.M. (2001). The effect of fiber treatment on the mechanical properties of unidirectional sisal-reinforced epoxy composites. *Composites Science and technology*, 61, 1437-1447.
- Rowell, R.M. (1990). *Opportunities for Lignocellulosic Materials and Composites*, In: Emerging technologies for materials and chemicals from biomass. Proceedings of symposium. Washington, DC. USA: American Chemical Society, 12-27
- Sain, M., Park, S.H., Suhara, F., & Law, S. (2004). Flame retardant and mechanical properties of nature fiber-PP composites containing magnesium hydroxide. *Polymer Degradation and Stability*, 83, 362–367.
- Sakata, I. Morita, M., Tsuruta, N., & Morita, K. (1993). Activation of wood surface by corona treatment to improve adhesive bonding. *Journal* of *Applied Polymer Science*, 49, 1251–1258.
- Satyanarayana, K.G., Arizaga, G.G.C., & Wypych, F. (2009). Biodegradable composites based on lignocellulosic fibers—An overview, *Progress in Polymer Science*, *34*, 982–1021.
- Savoia, M., Stefanovic, M., & Fragassa, C. (2016). Merging Technical Competences and Human Resources with the Aim at Contributing to Transform the Adriatic Area in Stable Hub for a Sustainable Technological Development. *International Journal of Quality Research*, 10(1), 1-16.
- Shukla, S.R., Rao, G.V.G., Athalye, A.R. (1993). Improving graft level during photoinduced graft-copolymerization of styrene onto cotton cellulose. *Journal of Applied Polymer Science*, 49, 1423–1430.
- Stark, N.M. (2006). Effect of weathering cycle and manufacturing method on performance of wood flour and high-density polyethylene composites. *Journal of Applied Polymer Science*, *100*, 3131-3140.
- Stark, N.M., & Matuana L.M. (2003). Ultraviolet weathering of photostabilized wood-flourfilled, *Journal of applied polymer science*. 90, 2609-2617
- Stark, N.M., & Matuana, L.M. (2004). Surface chemistry and mechanical property changes of wood-flour/high-density-polyethylene composites after accelerated weathering. *Journal of Applied Polymer Science*, 94, 2263-2273.
- Stilwell, D., Toner, M., & Sawhney, B. (2003). Dislodgeable copper, chromium and arsenic from CCA-treated wood surfaces. *Science of the Total Environment*, *312*, 123–131.
- Vasile, C., Darie, R.N., Cheaburu-Yilmaz, C.N., Pricope, G.M., Bracic, M., & Pamfil, D. (2013). Low density polyethylene–chitosan composites. *Composites Part B*, 55, 314–323.
- Wang, S., Lin, Y., Zhang X. & Lu, C. (2015). Towards mechanically robust cellulose fiberreinforced polypropylene composites with strong interfacial interaction through dual modification. *RSC Advances*, 5, 50660–50667.
- Woodhams, R.T., Thomas, G., & Rodgers, D.K. (1984). Wood fibers as reinforcing fillers for polyolefins. *Polymer Engineering and Science*, 24(15), 1166–1171.
- Xu, K.M., Li, K.F., Zhong, T.H., & Xie, C.P. (2014). Interface self-reinforcing ability and antibacterial effect of natural chitosan modified polyvinyl chloride based wood flour composites. *Journal of Applied Polymer Science*, 131(3), 39854.

- Xu, K.M., Li, K.F., Zhong, T.H., Guan, L., Xie, C.P., & Li, S. (2014). Effects of chitosan as biopolymer coupling agent on the thermal and rheological properties of polyvinyl chloride/wood flour composites, *Composites Part B*, 58, 392–399.
- Yeh, S.K., & Gupta, R.K. (2008). Improved wood–plastic composites through better processing, *Composites: Part A*, *39*, 1694–1699.
- Yeh, S.K., Agarwal, S., & Gupta, R.K. (2009). Wood–plastic composites formulated with virgin and recycled ABS. *Composites Science and Technology*, 69, 2225–2230.
- Zafeiropoulos, N.E., Williams, & D.R., Baillie, C.A. (2002). Engineering and characterisation of the interface in flax fibre/polypropylene composite materials. Part I. Development and investigation of surface treatments. *Composites Part A*, *33*, 1083–1093.
- Zhang, Y., Sjögren, B., Engstrand, P., & Htun, M. (1994) Determination of charged groups in mechanical pulp fibres and their influence on pulp properties. *Journal of Wood Chemistry and Technology*, 14, 83–102.
- Zheng, Y.T., Cao, D.R., Wang, D.S., & Chen, J.J. (2007). Study on the interface modification of bagasse fibre and the mechanical properties of its composite with PVC, *Composites Part A*, 38, 20–25.

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