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SEPTEMBER/OCTOBER 2018



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Chairman's Message:



Ray Boeman, Ph.D.

he year has gone by incredibly fast, and I personally have never been more excited and optimistic about the future of our industry, and this division. Key elements of shaping this future are engaging new ideas and fostering a growing infusion of young talent through strong support for educational initiatives and programs that enrich the experience of young professionals.

With respect to education, the division continues to expand its activities. We received several excellent proposals this year for our university grant program. Grants were awarded to Auburn University; University of Alabama, Tuscaloosa; University of Southern California; and University of Wisconsin. More information on the grant program can be obtained from the Division's Education Chair, Prof. Uday Vaidya (<u>uvaidya@utk.</u> <u>edu</u>). The division also supports the SPE PlastiVan (<u>www.4spe.org/spe-plastivan/</u>) for which we are expanding our sponsorship from three events to five in the coming year.

I am pleased to report that Creig Bowland, John Gillespie, and Dale Grove were reelected to their board seats and that Alex Kravchenko, an Assistant Professor at Old Dominion University, was newly elected to the board. I am also pleased to announce that Dale Brosius was re-elected as the Composite Division Councilor. Three of our board members received awards this year in recognition of their leadership. Prof. John Gillespie, Jr. was recognized as Fellow of the Society, and Prof. Uday Vaidya was recognized as Honored Service Member (HSM). The division's Chair-elect, Ian Swentek has been recognized by SAMPE with a Young Professional Emerging Leadership Award. Additionally, the Composites Division has again received the SPE Pinnacle Gold award reflecting significant and committed efforts of our board members.

Under the leadership of Shankar Srinivasan and Rich Caruso, the division organized and moderated five sessions and two keynote presentations for another successful ANTEC conference in May. Please note that the timing for ANTEC has been moved up for 2019 – it will be held March 18-21 in Detroit. As you read this you may very well be attending another tremendously successful ACCE which continues to be this division's keystone event held jointly with the Automotive Division.

I am convinced the Composites Division (www.4spe.org/composites) and the industry overall have a bright future. Please feel free to reach out to me or any of the board members to get more involved and to share your thoughts on where the future may take us.

Best regards, Ray







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Treasury Report



By: Tim Johnson, Treasurer



he Composite Division continues to maintain a positive budget. Currently the Division has cash on the order of \$94K and \$74K in investment. With increased income from the ACCE, and Sponsorship of the Newsletter that continues to exceed costs, the Division has steadily increased awards. Notably, the Board recently approved increased support of the SPE Plastivan Program for the coming year, and a substantial increase in the support of Education matching grants which will exceptionally draw on reserves.

The Composites Division continues to provide significant support for student activities at ANTEC and ACCE, and looks forward to the expanded outreach through the Plastivan and Education Grant programs.



Award Winners



By: Dr. Dale Grove



2018 Travel Award Winners

t is with great pleasure that I announce the travel award winners for 2018: Connor Armstrong from the University of Maryland and Zhaogui Wang from Baylor University. Connor is presently striving for his Master's degree as he studies the orientation behavior of carbon microfibers in 3D printing through divergent dies as well as

self-healing thermoplastic composites, while Zhaogui plans to get a PhD as he models various fiber orientation models combined with different matrix constitutive algorithms including complex viscoelastic matrix models.

I enjoyed meeting both of them, and I wish them continued success in their endeavors.

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Connor Armstrong



Zhaugui Wang

Award Winners continued...



2018 Harold Giles Award Winners

he 2018 Harold Giles Scholarship award winners are Mr. Yourri-Samuel Dessureault and Mr. Matthew Swift. Sixteen applications were received this year. It is always a difficult job to select winners based on the many talented applications that the judges received.

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Mr. Matthew Swift Undergraduate Student Winner



Mr. Yourri-Samuel Dessureault Graduate Student Winner

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Award Winners continued...



Mr. Dessurealt presently attends Florida State University and is pursuing a PhD degree in Industrial and Manufacturing Engineering. He has an under-graduate degree from Winona State University in Composite Materials Engineering. Active in societies including SPE and SAMPE, plus campus groups Diversity in Research and Engineering of Advanced Materials (DREAM) and Retaining Engineers Through Research Entrepreneurship and Advanced-materials Training (RETREAT), Yourri stood out amongst the other candidates. In addition to reaching out beyond himself, he maintained a high GPA (3.7/4.0). Currently, his research is investigating incorporation of carbon nanotubes (CNTs) into hybrid structures as a means of decreasing the resin layer between prepreg sheets, the overall weight of the structure, and increasing the multifunctional capabilities of the composite. Yourri is also the graduate student lead for the Florida A&M University Orion Spacecraft Collaboration with Lockheed Martin.

Mr. Matthew Swift attends Iowa State University and is pursuing a degree in Materials Engineering with Polymeric Specialization. Mathew was the Composites Manager for the Prisum Solar Car Program, to compete in the World Solar Challenge to race across the continent of Australia in 2017. He has worked as a research assistant in the Soft Materials and Structures Lab as well as the Osteoceramics Lab. He has also maintained an exceptional GPA (3.97/4).

The Composite Division Award Committee is delighted to award to Mr. Yourri-Samuel Dessureault and Mr. Matthew Swift the Harold Giles Graduate Student Scholarships for 2018. The Composite Division will continue to offer the Harold Giles Scholarship to worthy candidates in the future; the scholarship was developed to honor the late Dr. Harold Giles, a past Composite Division Awards Chair. As a former University Professor at the University of North Carolina, Azdel employee, and GE employee, Harold knew full well the value of scholarships to students. He was always a proponent of awarding worthy students and served the society well in this capacity. Harold would have been pleased to know that students like Yourri-Samuel Dessureault and Matthew Swift received this award. So, if you believe that you know of other worthy candidates, please apply in 2019.

Humbly submitted by, Timothy A. Johnson

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Award Winners continued...



2018 Educator of the Year Award

he Composite Division of the Society of Plastic Engineers proudly recognizes Dr. Luyi Sun from the University of Connecticut as the 2018 Educator of the



Dr. Luyi Sun Accepting the 2018 Educator of the Year Award

Year Award. What drew the judges' attention to this candidate was the large number of students (50 undergraduate researchers / 7 Masters Students / and 12 PhD candidates) that have worked under Dr. Sun's supervision, and the work that he has done in biopolymer, nanosheet, Li ion battery, spray coating, barrier, structural, fluoropolymer coating, and photocatalytic composites areas.

We hope to honor additional, well qualified Educators on the years to come, so if you are aware of such candidates, please enter them into 2019's Educator of the Year award.

Respectfully submitted,– Dr. Dale A. Grove Composite Division Awards Chair

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SPE Council Summary



By: Dale Brosius

May 7, 2018, Orlando, FL

he SPE Council meetings were held prior to ANTEC 2018 in Orlando, Florida. At the beginning of proceedings, President Raed Al-Zubi called for a moment of silence to honor the passing of two distinguished SPE members, Dr. Costel Denson and Mr. Gautam P. Shah.

The new SPE President, Dr. Brian Grady, introduced himself to Council and presented his thoughts on the state of the Society and the role of the SPE Foundation. He shared his list of priorities which include enhancing the ANTEC experience, committing to SPE's global presence and working on membership retention. He also restated the Executive Board's commitment to full transparency. In addition, this meeting marked the arrival of newlyelected Executive Board members who were seated between Council I and II:

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SPE Council Summary continued...



- Dr. Raymond Pearson (VP Technology & Education)
- Mr. Scott Eastman (VP Sections)
- Dr. Brian Landes (President-Elect)

Clear and relevant governance is the foundation of a well-run society and SPE continues to refine its policies and by-laws. Several items were amended and approved during Council thanks to the diligent leadership of Councilor Bruce Mulholland. At the Council Committee of the Whole (CCOW). Councilor Babli Kapur was re-elected as Chair.

All presentations and data discussed during Council meetings are available on The Chain on Leadership Lane. We encourage everyone to take the time to review this information to get a full understanding of the Society.

Financial Review

The 2017 audit process is complete and there were no findings. The payment of \$1.5MM from Wiley Publishing was re-booked as deferred income over a 10-year period following advice from the auditors. There were no questions from the floor.

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Teri Chouinard CBC, APR Teri@IntuitGroup.com

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The 2018 budget was presented to Council. This budget was approved by the Finance Committee and the Executive Board in March 2018. SPE is projecting a small loss for the year. CSE Farrey reviewed the cost control measures implemented to-date, including the new technology platform and the winding down of earlier contracts. Stephanie Clark continues to generate new non-dues revenue through corporate sponsorships and other programs.

At the time of writing, ANTEC registrations were at 1500. Chief Staff Executive Pat Farrey noted that there were changes to the ANTEC pricing model in 2018 and that 180 volunteer members will attend at no charge. Farrey stated that ANTEC was projected to generate \$140k in profit.

SPE Foundation

SPE Foundation Director, Eve Vitale, reviewed the mission and board structure of the organization. She presented the new Impact Report which illustrates the good works provided by the Foundation. She reviewed the PlastiVan successes in 2017 which included the addition of several new educators. The Foundation is looking for assistance in the Western US and with more corporate sponsors. The value of scholarships and grants YTD is \$54k.

Strategic Commentary

CSE Farrey outlined his 4 priorities for the Society:

- Technology
- ANTEC
- Staffing
- Profitability

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Chicago, IL · Silicon Valley, CA · Toronto, ON Phone: (855) 4TH-STATE or (855) 484-7828 infoptna@plasmatreat.com www.plasmatreat.com Farrey presented the new SPE organizational chart and reviewed future activities and strategic actions which include establishing the Plastics For Life[™] event as a major international industry award. He echoed President Grady's commitment to SPE's global members and volunteers after having spent time in Europe, Middle East and East Asia. He stressed the importance of SPE as a peoplecentric organization.

Farrey outlined the chapter support model as it exists today. He directly addressed the conflict and differences of opinion between what SPE HQ offers, what it charges for services, and what chapters perceive as valuable. Farrey presented detailed slides on all service offerings from SPE which can be viewed on The Chain. Farrey challenged councilors to review ANTEC programming and the new website as examples of improved functionality and new capabilities delivered by SPE staff. From a business perspective, SPE currently spends \$600k on servicing small events for a return of \$300k. This is not sustainable and is a primary cause of negative operational results for the society.

Related to technology implementation, proposed changes to the member dues model were hotly debated on the Council floor. Several Section councilors objected to a proposed variable pricing dues model that would allow individual chapters to assign a dollar value to their membership. The previous model automatically included 2 group affiliations, including a geography-based Section assignment. A majority of councilors ultimately voted to revert to the previous model.

Executive Board Reports

The Executive Board members continue to work in functional roles as outlined by the new governance model. As such, individual members provided detailed updates on their

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SPE Council Summary continued...



respective portfolios. Notably, the VP of Technology & Education presented new and exciting details on a partnership with Virginia Tech on Additive Manufacturing coursework. The technical review will take place in the 2nd half of 2018 with an initial course offering scheduled for 2019. This is an excellent opportunity for SPE to create real value for members while providing important technical oversight in a dynamic area for plastics professionals.

After several rounds of fireside chats and continued dialog, Sections and Divisions still have room to maximize the value they can deliver to SPE members. The EB and Council recognize that many members do not crosspollinate and more can be done to ensure vibrant TopCons and networking opportunities. Section and Division leaders can do more to connect with each other and learn about each other's events.

The next Council meeting will be held in Charleston, SC on September 21-22. Respectfully submitted, Conor P. Carlin EB Secretary VP Marketing & Communications

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Award Winning Paper

Hybrid Cellulose-Inorganic Reinforcement Polypropylene Composites: Lightweight Materials For Automotive Applications

Mariana D. Reale Batista, Lawrence T. Drzal

Chemical Engineering and Materials Science Department, Michigan State University, East Lansing, Michigan, 48824, USA Composite Materials & Structures Center, Michigan State University, East Lansing, Michigan, 48824, USA

Alper Kiziltas, Deborah Mielewski

Sustainability and Emerging Materials, Ford Motor Company, Dearborn, Michigan, 48124, USA

Abstract

Cellulose fibers are attracting considerable attention within the transportation industry as a class of reinforcing agents for polymer composites owing to their low cost, low density, high mechanical properties, and considerable environmental benefits. The objective of this study was to develop hybrid composites combining cellulose fiber with long glass fiber, short glass fiber or talc in a polypropylene (PP) matrix to optimize the overall composite properties. Tensile, flexural and notched Izod impact tests revealed that in general the mechanical properties decreased with increasing cellulose content, however, adding an optimum concentration of the cellulose fiber is a promising alternative to reduce or replace the utilization of inorganic fibers. For applications in automotive 'under-the-hood' and body interior components, the hybrid cellulose-inorganic reinforcement composite approach not only leads to superior weight and cost savings, but also environment benefits over the inorganic reinforced composites.

Introduction

The growing environmental awareness and the demand for the utilization of renewable sources to develop sustainable and recycled materials have promoted the incorporation of cellulose fibers as reinforcement for polymer composites [1]. Cellulosic fiber reinforced polymer composites have been used for many applications such as automotive components, aerospace parts, sporting goods and in the construction industry [2]. The interest in using this material is due to its sustainable nature, low cost, acceptable mechanical properties, elimination of abrasive damage to processing equipment, abundant availability and reduced health concerns [3-7].

Cellulose fiber has a lower density compared to glass fibers and talc fillers, approximately 1.5 g/cm³ versus 2.5 and 2.8 g/cm³, respectively. Therefore, its usage in the automotive industry is a central strategy for meeting light weighting and fuel economy standards. Reducing a vehicle's weight by 10 % can improve the fuel economy by 3 to 7 % [8] and contribute to attaining the CAFE standards. Despite the attractiveness of natural fiber reinforced



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polymer composites, they exhibit lower modulus and strength as well as inferior moisture resistance compared to synthetic fiber reinforced composites, such as glass fiber reinforced polymer composites [9].

Hybridization of cellulose fiber with inorganic fibers is one possibility to improve the mechanical properties of the composites. The advantage of combining two or more fiber types in a single matrix is that the unique properties of one type of fiber could complement what is lacking in the other [10,11]. Hybridization is also a means to combine different fiber properties to develop a multifunctional composite. In this sense, hybrid composites made with cellulose fibers possess advantages in weight reduction, sustainability and higher mechanical properties.

The objective of this study was the development of hybrid composites, investigating the synergistic effects of combining cellulose fiber with long glass fiber (LGF), short glass fiber (SGF) or talc in a polypropylene (PP) matrix. The focus was on increasing the biobased content by reducing the amount of inorganic reinforcement. The mechanical, thermal and morphological properties of the resulting composites were evaluated in terms of feasibility for automotive applications to decrease the environmental impact while maintaining the product safety, durability, and quality.

Experimental

Materials

PP homopolymer pellets were supplied from three commercial sources (Company X, Company Y and Company Z) listed in Table I. Cellulose was obtained from a commercial source combined with PP in the form of pellets from two sources of production of 2014 (Cellulose A) and 2016 (Cellulose B). The inorganic reinforcements were supplied in the form of master-batch pellets and their 'as received' composition can be observed in Table I.

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Table I: As received pellets compositions

Pellets compositions					
Composition	Comments				
Homopolymer PP pellets	Kindly obtained from local sources: Neat PP X from Company X (4.0 g/10 min, ASTM D1238) Neat PP Y from Company Y (37 g/10min, ISO 1133) Neat PP Z from Company Z (17 g/10min, ISO 1133)				
30 wt. % Cellulose filled PP pellets	Kindly provided by local source: Cellulose A from 2014 Cellulose B from 2016				
30 wt. % LGF filled PP pellets	Used in instrumental panel substrates				
40 wt. % (SGF / Mica) filled PP pellets	Used in console substrate				
42.5 wt. % Talc filled PP pellets	Used in head lamp housing				
33 wt. % SGF filled PP pellets	Used in console substrate				

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Processing

The composites were prepared by injection molding (Boy Machines Model 80M) and the total fiber mass content was fixed at 30 wt. %, with the cellulose concentration varying gradually from 0 wt. % up to 30 wt. %. Table II summarizes the composite compositions and sample identification codes, for each constituent source. To achieve the desired inorganic reinforcement/cellulose concentration, the as received master-batch pellets from Table I were mixed and diluted with neat PP as needed.

Table II: Sample identification codes and composite composition for each constituent source

LGF / Cellulose hybrid composites					
Sample code	LGF (wt. %)	Cellulose A (wt. %)	PP from Company X (wt. %)		
Neat PP X	0	0	100		
LGF/Cellulose A (30/0)	30	0	70		
LGF/Cellulose A (20/10)	20	10	70		
LGF/Cellulose A (15/15)	15	15	70		
LGF/Cellulose A (10/20)	10	20	70		
LGF/Cellulose A (0/30)	0	30	70		
(SGF / Mica) / Cellulose hybrid composites					
Sample code	(SGF/Mica) (wt. %)	Cellulose B (wt. %)	PP from Company Y (wt. %)		
Neat PP Y	0	0	100		
(SGF/Mica)/Cellulose B (30/0)	30	0	70		
(SGF/Mica)/Cellulose B (20/10)	20	10	70		
(SGF/Mica)/Cellulose B (15/15)	15	15	70		
(SGF/Mica)/Cellulose B (10/20)	10	20	70		
(SGF/Mica)/Cellulose B (0/30)	0	30	70		
Talc / Cellulose hybrid composites					
Sample code	Talc (wt. %)	Cellulose B (wt. %)	PP from Company Y (wt. %)		
Neat PP Y	0	0	100		
Talc/Cellulose B (30/0)	30	0	70		
Talc/Cellulose B (20/10)	20	10	70		
Talc/Cellulose B (15/15)	15	15	70		
Talc/Cellulose B (10/20)	10	20	70		
Talc/Cellulose B (0/30)	0	30	70		

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SGF / Cellulose hybrid composites					
Sample code	SGF (wt. %)	Cellulose B (wt. %)	PP from Company Z (wt. %)		
Neat PP Z	0	0	100		
SGF/Cellulose B (30/0)	30	0	70		
SGF/Cellulose B (20/10)	20	10	70		
SGF/Cellulose B (15/15)	15	15	70		
SGF/Cellulose B (10/20)	10	20	70		
SGF/Cellulose B (0/30)	0	30	70		

Prior to injection molding, the pellets were dried at 60°C overnight to reduce moisture content. Then they were mixed and fed into the hopper on the injection molding machine. No extrusion process was used in this research. Therefore, it is worth mentioning that it consists of an efficient one-step injection process that can speed up part production. To avoid cellulose degradation the maximum injection temperature was limited to 193°C (380 F). Composites were molded into ASTM test specimens and conditioned in a room at $23 \pm 2^{\circ}$ C and $50 \pm 5\%$ relative humidity for 7 days before conducting mechanical tests.



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Testing procedure and characterization

Mechanical test

Tensile and flexural tests were conducted in an Instron 3366 in general compliance to ASTM D638-10 and ASTM D790-10, respectively. The tensile test used a crosshead speed of 5.0 mm/min, a 5kN load cell and a 50 mm extensometer was attached to the gauge section of the specimen to measure the displacement. The stress at maximum load, strain at maximum load and Young's modulus were determined. At least six specimens were tested for each data set.

The flexural tests also used a 5kN load cell, at a rate of 1.0 mm/min and a support span length of 50 mm. The stress at 5% strain and flexural modulus were determined.

The impact strength of notched Izod specimens was measured according to ASTM D256-10. Tests were conducted on a Testing Machines Inc. 43-02-03 model impact test machine with a 2 lb. pendulum, and the results are the average from ten specimens of each data set.

All the mechanical tests were run in an environmentally conditioned room at 23 ± 2 °C and $50 \pm 5\%$ relative humidity.

Thermal characterization

Melting and crystallization behavior of the neat polymer matrix and the composites were measured using a TA Instruments Q2000 differential scanning calorimetry (DSC). To remove the thermal history, samples were heated from room temperature to 250 °C at a rate of 20 °C/min and held isothermally at 250 °C for 5 minutes. Then, they were cooled to -50 °C at a rate of 10 °C/ min and held at -50 °C for 5 minutes before reheating to 250 °C at a rate of 10 °C/min. The melting and crystallization behavior were collected from the heat flow versus temperature curves. Melting temperature (T_m) was assigned as the peak minimum of the endothermic melt transition, and crystallization temperature (T_c) as the peak maximum of the exothermic crystallization transition. The enthalpy of melting (Δ Hm) and enthalpy of crystallization (Δ Hc) were also measured from these curves. Specimen weight was in the range of 7 to 9 mg and the results were averaged from three specimens of each sample.

The thermal stability of the neat PP and the composites was investigated by thermal gravimetric analysis (TGA) curves using a TA Instruments Q500. The thermograms were obtained under constant air flow rate of 50 mL·min⁻¹ and the samples were heated up to 600 °C at a rate of 10 °C/min. The sample weight was in the range of 8 to 10 mg. Three replicates of each sample were performed.

Scanning electron microscopy (SEM)

A Carl Zeiss EVO LS 25 scanning electron microscope with accelerating voltage of 15 keV was used to observe the morphology of the impact fracture surface of the samples. The samples were sputter-coated with platinum to prevent surface charging.

Results and discussion

Mechanical properties

Tensile properties

Figures 1, 2 and 3 show the effects of different fiber combinations on the tensile properties of the composites, along with the neat PP. Overall, the addition of inorganic fibers and cellulose led to a considerable increase in the maximum tensile stress (Figure 1) and Young's modulus (Figure 2) in comparison to the neat PP. The inorganic fibers produced the greatest

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improvement in these properties. An increase of 232% and 401% for tensile stress and modulus, respectively, was observed for LGF/Cellulose A (30/0) composites compared to neat PP X (Figures 1a, 2a). As the cellulose content was increased to replace a portion of the LGF, SGF/ Mica or SGF, the tensile stress and modulus of the composites decreased. For these composites which contain glass fiber, this trend is expected due to the stronger and stiffer properties of the inorganic fiber compared to cellulose [12]. As reported in the work of Thwe and Liao [9], the tensile strength and modulus of hybrid composites of bamboo fiber and glass fiber in PP gradually increased with increasing glass fiber to bamboo fiber ratio.

Note that even with the incorporation of 30 wt. % cellulose, the tensile stress and modulus of the composites are still higher compared to

neat PP. The addition of 30 wt. % Cellulose B promotes an increase of 18% and 129% for tensile stress and modulus, respectively, in comparison with neat PP Y (Figures 1b, 1c, 2b, 2c). SGF/Cellulose B (15/15) hybrid composites increased the tensile stress and modulus by 58% and 171%, respectively, in comparison with neat PP Z (Figures 1d, 2d) and is suitable for body interior (console substrate, wiring harness) and under-the-hood (battery and power distribution box covers) applications.

The tensile stress at maximum load for the composites made of talc and cellulose did not follow the previous trend (Figure 1c). Talc/Cellulose B (10/20) composites exhibited an increase of 15% for the tensile stress in comparison with Talc/ Cellulose B (30/0) composites, which represents an advantage of the hybrid composite over the composite reinforced with talc only.

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The tensile strains at maximum load of the composites were markedly lower than neat PP (Figure 3) with no considerable difference in strain as a function of fiber content.

Overall, the mechanical properties in tension show that the hybridization of cellulose with inorganic reinforcements produced beneficial enhancements.

Flexural properties

Figures 4 and 5 show the flexural properties of the polymer matrix and the composites. Flexural properties exhibited a similar trend as the tensile results, revealing that in general the mechanical properties decreased as the amount of cellulose increased. The stress at 5% strain achieved a 59% enhancement for LGF/Cellulose A (0/30) compared to neat PP X (Figure 4a).

The presence of cellulose increased the stress at 5% strain for Talc/Cellulose B composites (Figure 4c) with the highest strength of 56.3 MPa for Talc/Cellulose B (10/20) hybrid composites. It represents an increase of 10% in comparison with Talc/Cellulose B (30/0) composites.

Similar to the tensile modulus results, all composites showed higher flexural modulus than neat PP (Figure 5).



Figure 1: Tensile stress at maximum load of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.



Figure 2: Young's modulus of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/ Cellulose B and d) SGF/Cellulose B composites along with neat PP.



Figure 3: Tensile strain at maximum load of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.



Figure 4: Stress at 5% strain of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/ Cellulose B and d) SGF/Cellulose B composites along with neat PP. *Specimens broke before 5% strain; maximum stress value used.

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Impact properties

The addition of cellulose fibers acted as stress concentrators and reduced the impact strength of the composites, as shown in Figures 6a, 6b and 6d. A decrease in impact strength of PP was also observed when sisal fibers were added to PP by unnotched Izod impact test [13].



Figure 5: Flexural modulus of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/ Cellulose B and d) SGF/Cellulose B composites along with neat PP.

Figure 6a shows that LGF/Cellulose A (15/15) composites exhibited an impact strength 91% higher than LGF/Cellulose A (0/30). As reported by Panthapulakkal and Sain [14] for hemp/ glass fiber/PP composites, the impact strength is enhanced with an increase in glass fiber content due to the improved resistance offered from the glass fibers in the composites. For the hybrid composites made of SGF/Cellulose B (Figure 6d) the impact strength is still preserved in comparison with the neat PP Z. The SGF/Cellulose B (20/10) composites increased the impact strength by 34% and the SGF/Cellulose B (10/20) composites still exhibited 16% increase compared to neat PP Z. No change in impact strength was observed when adding cellulose to talc composites (Figure 6c).

Morphological properties

Figures 7, 8, 9 and 10 show the impact fracture surface of LGF/Cellulose A, (SGF/Mica)/ Cellulose B, Talc/Cellulose B and SGF/Cellulose B composites, respectively, along with the neat PP. In the micrographs, the direction of impact is from right to left. The samples investigated are neat PP and composites reinforced with inorganic reinforcement/cellulose at 30/0, 15/15 and 0/30 (wt.%).



Figure 6: Impact strength of a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.

It can be observed that the basic deformation mechanism of all unmodified PP is shear yielding (Figures 7a, 8a, 9a, 10a). There was no significant difference between the surface morphologies of the Cellulose A and Cellulose B composites (Figures 7d, 8d, 9d, 10d). Since the cellulose fiber is hydrophilic the micrographs do not show evidence of interfacial interaction between the cellulose and the matrix, and this incompatibility may reduce the mechanical properties. This observation is in agreement with the reduced tensile, flexural and impact properties measured on the cellulose reinforced composites. Cellulose fiber fracture is also clear from Figure 7d.

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The micrographs of specimens reinforced with glass fiber (long or short) exhibited fiber pull out (Figures 7b, 8b, 10b) that indicates poor interfacial adhesion between the glass fiber and matrix. This behavior is in agreement with results reported by Arbelaiz et al. [11]. The distribution of the fibers in the matrix appears to be good however.

With the incorporation of 15 wt. % cellulose (Figures 7c, 8c, 10c), holes from the pulled out glass fibers are still observed. In the SEM micrographs glass fiber can be distinguished from cellulose by their difference in diameter. Good distribution of the hybrid fibers is observed.



Figure 7: SEM micrographs of a) Neat PP X, b) LGF/Cellulose A (30/0), c) LGF/Cellulose A (15/15) and d) LGF/Cellulose A (0/30).



Figure 8: SEM micrographs of a) Neat PP Y, b) (SGF/Mica)/Cellulose B (30/0), c) (SGF/ Mica)/Cellulose B (15/15) and d) (SGF/Mica)/ Cellulose B (0/30).



Figure 9: SEM micrographs of a) Neat PP Y, b) Talc/Cellulose B (30/0), c) Talc/Cellulose B (15/15) and d) Talc/Cellulose B (0/30).

Figure 8b identifies a mica particle and Figure 9b identifies the platelet talc filler. Micrographs show good talc distribution in the matrix with no large aggregates present.

Overall, the composites show good fiber distribution, confirming that the one step injection process with no compatibilizer successfully blended the hybrid materials in the PP. It shows the advantage of using these pre-compounded pellets to speed up the process production.

Thermal properties

TGA

Figure 11 summarizes the thermal stability of the composites, where Td is assigned as the temperature at the maximum rate of decomposition determined from the derivative thermogravimetric (DTG) curves. All composites are more stable than the neat PP and the Td of the inorganic reinforced composites manifested the greatest increases. The incorporation of 30 wt. % cellulose also increased the thermal stability relative to PP but to a lesser degree. The inorganic reinforcement/cellulose composites reinforced at 20/10, 15/15 and 10/20 (wt. %) have similar thermal stability. This provides an opportunity for hybrid composites made of cellulose to be considered for challenging conditions on under-thehood applications.

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Figure 10: SEM micrographs of a) Neat PP Z, b) SGF/Cellulose B (30/0), c) SGF/Cellulose B (15/15) and d) SGF/Cellulose B (0/30).

Thermal stability was also defined by the temperature at which 1% and 10% weight loss occurred, T_1 and T_{10} , respectively, as shown in Figures 12 and 13. All hybrid composites are more thermally stable than the neat PP. The composites that contain cellulose possess lower T_1 and T_{10} compared to the composites with inorganic reinforcements only. However, this behavior is not observed for LGF/Cellulose A composites (Figures 12a, 13a).

Figure 14 shows the residual weight percent at 587 °C. The residue decreased with cellulose content at this temperature and the residue is mainly from the inorganic reinforcements.

DSC

Figure 15 summarizes the T_c and T_m of PP and the composites. It indicates an increase in the T_c for all composites compared to neat PP, which reveals that the fibers act as nucleating agents leading to a faster crystallization of PP matrix. This is a beneficial development which can increase the rate of part production. The DSC measurements indicate that the addition of cellulose had minimal effect on the melting temperatures.

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Figure 11: Temperature at the maximum rate of decomposition determined from DTG curves for a) LGF/Cellulose A, b) (SGF/Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.



Figure 12: Temperature at 1% weight loss from TGA curves for a) LGF/Cellulose A, b) (SGF/ Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.



Figure 13: Temperature at 10% weight loss from TGA curves for a) LGF/Cellulose A, b) (SGF/ Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.

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Figure 14: Residual weight percent at 587 °C from TGA curves for a) LGF/Cellulose A, b) (SGF/ Mica)/Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites along with neat PP.

The ΔH_c and ΔH_m decreased with the incorporation of fibers (Table III), with no significant dependence on fiber loading. This trend is in agreement with the results of Huda et al. [15], where a reduction of the melting and crystallization enthalpies of the composites was observed with the addition of recycled newspaper cellulose fibers and talc compared to neat PP.

Composites reinforced with 30 wt. % cellulose have, marginally, the lowest enthalpies. It may be that cellulose was hindering polymer chain movement during heat cycling, as observed by Langhorst et al. [8].

Summary and Next Steps

In this study, we investigated the hybridization of cellulose fiber with several inorganic fibers to reinforce a PP matrix for automotive applications. Composites were processed by injec-



Figure 15: Crystallization and melting temperatures for a) LGF/Cellulose A, b) (SGF/Mica)/ Cellulose B, c) Talc/Cellulose B and d) SGF/Cellulose B composites.

tion molding with total reinforcement content kept at 30 wt. %. Overall, it was found that the hybrid fibers acted as an effective reinforcement. Good synergetic effects on the mechanical, thermal and morphological properties of the hybrid composites were observed.

Results from tensile, flexural and notched Izod impact tests show that in general the mechanical properties decreased with increasing cellulose content. However, composites with an optimum amount of cellulose fiber may reduce or replace a portion of the inorganic reinforcements in many applications. LGF/Cellulose A composites exhibited the best mechanical properties. However, when comparing thermal properties, (SGF/ Mica)/Cellulose B, Talc/Cellulose B and SGF/ Cellulose B composites exhibited similar or even superior properties.

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Composition	LGF/Ce	ellulose A	(SGF/Mica)/Cellulose B		Talc/Cellulose B		SGF/Cellulose B	
(wt. %)	$\Delta H_c (J/g)$	$\Delta H_m (J/g)$	$\Delta H_c (J/g)$	$\Delta H_m (J/g)$	$\Delta H_c (J/g)$	$\Delta H_m (J/g)$	$\Delta H_{c} (J/g)$	$\Delta H_m (J/g)$
Neat PP X	89.8 ± 3.3	86.4 ± 3.6	-	-	-	-	-	-
Neat PP Y	-	-	96.7 ± 0.2	95.8 ± 0.3	96.7 ± 0.2	95.8 ± 0.3	-	-
Neat PP Z	-	-	-	-	-	-	104.6 ± 2.9	104.0 ± 5.5
(30/0)	75.1 ± 0.6	75.7 ± 1.6	67.8 ± 1.8	67.3 ± 0.8	70.1 ± 1.4	69.9 ± 1.5	73.6 ± 1.6	72.0 ± 0.2
(20/10)	73.5 ± 1.5	73.4 ± 3.4	68.7 ± 3.1	67.8 ± 3.3	71.6 ± 1.7	68.6 ± 0.3	69.4 ± 3.0	67.9 ± 0.3
(15/15)	74.3 ± 3.8	63.6 ± 6.7	69.2 ± 2.8	66.8 ± 1.4	69.2 ± 2.7	64.2 ± 2.6	70.6 ± 3.8	71.3 ± 3.2
(10/20)	70.7 ± 3.2	68.1 ± 3.1	66.9 ± 2.8	68.0 ± 2.6	70.6 ± 0.7	68.0 ± 1.4	73.3 ± 1.0	71.6 ± 2.1
(0/30)	67.6 ± 3.3	63.0 ± 0.6	62.3 ± 5.0	62.5 ± 5.3	62.3 ± 5.0	62.5 ± 5.3	62.3 ± 5.0	62.5 ± 5.3
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Table III: Enthalpies of crystallization and melting for neat PP and composites

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The T_c for all composites increased in comparison to neat PP, revealing the fibers ability to act as nucleating agents and speed part production. SEM showed good fiber distribution, however due to the inherent lack of compatibility between cellulosic fibers and polymeric matrices such as PP, the use of a compatibilizer is recommended as future work to improve the adhesion of the fiber to the PP matrix leading to even greater mechanical and thermal performance of the hybrid composites.

Cellulose fiber polymer composite properties can be improved by the hybridization with inorganic reinforcements. This work shows that hybridization of cellulose fiber with inorganic reinforcements in polypropylene composites has the potential to reduce the use of inorganic fibers in different automotive applications, leading to weight and cost savings, and contributing to sustainability of composites.

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