Abstract
This article addresses the effect of Na+-Montmorillonite (MMT) on cryogenic mechanical properties of polypropylene (PP). Polypropylene/Montmorillonite (PP/MMT) nanocomposites were prepared by using twin-screw extruder incorporating the polypropylene grafted maleic anhydride (PP-g-mA) as compatibilizer is used for better dispersion of nanoclay in the polymer matrix. The dispersion and exfoliation of MMT in the nanocomposites were observed using transmission electron microscopy (TEM) and X-ray diffraction (X-RD). Mechanical behaviors of PP/MMT nanocomposites at both room temperature (RT) and liquid nitrogen temperature (77 K) were investigated in terms of tensile, flexural and impact properties. The mechanical results showed that the tensile strength of PP/MMT nanocomposites at both RT and 77 K reached the maximum at the 5wt. % MMT content, increased by 3.1% and 13.2%, respectively, as compared with those of pure PP sample. The notched impact strength at RT increased from 22.33 J/m for the pure PP matrix to 36.53 J/m for the nanocomposite with the 5wt% MMT content. The fracture surfaces of neat PP and PP/MMT nanocomposites were examined using Scanning Electron Microscopy (SEM). Finally, the dependence of the storage modulus and loss modulus of the MMT content was examined by Dynamic Mechanical Analyzer (DMA) at cryogenic temperature.

Keywords: Cryogenic Temperature, Mechanical Properties, Nanocomposites, Polymer

1. Introduction
The mechanical and thermal properties of polymers and composite structures can be altered through the use of various kinds of fillers. The dimensions of these fillers typically fall on a macroscopic (1 µm–1 nm) length scale. Fillers of this type increase the stiffness and heat distortion temperature of a polymer, primarily because the filler makes up a significant proportion of the total mass. However, macroscopic fillers usually cause decreases in strength, impact resistance, and processability. A new area of composites research has emerged in the last two decades that utilizes nanoparticle fillers to alter the properties of polymers. Definitive results have not yet been achieved, but trends show that when processed properly small amounts (≤5 wt. %) of nanoparticle fillers can increase the modulus, strength, toughness, resistance to chemical attack, gas impermeability, resistance to thermal degradation, and dimensional stability of polymeric materials. However, the majority of this work has focused on neat thermoplastics and it is only recently that thermosets have begun to be examined as potential matrices for nanocomposites. With the rapid development of the space industry and superconductive technologies, polymer matrix and their composites have also been widely used in these fields. Successful applications of these materials highly depend on their performances at extreme low temperature (such as liquid nitrogen and liquid helium temperature), particularly for reusable launch vehicles. Previous studies have shown that MMT can effectively improve properties of polymer matrix at room temperature (RT). It is expected that it may also improve these properties at cryogenic temperature. Generally speaking, cryogenic properties of polymer matrix can be significantly different from those at RT because of the effect of large thermal contraction. The property improvement of PP/MMT nanocomposites cannot be simply inferred from the
results of the property improvement of PP/MMT nanocomposites obtained at RT. To ensure the safety of the applications of these materials at the cryogenic temperature, it is indispensable to study cryogenic mechanical properties of PP/MMT nanocomposites. To our best of knowledge, no work has been reported on the cryogenic mechanical properties of PP/MMT nanocomposites. Therefore, in this paper, PP/MMT nanocomposites were prepared and their mechanical properties at both RT and 77 K were experimentally investigated and compared.

2. Experimental

2.1 Materials

The PP has grade of H110MA with density of 0.910g/cc and melt flow index (MFI) of 11g/10 min, purchased from reliance Ltd, India. The coupling agent PP-g-MA purchased from Exxon mobile India Pvt. Ltd, India, with the grade name Exxlor PO 1020, with MFI 125gm/10 min with 0.75% of grafting. Nanoclay i.e. unmodified MMT having CEC 92.6 meq/100g, purchased from southern clay products Inc, USA.

2.2 Preparation of PP/MMT Nanocomposites

Melt mixing of PP, PP-g-MAH (10 wt. %) and unmodified MMT of 1, 3, 5 and 7wt. % carried out in counter rotating twin screw extruder. Prior to melt mixing, the PP and the nanofillers were preheated in a vacuum oven at 60oC for a time of 6 hours. PP was fed into the twin screw extruder at the rate of 5 kg/hour and the nanofillers were introduced subsequently at the melting zone. The process of manufacturing of PP/MMT nanocomposites were carried out at a two screw speed of 150 rpm and maintain the temperature difference of barrel in twin screw extruder were 160, 170 and 180°C between feed zones to die zone, followed by granulation in a pelletizer and drying. These pellets were injection molded by using injection moulding machine. The manufacturing of test specimens of tensile, flexural and impact strength as per ASTMD by using injection moulding machine with clamping force 800K fitted with dehumidifier at a temperature range of 195-220°C.

2.3 Characterization Methods

2.3.1 X-ray Diffraction

2.3.2 SEM and TEM

The SEM images of the PP and PP/MMT nanocomposites, the fractured surfaces were coated thin layers of gold about 10 Ao. All SEM images were taken from JEOL, JSM 840A scanning electron microscope with an accelerating voltage of 10kv. The TEM images of unmodified MMT and PP/MMT nanocomposites, the thin sections ranging from 70-100nm in thickness were cut with a diamond knife using Reichert-Jung FCH ultra cryomicrotome cutter. The all TEM images were taken from a JEOL-EM-2000 FX Electron Microscope with 200 kV accelerating voltage.

2.3.3 Mechanical Testing

The tensile samples were prepared according to the recommendation of ASTM D638. The tensile properties of PP/MMT nanocomposites at RT and 77 K were measured by a using universal testing machine (UTM) LR-100K (Lloyd Instrument Ltd U.K.) with a 10 kN load cell, with a crosshead speed of 2 mm/min. The cryogenic temperature condition was achieved by dipping clamps and samples in a liquid nitrogen filled cryostat designed in our laboratory. The entire test was conducted while the sample and its loading fixture were submerged in liquid nitrogen. A schematic diagram of the loading fixture is shown in Figure 1. The flexural samples were prepared according to the recommendation of ASTM D-790. The flexural properties of PP/MMT nanocomposites at RT and 77 K were
measured by using Universal Testing Machine (UTM) LR-100K (Lloyd Instrument Ltd. UK), with a cross head speed of 1.3 mm/min and a span length of 50 mm. The cryogenic temperature condition was achieved by dipping clamps and samples in a liquid nitrogen filled cryostat designed in our laboratory. The entire test was conducted while the sample and its loading fixture were submerged in liquid nitrogen. A schematic diagram of the loading fixture is shown in Figure 2. The notched impact strength was measured with a Reger Impact Tester according to ASTM-D256. Impact testing at 77 K was performed with the samples dipped in a liquid nitrogen filled cryostat for over 10 min and was completed in a couple of seconds after taking the specimens out from the cryostat. At least five specimens were tested for each composition. The temperature inside the samples was examined to have no obvious change after the samples were taken out from the cryostat for a few seconds using cryogenic thermocouples embedded inside the samples. Therefore, the temperature for the impact testing was approximately considered to be 77 K.

2.3.4 Dynamic Mechanical

Viscoelastic properties such as storage modulus and loss modulus as a function of temperature were measured in a dynamic mechanical thermal analyzer (Netzsch DMA 242-C) with the temperature range from 200°C-150°C. The frequency used was 1 Hz and the heating rate, 10°C/min. The specimen dimension was 60X10X3 mm.

3. Result and Discussion

3.1 X-ray Diffraction Analysis

Figure 3 shows the XRD curve of unmodified MMT clay. In this Figure 3, the peak shifts from 2θ = 6.98°, corresponding to basal d-spacing 12.6 Å. XRD curves of PP and PP/MMT nanocomposites as shown in Figure 4 with the help of coupling agent the polymer intercalated between the intercalary spacing of layered silicates and the clay layers to move apart. The absence of peak showed the separation of clay layers and the formation of nanocomposites at 1 wt. % to 5 wt. % nanofiller reinforcement. This indication by TEM studies that intercalated structures have formed in 1 wt.% and 3 wt.% of nanofiller in PP/MMT nanocomposites. PP infused with 5 wt. % of MMT has exfoliated nanocomposites structures.
Cryogenic Mechanical Properties of PP/MMT Polymer Nanocomposites

For the ease of comparison, TEM morphologies of the raw MMT and PP/MMT nanocomposites (1 wt%, 3 wt%, 5 wt% and 7 wt%) are presented in Figure 5a-e. The Figure 5a-e showed that the size of unmodified MMT clay particles is about 20 nm thick and 200–500 nm length. The Figure 5a-c showed the nanofiller has been further intercalated and exfoliated in the PP/MMT nanocomposites with the nanofiller reinforcement of 1 wt%, 3 wt% and 5 wt%. From this result observed macromolecules of polymers formed the interlayer of the unmodified MMT during melt mixing process. The reason is that netlike macromolecules formed the interlayer of the MMT during the melting process. The d-spacing of unmodified MMT nanoclay is increased, which is consistent with the XRD curves (Figure 5c). However, some of the MMT actually consists of several layers, indicating that they were not fully exfoliated to the single layer. It is probably because they were not organically modified due to the charge of the pristine clay layer varying from layer to layer. When the content of MMT in composite was increased to 7 wt%, the conglomerating probability of MMT and the quantity of non-exfoliated MMT inside PP matrix increased. Therefore, it can be seen in Figure 5d that serious aggregations occurred everywhere at the MMT content of 7 wt%. Similar results can also be drawn from Figure 5d.

Figure 3. XRD curves of PP and PP/MMT composites.

Figure 4. XRD curves of PP and PP/MMT composites.

3.2 TEM Analysis

For the ease of comparison, TEM morphologies of the raw MMT and PP/MMT nanocomposites (1 wt%, 3 wt%, 5 wt% and 7 wt%) are presented in Figure 5a–e. The Figure 5a-e showed that the size of unmodified MMT clay particles is about 20 nm thick and 200–500 nm length. The Figure 5a-c showed the nanofiller has been further intercalated and exfoliated in the PP/MMT nanocomposites with the nanofiller reinforcement of 1 wt%, 3 wt% and 5 wt%. From this result observed macromolecules of polymers formed the interlayer of the unmodified MMT during melt mixing process. The reason is that netlike macromolecules formed the interlayer of the MMT during the melting process. The d-spacing of unmodified MMT nanoclay is increased, which is consistent with the XRD curves (Figure 5c). However, some of the MMT actually consists of several layers, indicating that they were not fully exfoliated to the single layer. It is probably because they were not organically modified due to the charge of the pristine clay layer varying from layer to layer. When the content of MMT in composite was increased to 7 wt%, the conglomerating probability of MMT and the quantity of non-exfoliated MMT inside PP matrix increased. Therefore, it can be seen in Figure 5d that serious aggregations occurred everywhere at the MMT content of 7 wt%. Similar results can also be drawn from Figure 5d.

Figure 5. TEM micrograph of PP/MMT composites a)1 wt%, b) 3 wt%, c) 5 wt%, d) 7 wt% and e) raw MMT.
3.3 Mechanical Properties

3.3.1 Tensile and Flexural Properties

Figure 6 plots the tensile strength of the samples at RT and 77 K. The unmodified MMT nanofiller were reinforced into the polymer at different weight percentage from zero to 7 wt%. The results showed that nanofiller reinforcement into the polymer has marginal influence on the tensile strength of PP/MMT nanocomposites. The nanofiller at 5 wt% increased the tensile by 3.1% and 13.2%, at both RT and 77 K, when compared with pure PP. Figure 7 plots the flexural strength of the samples at RT and 77 K. The MMT nanoclay was reinforced into the polymer at different weight percentage from zero to 7 wt%. The results showed that nanofiller reinforcement has significant influence on the flexural strength of PP/MMT composites. As the content of MMT increases, the flexural strength at both RT and 77 K reaches maximum at the MMT content 3 wt%, and increased by 11.5% and 21.2%, respectively, when compared with those of pure PP samples. It can be seen from Figure 4b-c that the intercalated and exfoliated MMT layers served as the short fasciculus inside the composite. The addition of short fasciculus to the PP matrix could reinforce the PP matrix and contributed to the increase of the tensile and flexural strength of the PP/MMT composite at the 5 wt% and 3 wt% MMT content. The tensile and flexural strength at 77 K was generally higher than that at RT at the same content of MMT. The strength of PP matrix becomes higher at 77 K and the interfacial adhesion becomes stronger at 77 K than at room temperature, leading to the higher strength. The thermal shrinkage caused by the big differences in the thermal expansion coefficients of PP and MMT would result in stronger interfacial adhesion strength between MMT and PP matrix at 77 K than at RT. On the other hand, the coefficient of thermal expansion (CTE) for MMT at RT was reported to be about 0.5x10^-5 K^-1. And the CTE for MMT at cryogenic temperature should be somewhat lower than this value because thermal expansion coefficient is closely related to thermal oscillation while intensity of thermal oscillation becomes weaker as temperature decreases, leading to a lower CTE. As a consequence, the ‘short fasciculus’ were clamped more tightly by the PP matrix at 77 K than at RT, resulting in higher tensile and flexural strength at 77 K than that at RT. The tensile strength at RT and 77 K decreased when the MMT content was further increased to 5 wt% and 7 wt%. This might be mainly attributed to the fact that the percentage of aggregated (namely non-exfoliated) MMT was higher at the higher MMT content (see Figure 3d). The micro-sized holes to be shown in Figure 8b were indicative of pull-out of aggregated MMT particles. When the composite samples were tested under tensile loading, cracks might initiate at the aggregated MMT particles and propagated throughout the samples. Therefore, the tensile strength was lower at the higher MMT concentration.

![Figure 6](image6.png)

**Figure 6.** Effect of MMT content on tensile strength of PP/MMT composites at RT and 77 K.

![Figure 7](image7.png)

**Figure 7.** Effect of MMT content on flexural strength of PP/MMT composites at RT and 77 K.
Figure 8 plots the tensile modulus of the nanocomposites at RT and 77 K. The tensile modulus of PP and PP/MMT nanocomposites at RT and 77 K was observed to increase almost linearly as the amount of nanofiller up to 5 wt% increased by 40.8% and 68.6%, respectively, when compared with those of pure PP samples. Figure 9 plots the flexural modulus of the PP and PP/MMT composites, respectively, when compared with those of pure PP samples. This is mainly because MMT has a higher modulus than PP matrix. Moreover, Figure 8 shows that the modulus at 77 K was much higher than that at RT at the same MMT content. This is because on the one hand, the modulus of the PP matrix at 77 K was higher than that at RT (see Figure 5 at zero MMT content) due to the tight arrangement of polymer molecules. On the other hand, any material would show higher modulus at lower temperature, so the modulus of MMT would also be higher at 77 K than at room temperature, contributing to the higher modulus of the composites at 77 K than at RT.

### 3.3.2 Notched Impact Strength

Figure 10 plots the impact strength of the nanocomposites at RT and 77 K as a function of MMT content. The notched impact strength at RT increased from 22.33 J/m of the pure PP to 36.53 J/m by the incorporation of MMT particles for the nanocomposite containing the 5 wt% MMT content while decreased for the composites containing the 7 wt% MMT content.

Toughness improvement of nanocomposites can be attained when MMT in pure PP can resist the propagation of cracks at the 5 wt% MMT content at RT as indicated by the increase of notched impact strength. The rougher surfaces of the nanocomposites in Figure 5 c-d than those of pure PP matrix in Figure 5 a-b are indicative of the improvement in toughness. Moreover, the notched impact strength at 77 K was shown to be lower than that at RT except for the pure PP matrix. The sharp decrease of the notched impact strength at 77 K for the cases of all PP/MMT nanocomposites may be caused by the thermal stresses due to the different contractive coefficients between the organic and inorganic phases when cooled down to 77 K. The increase of the notched impact strength at 77 K for the pure PP matrix may be attributed to tight arrangement of molecules and no different contractive coefficients in the homogeneous phase.
3.3 SEM Analysis

The fracture surfaces of the nanocomposites were observed using SEM. Figure 11a-j displays some representative SEM images of the fracture surfaces of the notched impact samples at RT and 77 K. Figure 11 shows that the surfaces of the pure PP samples were smooth and featureless, representing brittle failure of homogeneous materials. As shown in the SEM images for the PP/MMT nanocomposite samples with the 1 wt%, 3 wt%, 5 wt%, and 7 wt% MMT content, the fracture surfaces were significantly different from those of pure PP matrix samples. The fracture surfaces were broken into small and rough fracture pieces, contributing to improvement of the toughness of the nanocomposites. However, the improved toughness by the rough fracture surfaces for the cases of 1 wt%, 3 wt%, and 5 wt% would be overwhelmed by the effect of aggregation of MMT particles since micro cracks would relatively easily initiate at the micro sized aggregated MMT particles, resulting in a reduced macroscopic toughness as shown in Figure 10. On the other hand, 77 K were smoother than those at RT at the same content of MMT except for the pure PP matrix, which represents more brittle fracture of the nanocomposites at 77 K.

3.4 Dynamic Mechanical Analysis

Figure 12 plots the storage modulus of the PP and PP/MMT nanocomposites at RT and 77 K. The results showed that unmodified MMT clay nanofiller have marginal influence on the storage modulus of PP/MMT respectively, when compared with those of pure PP samples. Figure 13 plots the loss modulus of the samples at RT and 77 K as a function of weight percentage of MMT from zero to 7 wt%. The results show that MMT content has no influence on the loss modulus of PP/MMT composites. As the content of MMT increases, the loss modulus at both RT and 77 K reaches minimum at the MMT content up to 5 wt%, and decreased by 3.1% and 13.2%, respectively, when compared with those of pure PP samples.

4. Conclusion

The cryogenic mechanical properties and dynamic mechanical analysis of PP/MMT nanocomposites prepared by adding the MMT nano filler content to pure PP matrix have been studied. XRD and TEM results showed that MMT can be exfoliated at the 5 wt% MMT content.
22.33 J/m for the pure PP matrix to 36.53 J/m for the nanocomposite with the 5 wt% MMT content. It was also observed that the notched impact strength at 77 K was lower than that at RT except for the pure PP matrix. The storage modulus of PP/MMT nanocomposites at both RT and 77 K reaches maximum at the MMT content up to 3 wt%, and the loss modulus of PP/MMT nanocomposites at both RT and 77 K reaches minimum at the MMT content up to 5 wt%.

5. References