Influence of SLES-layered double hydroxides on the mechanical and biodegradation properties of poly (lactic acid) nanocomposites

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A B S T R A C T
Petroleum based polymers have great attention due to its excellent properties. Unfortunately, its uses give detrimental impacts to the health of our environment. Poly (lactic acid), PLA is recognised as one of the potential biodegradable polymers as it exhibits almost similar properties to non-biodegradable polymers. In this report, nanocomposites composed of PLA and sodium lauryl ether sulphate modified layered double hydroxides, SLES-LDH were prepared by solvent casting method to enhance PLA tensile strength. The pristine LDH used was synthesised by a co-precipitation method and then modified by surfactant via an ion exchange process. X-ray diffraction result showed LDH interlayer spacing increased from 8.10Å to 34.23Å indicated intercalation of the surfactant molecules into LDH was successfully carried out. The result of tensile strength test showed the addition of 1 wt% of SLES-LDH in the PLA improved 26% of its tensile strength compared to that of the pure sample. To study the presence of SLES-LDH on the product biodegradability, burial test was carried out. It reveals that PLA nanocomposites degrade faster than the starting polymer and the degradation is SLES-LDH content dependent.

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1. Introduction
Most commonly used polymers such as polyethylene terephthalate, polypropylene and polystyrene, are petroleum based and non-biodegradable materials. Therefore, use of these products should be avoided or minimised in order to reduce their impact to environment deterioration. Consequently, biodegradable polymers received a lot of attention and they are good alternatives for the petroleum based materials. Poly (lactic acid), PLA is a thermoplastic polymer which is biodegradable and can be produced by many renewable resources (Anderson et al., 2008) such as corn, potato and sugar beet. Therefore, it is an attractive alternative to conventional petrochemical-based polymers. This polymer degrades naturally and completely into water and carbon dioxide without harmful products (Bordes et al., 2009). In addition, PLA has similar properties with non-biodegradable polymers. However its high stiffness and brittleness have to be modified to allow it to be used more applications. One of the approaches to modify the PLA properties to make it more useful industrial applications is to blend it with stearic acid modified LDH (Mahboobeh et al., 2010; Mahboobeh et al., 2012). LDH is anionic clay which consists of positively charged metal hydroxide sheets with exchangeable anions and water molecules. The general composition of LDH can be represented by \[ M_1-x^2 + M_2^{2+}[(OH)x]_2 + A^{n+}/mH_2O \], where \( M^{2+} \) and \( M^{3+} \) are divalent and trivalent cations, and \( A^a \) is an exchangeable anion. This compound has been given much attention recently because it is facile, easy to prepare, as well as environmental friendly (Williams and O’Hare, 2006; Katiyar et al., 2010; Mahboobeh et al., 2012; Livi et al., 2012). Hydrotalcite is naturally occurred LDH (Hochstetter, 1842) but its synthesis was attempted since 1942 (Feitknecht and Gerber, 1942). Modification of LDH by an ion exchange process facilitates intercalation of bulky organic anions into its layers, resulting in enhancement of its hydrophobicity property. It is reported that anionic surfactants with at least one anionic end group and
long hydrophobic tail are the best compounds to be intercalated resulting an increase of interlayer spacing of LDH which then accommodates intercalation of large species (Maiti et al., 2008; Zhao et al., 2003-2009). Other approaches include modification by ionic liquids method (Livi et al., 2012) and grafted TiO₂ particles (Luo et al., 2009). In this report we describe the results of our attempt to modify LDH by SLES. The product of this modification is then used as an additive for PLA nanocomposites preparation.

2. Materials and methods

2.1. Materials

Magnesium nitrate hexahydrate and aluminium nitrate nonahydrate (Merck, Germany) were used to synthesis pristine LDH. PLA (Bio-Polymer 3052D) was supplied by NatureWorks LLC, USA. SLES was supplied by Bekee (M) Sdn Bhd, Malaysia.

2.2. Synthesis of pristine LDH

Synthesis of LDH was carried out by heating at 70 °C, adjusted using 1M sodium hydroxides solution 500 mL aqueous solution containing 38.44 g of magnesium nitrate hexahydrate and 18.76 g of aluminium nitrate nonahydrate at 100 rpm for 24 hours. The LDH obtained was filtered, washed thoroughly and dried at 60 °C for 24 hours.

2.3. Modification of LDH by SLES

SLES-modified LDH was prepared by shaking required amount of LDH in 500 mL of SLES solution at 75 °C for 24 hours. The SLES solution was prepared by dissolving the required amount of this compound in 500 mL water at 50 °C and 500 rpm. The modified LDH was then filtered, washed from several times and dried an oven for 24 hours and 60 °C.

2.4. Preparation of PLA/SLES-LDH Nanocomposites

PLA and its nanocomposites sheets were prepared using solvent casting method. Ten grams of PLA was first dissolved in 120 mL chloroform at 40 °C. For the composite preparation, a chosen amount of modified LDH was then added into the prepared PLA solution. Then the mixtures were stirred vigorously for 3 hours, sonicated for another 30 minutes, casted in a petri dish and allowed to dry at room temperature until the film was obtained. The amount of the LDH used was 1, 3, 5, 7 and 10wt%.

2.5. Characterization

(a) X-Ray diffraction method: X-ray diffraction patterns of PLA samples were recorded using a XRD 6000 Diffractometer, Shimadzu. The instrument was operated at 30 kV/30 mA and produced Cu-Kα radiation (λ = 0.15405 nm). All samples were scanned at a rate 0.2° 20/min in a range from 2 to 65°.

(b) Fourier transform infrared spectra: Spectra of samples sheets were recorded using a Perkin Elmer System FTI 1650 Spectrum BX, Spectrometer using an average 10 scan in the range from 400 to 4000 cm⁻¹.

(c) Tensile measurement: Five dumbbell shape samples were prepared using ASTM Standard D638-V. The thickness and width of the gauge section of each specimen were calculated. The test was carried out by Instron Universal Testing Machine to obtain value of tensile strength, tensile modulus and elongation at break.

(d) Biodegradation test: The biodegradation test of the PLA film was carried out by burying the samples in the soil at our campus in the Sungai Besi Camp, Kuala Lumpur Malaysia for six months. Each film sample was cut into 1 cm × 1 cm dimension and buried in the soil at a depth of about 8 cm from the ground surface for a chosen period. It was then washed with distilled water and dried in an oven at 35 °C. The weights before and after the burial of each sample were recorded.

3. Results and discussion

3.1. Characterization of SLES-LDH

In this work, preparation of LDH was successfully modified by SLES via ion exchange process as shown in Fig. 1. The modification of LDH was conducted by exchanging the original anion, NO₃⁻ to long alkyl surfactant anion, CH₃(CH₂)₁₀CH₂(OCH₃CH₂)₂OSO₃⁻ in order to enlarge the interlayer spacings of LDH, hence enhance its hydrophobicity.

Fig. 1: Schematic representation of ion exchange process

Fig. 2 shows the x-ray diffraction results of pristine LDH and SLES-LDH. Sharp peaks in both pristine LDH and SLES-LDH pattern indicate that they are crystal with well-defined interlayer structure (Jamal et al., 2015). The peak of SLES-LDH was at 2θ of 25.8° which smaller than that of the pristine LDH shows the intercalation of surfactant anions into LDH interlayers was successfully carried...
out; therefore its interlayer distance is bigger than that of the original material. The interlayer spacings can be calculated using Bragg’s equation where \( n \lambda = 2d \sin \theta \) where \( n \) is 1, \( \lambda \) = wavelength of Cu-K\( \alpha \) radiation, \( d \) = interlayer spacing of pristine LDH and SLES-LDH and \( \theta \) = diffraction angle. The calculation shows the interlayer spacings for the pristine LDH is 8.10 Å while for the SLES-LDH is 34.23 Å. Increase of the interlayer spacing due to intercalation of a long hydrophobic anionic surfactant chain has been reported earlier (Maiti et al., 2008).

Fig. 2: XRD patterns of pristine and modified LDH

Fig. 3: FTIR spectra of pristine and modified LDH and SLES-LDH

Fig. 4: Effect on SLES-LDH content on tensile strength and tensile modulus of PLA/SLES-LDH nanocomposite

Fig. 5: Effect of SLES-LDH content on elongation at break of PLA/SLES-LDH nanocomposites

3.2. Characterization of PLA/SLES-LDH nanocomposites

Tensile strength and tensile modulus and elongation at break of neat PLA and its nanocomposites are displayed graphically in Fig. 4 and 5 respectively. Addition of 1 wt% SLES-LDH in the PLA improves the tensile strength of PLA/SLES-LDH nanocomposites by about 26% which is from 19.0 MPa to 23.0 MPa, and attained the highest strength with 1 wt% SLES-LDH for PLA (23 MPa). It is reported that dispersion of SLES-LDH in PLA resulted in strong interfacial interaction between polymer matrix and layered silicates, which produces higher mechanical strength when compared to conventional microcomposites (Ren, 2010). However, further increase amount of SLES-LDH slightly does not affect the tensile strength which can be considered as no improvement in tensile strength. The addition of SLES-LDH also shows the increment of tensile modulus of PLA nanocomposites compared to neat PLA.

For Fig. 5, the elongation at break of PLA decreases with the addition of SLES-LDH. The presence of SLES-LDH reduced the mobility of PLA chains, thereby reducing the flexibility of PLA. Consequently, higher SLES-LDH loading produced high tensile strength with low elongation.
SEM was used to observe surface morphology of the fractured tensile strength samples. Fig. 6a and 6b displays the comparison of surface micrograph of neat and PLA/1wt% SLES-LDH nanocomposites.

Fig. 6a shows the smooth and brittle surface of neat PLA polymer. In contrast, rough and stretch surface can be seen in PLA nanocomposite fractured surface (Fig. 6b) indicated that strong characteristic in PLA when added SLES-LDH. The addition of SLES-LDH improves the interaction between additive and polymer, showed that the nanocomposites are compatible with the SLES-LDH, and resulted in the improvement of tensile strength.

Fig. 7 shows the weight loss percentages of the pure PLA and PLA nanocomposites buried in the soil for up to six months. It displays that degradation rate the composites is higher than that of the pure PLA. It also indicates the weight loss of PLA nanocomposite increases as the weight percentage of SLES-LDH in the composites is enhanced. Similar observations was reported by Mahboobeh et al. (2012) when they studied the degradation of PLA/stearate modified LDH composites and Ray and Okamoto (2003) when the worked with the PLA/modified MMT degradation. These results also reveal that the pure PLA degrades slowly; only about 1% of weight loss is recorded after 6 months it is buried in the soil. In contrast, the PLA containing 10wt% SLES-LDH degrades faster whereby 17% of the weight loss is recorded the same period of the exposure. It is reported that PLA degradation is a two-step reaction. Initially, high molecular weight of PLA is hydrolysed to produce lower molecular weight compounds. This step is temperature and humidity dependent. Then, the degradation process is continued with the presence of microorganism in the environment to convert them into carbon dioxide, water and humus (Tsujii and Ikada, 2000; Gruber and O’Brien, 2002).

![Fig. 6: (a) Neat PLA; (b) PLA/1 wt% SLES-LDH nanocomposite](image)

![Fig. 7: Time course for biodegradation of pure PLA and PLA nanocomposites with various weight ratios of LDH](image)

### 4. Conclusion

PLA/SLES-LDH nanocomposites were synthesized successfully via solvent casting method. The used modification of LDH by SLES was prepared by ion exchange method. With the addition of 1wt% SLES-LDH in PLA, the enhancement of tensile strength was obtained around 26% but poor result on its flexibility. For biodegradability study, fast degradation can be obtained in PLA nanocomposites compared to its pure with 17% mass loss after six months buried in soil. However, the degradation rate depends on many factors such as moisture and temperature.

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### References

Anderson KS, Schreck KM, and Hillmyer MA (2008). Toughening polylactide. Polymer Reviews, 48(1): 85–108.

Bordes P, Pollet E, and Averous L (2009). Nanobiocomposites: biodegradable polyester/nano clay system. Progress in Polymer Science, 34(2): 125-155.
Feitknecht W and Gerber M (1942). Double hydroxides and basic double salts. II Mixed precipitates from calcium-aluminium salts solutions. III Magnesium-aluminium double hydroxides. Helvetica Chimica Acta, 25: 106-131.

Gruber P and O’Brien M (2002). Polylactides “NatureWorksTM PLA. In: Doi Y and Steinbüchel A (Eds.), Biopolymers polyesters III applications and commercial products: 235-249. Wiley-VCH, Weinheim, Germany.

Hochstetter C (1842). Untersuchung über die Zusammensetzung einiger Mineralien. Advanced Synthesis and Catalysis, 27(1): 375-378.

Jamal SH, Yunus WMZW, Azilah N, Kasimb M, Khima OK, and Ahmadc M, (2015). Preparation and characterization of sodium lauryl ether sulphate modified layered double hydroxides. Jurnal Teknologi (Science and Engineering), 76(3): 109-113.

Katyar V, Gerds N, Koch CB, Risio J, Hansen HCB, and Plackett D (2010). Poly L-lactide-layered double hydroxide nanocomposites via in situ polymerization of L-lactide. Polymer Degradation and Stability, 95(12): 2563-2573.

Livi S, Bugatti V, Estevez L, Duchat-Rumeau J, and Giannelis EP (2012). Synthesis and physical properties of new layered double hydroxides based on ionic liquids: Application to a polylactide matrix. Journal of Colloid and Interface Science, 368(1): 122-129.

Luo YB, Wang XL, Xu DY, and Wang YZ (2009). Preparation and characterization of poly [lactic acid]-grafted TiO2 nanoparticles with improved dispersions. Applied Surface Science, 255(15): 6795-6801.

Mahboobeh E, Wan Yunus WMZ, Hussein Z, Ahmad M, and Ibrahim NA (2010). Flexibility improvement of poly (Lactic Acid) by stearate modified layered double hydroxide. Journal of Applied Polymer Science, 118(2): 1077-1083.

Mahboobeh E, Wan Yunus WMZ, Ibrahim NA, and Shameli K (2012). Degradability enhancement of poly (Lactic Acid) by stearate-Zn3Al LDH nanolayers. International Journal of Molecular Sciences, 13(7): 7938-7951.

Maiti M, Bhattacharya M, Bhowmick AK, (2008). Elastomer nanocomposites. Rubber Chemistry and Technology, 81(3): 384-469.

Ray SS and Okamoto M (2003). Polymer/layered silicates nanocomposites: A review from preparation to processing. Progress in Polymer Science, 28(11): 1539-1641.

Ren J (2010). Biodegradable poly (Lactic Acid): synthesis modification, processing and applications. Tsinghua University Press, Beijing, China.

Tsuji H and Ikada Y (2000). Biodegradable polyesters for medical and ecological applications. Macromolecular Rapid Communications, 21(3): 117-132.

Wang B, Siahaan TJ, and Soltero RA (2005). Drug delivery: Principles and applications. John Wiley and Sons, Hoboken, USA.

Williams GR and O’Hare D (2006). Towards understanding, control and application of layered double hydroxide chemistry. Journal of Materials Chemistry, 16(30): 3065-3074.

Wypch G (2004). Handbook of plasticizers. ChemTech Publishing, Toronto, Canada.

Zhao N, Shi S, Lu G, and Wei M (2008). Polylactide (PLA)/layered double hydroxides composite fibers by electrospinning method. Journal of Physics and Chemistry of Solids, 69(5): 1564-1568.