

Thermal stability of PMMA–clay hybrids

TANUSHREE CHOUDHURY* and NIRENDRA M MISRA

Department of Applied Chemistry, Indian School of Mines University, Dhanbad 826 004, India

MS received 9 December 2008

Abstract. Materials with small particle size are being extensively used in composites and hybrid materials. Exfoliated clay–polymer hybrids show enhanced properties. Exfoliation of clay platelets can be affected by selecting dispersing agents. In the present work, clay dispersed by natural dispersant (soap stone powder), cetyl trimethyl ammonium bromide (CTAB) dispersed clay and acid clay (amorphous clay) are taken. They are then polymerized with poly methyl methacrylate (PMMA) by solution intercalation method. The thermal stability of these different clay–PMMA hybrids have been studied and compared with that of pure PMMA by differential scanning calorimeter (DSC). The bonding of clay with PMMA has been studied by IR. Morphology of clay–PMMA hybrids has been shown by SEM and XRD which indicate partially exfoliated structure in T606-4 and intercalated structures in T606-6 and T606-2.

Keywords. Dispersants; DSC; exfoliation; hybrid; nanoclay; poly (methyl methacrylate); SEM; thermal stability; XRD.

1. Introduction

Clay minerals are hydrated alumino-silicates having layered structure. Clay particles that have the sizes of a few nanometers thick, if properly dispersed in the polymer matrix impart unique combination of physical and chemical properties that make these composites suitable for making coatings and films for a variety of industrial applications. Clay minerals are composed of one central octahedral layer sandwiched between two tetrahedral layers. Isomorphous substitution of tetravalent Si by trivalent Al in tetrahedral layer and trivalent Al by divalent Mg in octahedral layer leads to charge deficiency which is compensated by compensating cations on the layer surfaces. These cations then get easily exchanged by other cations in solution. Thus clay minerals bear potential of forming organic–inorganic hybrid materials.

There are three types of polymer–clay hybrids as reported in literature (Ruiz-Hitzky 2004). They are (i) intercalated hybrids which result from the intracrystalline insertion of organic compounds inside the layers of certain lamellar solids, (ii) exfoliated or delaminated hybrids which are formed when the layers of clay are delaminated and the resulting platelets are homogeneously dispersed throughout the polymer matrix and (iii) sol–gel hybrids which refer to organically modified silicates or ceramics.

Exfoliated polymer clay hybrids exhibit improved modulus, lower thermal expansion coefficient, gas permeability, higher swelling resistance and enhanced ionic

conductivity compared with pristine polymers due to nanoscale structure of hybrids and synergism between polymer and silicate (Xu *et al* 2003). Delamination (or exfoliation) of clay depends on the dispersion of clay, and properties of the hybrids depend on the interaction of delaminated clay with the polymer or fibre. In order to obtain enhancement in these properties, surface of the clay which is hydrophobic in nature needs to be modified so that it can interact with hydrophobic polymers. Surface of the clay is generally modified by cation exchange process. The ability of clays to exchange cations between each of their individual layer and retain them is a unique characteristic. The intercalated cations can be exchanged by other cations by treatment of other cations in solution. Thus cation exchange capacity can be defined as the maximum amount of cations that a given amount of clay can take up and is constant (Van Olphen 1963). The clay platelets are held together by strong electrostatic forces of attraction between its layers and inter-gallery cations (Ma *et al* 2003). As a result of modification of its surface by cation-exchange process, the layers get separated or delaminated due to reduction in surface energy. Alkyl ammonium cations incorporated in organosilicates lower the surface energy of inorganic host and improve the wetting characteristics with polymer (Sinha Ray and Okamoto 2006). It provides functional group that can react with polymer or initiate polymerization of monomers to improve interfacial strength between inorganic host and polymer (Sinha Ray and Okamoto 2006).

The polymer clay hybrids are obtained by the following three main methods: (i) *in situ* polymerization, (ii) exfoliation-adsorption and (iii) melt intercalation (Ale-

*Author for correspondence (tanu_c79@yahoo.co.in)

xandre and Dubois 2000). Among thermoplastic–clay hybrids, PMMA has been widely used as a model matrix because it can be easily polymerized in bulk (*in situ*) solution, suspension and emulsion techniques (Xu *et al* 2006). Numerous works have been reported in literature on PMMA as PMMA–clay hybrids have a potential advantage in reducing flammability, gas permeability and improving thermal and mechanical properties without any loss in optical clarity (Zheng *et al* 2005).

In the present work, we examine hybrids of poly (methyl methacrylate) (PMMA) with three different types of clay (clay dispersed by natural dispersant, CTAB treated clay and acid or amorphous clay). Hybrid formation is evaluated by scanning electron microscope (SEM). Bonding between clay and PMMA is being shown by FTIR. The thermal stabilities of the resulting hybrids have been compared by DSC. The present study was stimulated by the observation of earlier works where it is shown that surfactant treated clay has a major role on the thermal stability of PMMA as compared to pristine clay.

2. Experimental

2.1 Materials

Clay used in this experiment was mainly Bikaner bentonite. Cetyl trimethyl ammonium bromide (CTAB) used for dispersing clay was acquired from Ranbaxy Fine Chemicals Ltd. Polymer PMMA used for intercalation was of molecular weight 15,000 and bought from Acros Organics. Natural dispersant or soap stone powder was acquired from natural source.

2.2 Modification of clay

Clay was made organophilic by CTAB and a natural surfactant. To 1 g clay, 0.1% dispersant solution (0.01 g of dispersant in 10 ml water) was mixed under constant stirring for 4 h so that clay gets well dispersed in it. One drop of clove oil was added to the naturally dispersed clay suspension to keep it bacteria free. The resulting suspension was centrifuged and dried in room temperature.

2.3 Preparation of acid/amorphous clay

Known amount of clay was taken and kept in furnace till the temperature reached 600°C and it was immediately taken out and quenched in 0.1 N HCl solution. The resulting solution was centrifuged and dried at room temperature.

2.4 Synthesis of PMMA/clay hybrids

PMMA–clay hybrid was prepared by solution-intercalation method (Kim *et al* 2005). 10 wt% clay was used to

prepare the hybrid. 1 g PMMA was dissolved in acetone and stirred vigorously in a round bottomed flask fitted with stopper at 40°C for 30 min. The organophilic clay (10 wt%) was added to the polymer solution and continued to stir for 1 h. The solution product was then filtered off and washed thoroughly. The product was then dried at room temperature.

2.5 Instrumentation

DSC of PMMA–clay hybrid was performed on differential scanning calorimeter Perkin Elmer DSC-7 under nitrogen atmosphere at a scan rate of 20°C/min. IR was done on Perkin Elmer FTIR Spectrometer Spectrum 2000. SEM of PMMA–hybrids was taken using Hitachi S-3400N, Japan, using 3 kV. XRD of PMMA–clay hybrids was done using X'Pert Software, Phillips, powder diffractometer equipped with Cu-K α generator ($\lambda = 1.5405600$). Generator tension was 35 kV.

3. Results and discussion

3.1 DSC analysis

DSC showed a 9.12°C (as indicated in table 1a) increase in the glass transition temperature (T_g) of the hybrid prepared with clay (dispersed by natural dispersant-T606-4) as compared to pure PMMA (T703-10). This result was higher than hybrid prepared with acid clay (T606-2) which showed an increase in T_g of 7.47°C while CTAB dispersed clay–PMMA hybrid (T606-6) showed an increase of 6.45°C. Thus clay dispersed by natural dispersant (T606-4) provides greater thermal stability to PMMA as compared to the other two clays.

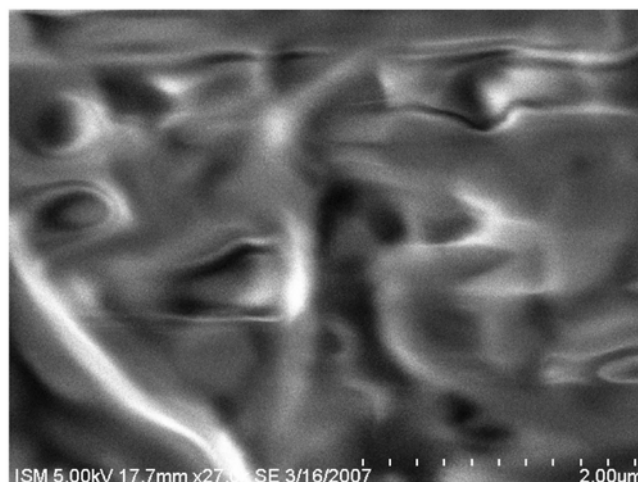


Figure 1. SEM of PMMA–clay hybrid dispersed by natural dispersant showing partially exfoliated structure.

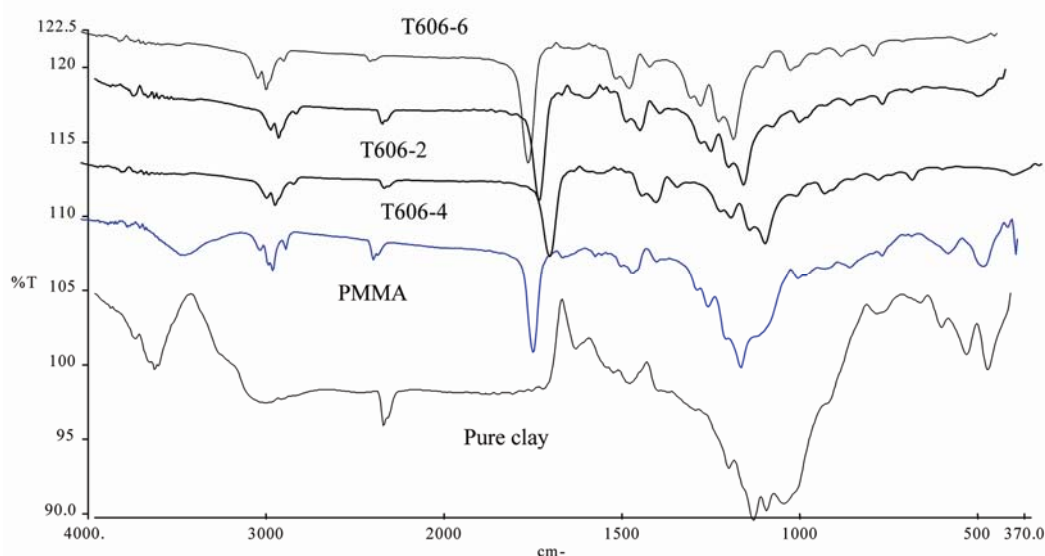


Figure 2. FTIR spectra of PMMA–clay hybrids.

Table 1a. Table showing T_g of PMMA–clay hybrids.

Sl. no.	Type of hybrid	Structure	T_g (°C) (DSC)
1.	Pure PMMA	–	97.12
2.	PMMA + clay (dispersed by natural dispersant)	Partially exfoliated	106.24
3.	PMMA + acid clay	Intercalated	104.59
4.	PMMA + CTAB clay	Intercalated	103.57

Table 1b. Table showing d -spacing of various organically modified PMMA–clay hybrids.

Sl. no.	Sample name	d -spacing (nm)	Angle (2θ)
1.	T606-4	3.56	2.47
2.	T606-2	2.85	3.09
3.	T606-6	2.74	3.21
4.	Pure MMT	1.6	5.44

3.2 SEM analysis

Figure 1 shows SEM micrograph of PMMA–clay hybrid where clay has been dispersed by natural dispersant (T606-4). It is a partially exfoliated structure showing individual clay layers (not in registry) with PMMA embedded in the galleries of clay. PMMA–clay hybrids of acid clay (T606-2) and CTAB treated clay (T606-6) show intercalated structures as can be seen from d -spacing data supported by XRD. Clay dispersed by natural dispersant might have helped in increasing the interlayer spacing of clay thus fastening intercalation of PMMA within the galleries of clay leading to partially exfoliated structure.

3.3 FTIR analysis

IR spectra of all PMMA–clay hybrids give a distinct absorption peak at 1730 cm^{-1} (C=O stretching), 2996 cm^{-1} and 2952 cm^{-1} ($-\text{CH}_2$ stretching), wide absorption peak at $3400\text{--}3600\text{ cm}^{-1}$ (H-bond). Characteristic montmorillonite peak was observed between 900 and 1000 cm^{-1} (Si–O–Si stretch) which was very much reduced in the spectrum recorded for the hybrids as can be seen in figure 2. This may be due to the fact that PMMA being grafted on the clay surface.

3.4 XRD analysis

The d -spacing observed for virgin MMT is 1.6 nm ($2\theta = 5.44^\circ$) which always increases upon organic modification (Jash and Wilkie 2005; Stadtmueller *et al* 2005). The d -spacings for the various organically modified clays are shown in table 1b. It is thus clear that d -spacing is greatest in T606-4 (PMMA–clay hybrid where clay has been dispersed by natural dispersant), thus indicating that exfoliated structure may have been attained.

4. Conclusions

Synthesis of PMMA/clay hybrids by solution intercalation method using three different types of clays (clay dispersed by natural dispersant, CTAB treated clay, acid/amorphous clay) resulting in partially exfoliated and intercalated structures, respectively was described. The different morphologies of the hybrids resulted in significant thermal property improvements in comparison to pristine polymer. Partially exfoliated hybrid of PMMA

with clay (dispersed by natural dispersant-T606-4) exhibited an increase in T_g 9.12°C higher than PMMA while for intercalated structures of PMMA with CTAB clay (T606-6) and acid clay (T606-2) showed only 6.45°C and 7.47°C increase in T_g . Using PMMA/clay hybrids, ion selective membranes can be developed for various metal ion detections. Other applications lie in film packaging and gas storage (Meneghetti and Qutubuddin 2006).

References

- Alexandre M and Dubois P 2000 *Mater. Sci. & Eng.* **28** 1
Jash P and Wilkie C A 2005 *Polym. Degrad. & Stab.* **88** 401
Kim H B, Choi J S, Lee C H, Lim S T, Jhon M S and Choi H J 2005 *Eur. Polym. J.* **41** 679
Ma J, Xiu J, Ren H J, Yu Z and Mai Y 2003 *Polymer* **44** 4619
Meneghetti P and Qutubuddin S 2006 *Thermochim. Acta* **442** 74
Ruiz-Hitzky E 2004 *Organic-inorganic materials: From intercalation chemistry to devices*, in *Functional hybrid materials* (eds) Pedro-Gomez Romero and Clement Sanchez (Weinheim: Wiley VCH) pp 15–49
Sinha Ray S and Okamoto M 2003 *Prog. Polym. Sci.* **28** 1539
Stadtmueller L M, Ratnac K R and Ringer S P 2005 *Polymer* **46** 9574
Van Olphen H (ed.) 1963 *Clay colloid chemistry* (London: Wiley Interscience)
Xu W, Liang G, Zhai H, Tang S, Hang G and Pan W 2003 *Eur. Polym. J.* **39** 1467
Xu Y, Brittain W J, Xue C and Eby R K 2006 *Polymer* **45** 3735
Zheng X, Jiang D D and Wilkie C A 2005 *Thermochim. Acta* **435** 202