

## Melt rheological behavior and batch foaming of modified PMMA

Yuki Kawahara<sup>1</sup>, S.K.Sukumaran<sup>1</sup> and Masataka Sugimoto<sup>1\*</sup>

Yuichi Shindo<sup>2</sup>, Kohhei Nishino<sup>2</sup>

<sup>1</sup> Yamagata University, 4-3-16 Jonan, Yonezawa, Yamagata, 992-8510, Japan

<sup>2</sup> Denka K.K., 6 Goi-minamikaigan, Ichihara, Chiba, 290-8588, Japan

(\*sugimoto@yz.yamagata-u.ac.jp)

### Abstract

The heat resistance of poly(methyl methacrylate)(PMMA) can be enhanced by blending styrene-methyl methacrylate-maleic anhydride copolymer (SMA) which is compatible with PMMA, without substantially changing the mechanical properties of original PMMA. Here from the viewpoint of the processability of the blends, we studied the rheological properties and foaming behavior by using CO<sub>2</sub> as a foaming agent.

We used commercial PMMA ( $M_w = 100,000$  and  $T_g = 108$  °C) and SMA ( $M_w = 160,000$  and  $T_g = 132$  °C). SMA/PMMA (50/50 wt%) blend (B50) was extruded by a single screw extruder ( $\phi=40$ mm) at 240 °C and screw rotation speed of 100 rpm. The blend was transparent and considered to be compatible ( $M_w$  140,000,  $T_g$  120 °C).

CO<sub>2</sub> was absorbed to the sheet samples in the pressure vessel of 25 °C for 8 hr. Adsorption pressure was adjusted so that CO<sub>2</sub> concentration of each specimen was to be 14 wt%. After the saturation, samples were soaked in the oil bath for 1 min to induce foaming at  $T_g + 10$  °C for each specimen (SMA: 142 °C, B50: 130 °C, PMMA: 118 °C). Then, samples were cooled in the water bath of 15 °C for 1 min.

Fig.1 shows SEM images of PMMA, B50, and SMA foamed at  $T_g+10$  °C and 15MPa of CO<sub>2</sub>. SMA indicated much smaller cell size than that of PMMA, the blends B50 also showed fine cells. Fig.2 summarizes the change in the cell diameter  $d$  and the expansion ratio  $E_r$  as a function of the blend ratio of SMA. Although  $d$  simply decreased as increasing the SMA content,  $E_r$  showed an increase with the SMA content and reached a peak at 50 wt% (B50). Above the peak,  $E_r$  was decreased.

We also studied rheological behaviors, such as extensional viscosity and dynamic viscoelasticity, to discuss the foamabilities of SMA and SMA/PMMA blend. This will be presented at the conference.

**Keywords** :“ Carbon dioxide”, “foaming”, “modified acrylic copolymer”

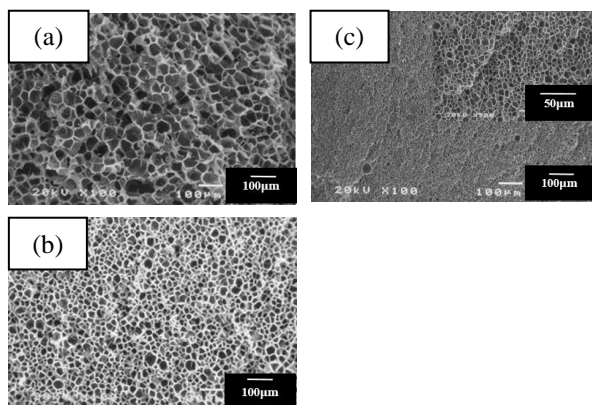


Fig.1 SEM images of foams with CO<sub>2</sub> at  $T_g+10$  °C:  
(a) PMMA, (b) B50, and (c) SMA

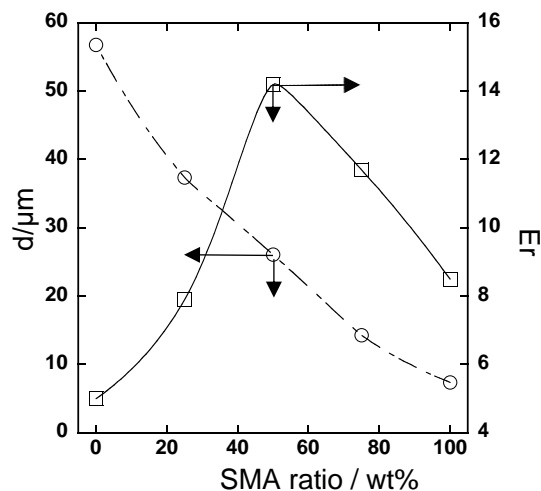


Fig.2 Effect of PMMA/SMA composition on average cell diameter  $d$  and expansion ratio  $E_r$ .