

Abstract Submitted  
for the MAR15 Meeting of  
The American Physical Society

**Bottlebrush additives drive formation of vesicle chains in polymer blends** HUI ZHEN MAH, PANTEA AFZALI, Univ of Houston, RAFEAL VERDUZCO, Rice University, GILA STEIN, Univ of Houston — The effects of bottlebrush polymer additive with poly (styrene-*r*-methyl methacrylate) side-chains on the thin film morphology of polystyrene (PS) and poly (methyl methacrylate) (PMMA) blends were studied. Results were compared to PS/PMMA blends with diblock copolymer PS-*b*-PMMA compatibilizer and without any additive. Thin films were spin casted from toluene onto a “neutral” silicon surface and then annealed at a fixed temperature of 150°C for a range of times (up to 85 mins). The morphology of the films was characterized using optical microscopy and atomic force microscopy. In the absence of any additive, the PS/PMMA blend rapidly de-mixes to form macroscale domains, while high loadings of the PS-*b*-PMMA additive can compatibilize the blend and suppress phase separation. However, the bottlebrush polymer additive drives the formation of well-organized vesicle chains in the PS/PMMA blend films. This morphology is favored by entropic considerations as the bottlebrush polymers are more stable than linear chains at the PS/PMMA interface and the brush like surface attracts.

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Date submitted: 13 Nov 2014

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