

Tuning Polymer-Grafted Particle Monolayer Structure at the Air-Water Interface by Introducing Anisotropic Features

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Abstract

Fabrication of anisotropic material is one of the important topics and we examined to introduce “anisotropic” nature by spreading polymer-grafted particle on the medium with polymer-reactive potential. Poly (*tert*-butyl methacrylate) (P*t*BMA) was polymerized from polystyrene latex (PSL) surface by ATRP to give P*t*BMA-grafted PSL (PSL-P*t*BMA). Particle monolayer was formed at air-water and air-acidic water interfaces and the monolayer characteristics were compared by π -A isotherm measurements, SEM observations, and contact angle measurements. π -A isotherms, in particular, indicates that the interaction between polymer chains become stronger by changing the subphase condition, which suggests that anisotropic particle monolayer formation.

Keywords

Anisotropic Particle, Polymer-Grafted Particle, Air-Water Interface, Polymer Brush, Particle Monolayer

1. Introduction

A particle at interface is a simplest system concerning interfaces and has wide variety of interests ranging from basic science to industrial process [1]. Among the particulate fabrications, in particular, there has been a great interest in creating materials with anisotropic property. One is to fabricate materials with anisotropic shape such as disc or rod. Another is to make anisotropic nature with spherical shape, *i.e.*, Janus particles [2] [3]. In the latter category, preparation process is further categorized into two. One is to making use of association behavior like micelle or emulsion including non-equilibrium status [4] [5] [6]

[7] [8]. The other is to utilizing the planner interface, for example, air/water interface or liquid/liquid interface [9] [10] [11] [12], which concerns the topic of this paper.

In this paper, we describe the fabrication of polymer-grafted particle monolayer with anisotropic features introduced at the air-aqueous medium interface. We have been studying polymer-grafted particle system at the air-water and oil-water interfaces [13]-[18]. We systematically investigated the effects of molecular weight, graft density, core particle, polymer species, and interface on the polymethacrylate-grafted particle monolayer structure by π -A isotherm measurement, SEM observation and X-ray reflectivity measurement. So far, we found that polymethacrylate-grafted particles forms monolayer at air-water interface and grafted polymethacrylate chains are fairly extended at the interfaces. On the basis of these previous findings, we verified to fabricate anisotropic polymer-grafted particle by utilizing potential reactivity at the air/water interface.

2. Experimental Method

2.1. Synthesis and Characterization of *PtBMA-Grafted Polystyrene Latex*

Polystyrene latex with chloromethyl groups was synthesized by soap-free emulsion polymerization as previously described [16]. The averaged diameter of polystyrene latex was estimated by dynamic light scattering with DLS-8000 (Otsuka Electronics Co., Ltd., Hirakata, Japan). Polydispersity was confirmed to be within 1%. Graft polymerization of PMMA from polystyrene latex surface via copper-mediated ATRP was carried out as follows [15] [19]: The polystyrene latex dispersion of DME was mixed with freshly distilled *PtBMA*, CuBr (I) and *N,N,N',N'',N'''*-pentamethyldiethylenetriamine (PMDETA). The mixture was carefully bubbled and degassed by ultrasonification under argon atmosphere. Then free initiator, α -chloro-*p*-xylene, was added to the mixture. Typical feed is 40 mg of polystyrene latex, 6.6 ml of dry MMA, 7 mg of CuBr, and 32 mg of PMDETA with 6.4 mg of free initiator. The solution was degassed by four or five freeze-pump-thaw cycles and the ampule was sealed off completely with argon purge. The polymerization was carried out at 80°C for 4 days to obtain *PtBMA-grafted* polystyrene latex (PSL-*PtBMA*).

Molecular weight and polydispersity of free polymer were confirmed by GPC equipped with column oven (CTO-10A/10AC, Shimadzu, Kyoto, Japan) and refractive index detector (RI153, JASCO Corp., Tokyo, Japan). Obtained polymer-grafted particle was firstly characterized by dynamic light scattering (DLS). Time correlation function was fitted to single exponential function, and hydrodynamic diameter and high monodispersity was confirmed.

Estimation of graft density is performed by utilizing IR spectrum measurement as follows: Calibration curves for *P(tBMA)* and PSL was obtained respectively with 2-cyanoacetoamide as internal standard as same as previous study

[16]. The following peaks were picked up for calibration curves; 1475 cm^{-1} (PSL, benzene ring in-plane vibration), 1140 cm^{-1} (P β BMA, C-O stretching), and 2270 cm^{-1} (cyanoacetoamide, $\text{C}\equiv\text{N}$ stretching). The linear relationships were obtained between molar ratio of [P β BMA]/[2-cyanoacetoamide] (or [styrene]/[2-cyanoacetoamide]) and peak area ratio [C-O stretching]/[$\text{C}\equiv\text{N}$ stretching] (or [benzene ring in-plane vibration]/[$\text{C}\equiv\text{N}$ stretching]). By IR measurement of PSL-P β BMA as in **Figure 1**, the amount ratios of P β BMA to PSL were obtained for each sample. Then, the graft density of P β BMA was calculated with the two assumptions that 1) the molecular weight of grafted P β BMA is equal to M_n obtained by GPC, and 2) polystyrene latex was made of only polystyrene with density of 1.047.

2.2. Preparation of PMMA-Grafted Polystyrene Latex Monolayer and Measurement of π -A Isotherms

The preparation of particle monolayer was carried out as previously reported [13]-[18]. The chloroform solution of $2.5 \times 10^{-4}\text{ g/ml}$ PSL-P β BMA was spread by microsyringe on ultra-pure water surface and HCl solution (pH3), respectively in Langmuir-Blodgett (LB) trough ($150\text{ mm} \times 160\text{ mm}$). After evaporation of solvent, the water surface was compressed and expanded by Teflon[®] barrier at 0.025 mm/sec to measure π -A isotherm at 25°C by USI-3-00 (USI System, Fujioka, Japan). At various surface pressures on a second compression, particle layer was deposited on glass plate by LB method at the speed of 10 mm/min . We observed the deposited particle layers by SEM (SM-7000FSK, JEOL Ltd., Tokyo, Japan). In order to estimate the basic property of a deposited layer on a substrate, contact angle measurement was measured by dropping $0.5\text{ }\mu\text{L}$ pure water on a substrate and observing the drop shape by a digital camera with a close-up lens [16].

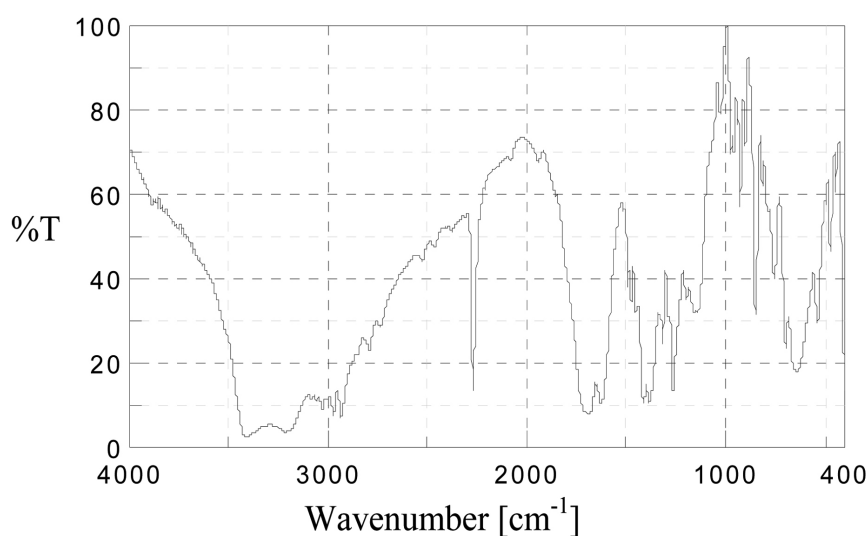


Figure 1. IR spectrum for PSL-P β BMA added with cyanoacetoamide with internal standard.

3. Results and Discussion

The characteristics of core PS particle and PSL-P β BMA are summarized in **Table 1** and **Table 2**, respectively. Important parameters for PSL-P β BMA, molecular weight of grafted polymers, graft density and hydrodynamic size in the solution, are summarized in **Table 2**. Although we obtained P β BMA homopolymer with high monodisperse nature ($M_w/M_n = 1.15 - 1.24$) in the comparable system without PSL, P β BMA molecular weight with PSL showed wider molecular weight distribution. Since the hydrodynamic size of PSL used here is 118 nm, P β BMA brush layer with ca.10-nm thickness covers around PSL latex core. The graft density of 0.14 is relative high graft density [13]-[18].

PSL-P β BMA of well-characterized sample is applied to monolayer formation at the air/water interface. **Figure 2** shows the π -A isotherm of PSL-P β BMA at the air-water interface as curve 1. The π -A isotherm is reproducible and shows the increase of surface pressure at a very large occupied area. For example, expected occupied area with PSL particles with closest packing is indicated by straight line in **Figure 1**, which shows more than 1/10 smaller area. As also discussed in the

Table 1. Characteristics of PSL.

Sample code	D_h /nm	Polydispersity
PSL	118	0.050

Table 2. Characteristics of PSL-P β BMA.

Sample code	M_n	M_w/M_n	Graft density/nm ⁻²	D_h /nm
PSL-P β BMA	76,000	1.41	0.113	144

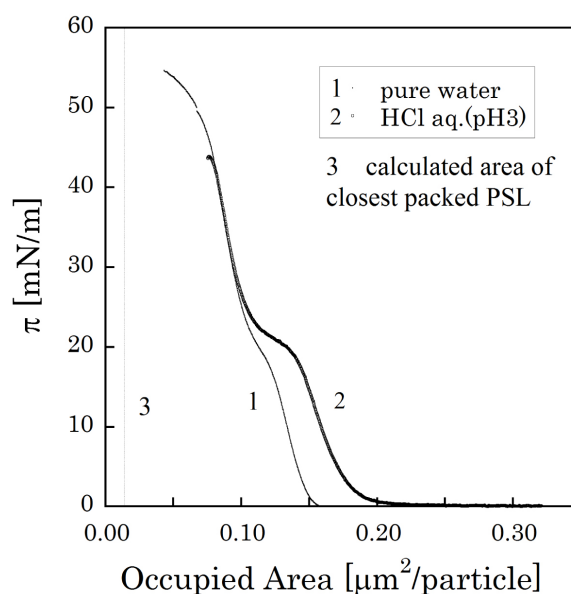


Figure 2. π -A isotherms for PSL-P β BMA on a pure water and on a HCl aqueous solution. Straight line indicates the occupied area of PSL core at closest packing.

former our papers, this indicates that polymer chains grafted from particles are much extended on the water surface [15] [16]. In this data, the occupied area at 1 mN/m corresponds to the interparticle distance of 388 nm assuming closest packing of particle at the air-water interface. On the other hand, calculated interparticle distance with stretched conformation of $M_n = 76,000$ is to be 386 nm. These two values are in agreement. We do know that the molecular weight used here is relative value and contains errors, however, the agreement of these values supports the extended fairly conformation of P β BMA on a water surface similar to PMMA system [16].

SEM images of deposited monolayer by LB method at several constant surface pressures are presented in **Figures 3(a)-(c)**. These SEM images support the monolayer formation, not multilayer. We can see individual PSL core with separation of more than 150 nm.

In order to introduce “anisotropy” in the particle monolayer by utilizing the nature of air-water interface, monolayer formation of PSL- β BMA is conducted on acidic aqueous solution of pH3. Partial proceeding of P β BMA hydrolysis in water subphase may occur because the hydrolysis of P β BMA into PMAA is irreversible reaction while P β BMA remains at the air side. However, the acidic condition here is milder than usual hydrolysis condition for P β BMA [20] [21]. Thus, small fraction of β BMA monomer units would be hydrolyzed to give methacrylic acid units.

In **Figure 2**, π -A isotherm obtained on acidic subphase is also presented as curve 2. The π -A isotherm on acidic subphase shows clear shift to larger occupied area before plateau region. On the other hand, two π -A isotherms at different conditions match after plateau region. Based our previous data, below the plateau region the interaction between polymer chains grafted from particle is observed while the interaction between hard spheres is observed above the plateau region [16]. Considering the interpretation of π -A isotherm, it is reasonable

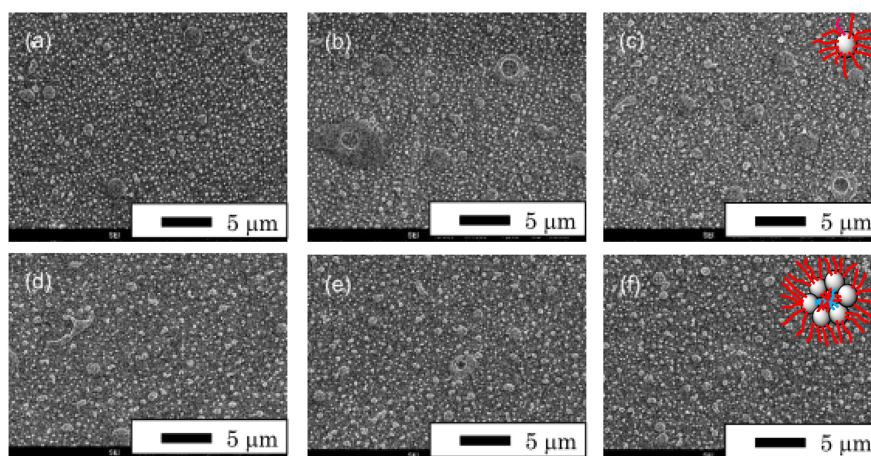


Figure 3. SEM images of deposited PSL-P β BMA monolayer on glass substrates at constant surface pressures ((a), (d): 10 mN/m, (b), (e): 15 mN/m, (c) (f): 30 mN/m) from pure water (a)-(c) and HCl aq. solution (d)-(f). Schematic figures are also shown.

that the difference between the π -A isotherms are observed only below the plateau region. In addition to that, assuming the hydrolysis proceeds in some part, the interaction between PMAA (the resultant of hydrolysis of P β BMA) is enhanced little bit by the electrostatic interaction (although almost MAA is not dissociated in acidic pH range), which agrees with the tendency that surface pressure is increased at the larger occupied area.

SEM images of deposited monolayer by LB method at several constant surface pressures are presented in **Figures 3(d)-(f)**. SEM images obtained at acidic condition shows more aggregated states than the ones at the air-water interface. The anisotropic nature of PSL-P β BMA at acidic condition would favors formation of large assembly of polymer-grafted PSL although we cannot ascertain whether the large aggregates are formed at the water surface or at the glass substrates.

We also estimate the surface property of deposited particle monolayer on substrate. **Figure 4** shows variation of contact angle of water droplet on the particle monolayers with varying surface pressure. The values of contact angles are similar at the two different subphase conditions above 5 mN/m. This is rather reasonable that the outer surface would be P β BMA because particle monolayer is deposited as Y type. In the studied range, aggregation shown in **Figure 3** is not so effective to the contact angles. By changing the substrate to more hydrophobic one, we could tune the contact angle by PSL-P β BMA from acidic subphase.

4. Conclusion

In the view of fabricating anisotropic particles monolayer, we investigated the behavior of PSL-P(*t*-butyl methacrylate) at the air/acidic water interface. P β BMA was polymerized from monodisperse polystyrene latex (PSL) surface by ATRP to give monodisperse P β BMA-grafted PSL (PSL-P β BMA). PSL-P β BMA monolayer was characterized by π -A isotherms and SEM. PSL-P β BMA monolayer deposited from acidic condition shows more aggregated state on glass substrate rather than the ones deposited from pure water, however, the contact angle of the both

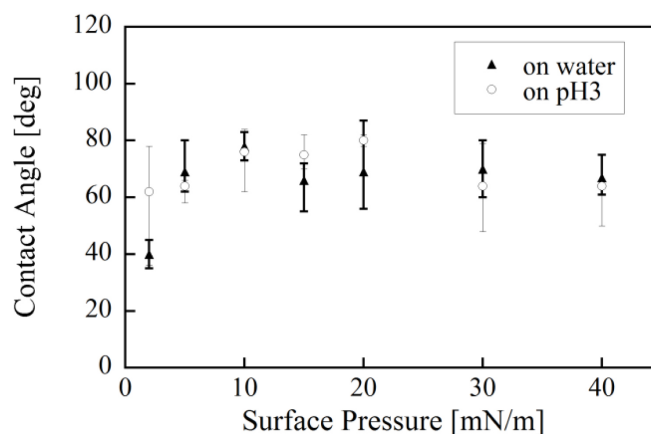


Figure 4. Contact angle of water droplets measured on PSL-P β BMA monolayer deposited on glass substrates at constant surface pressures with different subphase conditions.

deposited film shows similar values.

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