Thin Films of Block Copolymers as Planar Optical Waveguides**

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Block copolymers (BCPs) have attracted significant attention as nanostructured materials, since they self-assemble to form well-defined, ordered, periodic nanoscale morphologies depending on the relative volume fraction of the constituent blocks.[1-2] The versatility of this unique class of polymers offers tremendous potential for their use as templates and scaffolds for applications in microelectronics, separation devices, optics, and optoelectronics.[3-8] While polymer-based optical-waveguide materials have been widely discussed,[9] little attention has been paid to BCP systems[10] in regard to their applications in the field of optical elements. Since the domain size and chemical nature of BCPs can be tailored by controlling the molecular weight and by incorporating specific functionality, BCPs may be ideal candidates for waveguide materials with tunable periodic dielectric constants and optimized physicochemical properties in integrated optical devices. Here, we demonstrate that thin films of BCPs with controlled microdomain orientation can be employed as planar optical waveguides and we investigate their waveguiding properties by optical waveguide spectroscopy (OWS).[11-13]

Resonance coupling between surface plasmons (or plasmon surface polaritons) and incident photons can occur at a metal/dielectric interface, especially in the experimental setup

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known as the Kretschmann configuration (Fig. 1) \[^{11,12,14,15}\] If the thickness of the dielectric layer is further increased (in organic polymeric layers typically thicker than \(\sim 200\) nm), waveguide optical modes, in addition to surface plasmon resonance, can be observed\[^{11,12,15}\]. In this communication, thin films of \(\text{Ag}\) and \(\text{SiO}_2\) layers were prepared by a sol–gel process according to a method reported previously\[^{17,18}\]. The hydrolyzed solution of the sol–gel precursors was delivered onto the 3-MPS-modified Ag surface, and the sample was spun at \(\sim 3400\) rpm for 1 min, producing films with a thickness of \(\sim 33.1\) nm. The Ag/3-MPS/SiO\(_2\) surface was stored in a desiccator at room temperature for a minimum of 2 days to complete the condensation process, and to remove any residual solvent.

An energetically neutral surface for PS-\(b\)-PMMA films was prepared on the SiO\(_2\) layer using a hydroxy terminated poly(styrene-co-methyl methacrylate) random copolymer (PS-\(r\)-PMMA) via the covalent coupling between the hydroxyl groups and the SiO\(_2\) surface.\[^{19,20}\] PS-\(r\)-PMMA was anchored to the SiO\(_2\) surface by spin-coating a 2 wt.-\% toluene solution at 1000 rpm, followed by annealing under vacuum at 165 °C for 3 days. After rinsing with toluene a well-defined, thin layer of PS-\(r\)-PMMA with a thickness of \(\sim 4.2\) nm remained.

Thin films composed of PS-\(b\)-PMMA and PMMA homopolymer mixture, containing 25 wt.-\% PMMA homopolymers with respect to the amount of PMMA block, were prepared by spin-coating 5 wt.-\% toluene solutions at 1000 rpm onto the PS-\(r\)-PMMA surface. The films were annealed at 155 °C for 2 days under vacuum, and then quenched to room temperature. Figure 2A illustrates the phase contrast atomic force microscopy (AFM) image of the surface of a PS-\(b\)-PMMA/PMMA film deposited onto a glass substrate as schematically shown in Figure 1B. The darker and brighter areas are PS matrices and PMMA microdomains, respectively. Hexagonally packed arrays of cylindrical PMMA microdomains oriented normal to the substrate are seen over a wide area of the film. The average center-to-center distance (\(L_{cc}\)) between PMMA microdomains was \(\sim 46\) nm. Deep UV exposure of the films, followed by acetic acid rinsing, was used to selectively remove PMMA microdomains and produce an ordered, nanoporous template. Figure 2B shows a height contrast AFM image of...
the arrays of nanopores (darker areas) in a crosslinked PS matrix after removal of the PMMA microdomains.

The waveguide patterns of the initial BCP film (blue full curves) are displayed in Figures 3A,B for s- and p-polarization, respectively, where characteristic transverse electric (TE) and transverse magnetic (TM) modes are observed. The best fits (blue dotted curves) to the experimental curves were obtained from Fresnel calculations to give the dielectric constants (\(\varepsilon\)) and film thicknesses (\(h\)), respectively.

Best fits to the waveguide patterns of the initial BCP film were not included. Shifts of the resonance coupling modes due to pore generation are indicated by arrows.

Broadening of the coupling dip and the decrease in amplitude of the experimental data compared with the calculated curves is attributed to the distribution in pore diameter and to the surface roughness of the film, which may disrupt the spatial coherence in the waveguide coupling conditions across the measured areas (~1 mm²), corresponding to the laser-beam size. The guided coupling modes of the BCP layer are entirely dictated by its effective dielectric constant (\(\varepsilon_{\text{eff}}\)) and its film thickness. The symmetry of the microdomain morphology: an array of cylinders with the axis aligned normal to the substrate surface dictates that one of the principal axes of the dielectric function, \(\varepsilon_{\text{eff}}\) \(= [\varepsilon_x, \varepsilon_y, \varepsilon_z]\), is parallel to the cylinder axis and that the two other axes are parallel to the film surface. As shown in Figure 2A, the distribution of the cylindrical domains is isotropic, so we assume \(\varepsilon_x = \varepsilon_z\). Therefore, three parameters, \(\varepsilon_x\) (\(= \varepsilon_y\)), \(\varepsilon_z\), and \(h\) need to be determined from the s- and p-polarized waveguide mode spectra.

The dielectric constants and the film thicknesses, obtained from fitting the experimental data in Figure 3 and using the Fresnel calculation, are as follows: \(\varepsilon_x = \varepsilon_y = 2.402\), \(\varepsilon_z = 2.429\), and \(h = 349\) nm.\(^{[11]}\) The thicknesses of the underlying layers (Cr, Ag, 3-MPS, SiO₂, and PS-r-PMMA) were determined separately by surface plasmon resonance (SPR) spectroscopy with the refractive indexes of the 3-MPS, SiO₂, and PS-r-PMMA layers being 1.444, 1.460, and 1.549, respectively.

The pore generation by UV and acetic acid etching was detected via shifts of the mode coupling angles to lower values (indicated by arrows) due to a decrease in \(\varepsilon_{\text{eff}}\) (note that \(\varepsilon_{\text{air}} = 1.0\) is much lower than \(\varepsilon_{\text{PMMA}} = 2.22\)), as shown in Figure 3 (green full curves). The degree of shift (\(\Delta \phi\)) for the TM₁ mode was measured to be 1.9° (from \(\phi = 38.9°\) to \(\phi = 37.0°\)) and that for the TE₁ mode was 1.1° (from \(\phi = 54.4°\) to \(\phi = 53.3°\)). The TM₂ mode was, however, convoluted with the critical angle (\(\phi_c\)) due to total internal reflection, and is not used for the analysis. Fresnel fitting (curves not shown) gives \(\varepsilon_x = \varepsilon_y = 2.377, \varepsilon_z = 2.385,\) and \(h = 341\) nm, for the porous template. From the decrease in the refractive index compared with the initial film, it is evident that sections of the PMMA chains were removed. For comparison, the optical constants (\(\Psi\) and \(\Delta\)) of the films were measured by variable angle spectroscopic ellipsometry (VASE; J. A. Woollam Co., Inc.) at an incidence angle of 70° and best fit to the experimental data using Wvase32 software. The effective dielectric constants of the initial BCP film and the porous template were calculated to be ~2.40 and 1.85, respectively.

\(\varepsilon\) values can also be calculated from the effective medium theory, suitable for the present case, where the domain size is much smaller than the wavelength of guided light, using the following equations\(^{[21,22]}\)

\[
\varepsilon_x = \varepsilon_y = \varepsilon_S + \frac{\varepsilon_D(\varepsilon_D - \varepsilon_S)}{\varepsilon_S - 1/2 \varepsilon_D(\varepsilon_D - \varepsilon_S)/\beta}
\]

\[
\varepsilon_z = f_D \varepsilon_D + f_S \varepsilon_S
\]

where \(f\) is the volume fraction,

\[
\beta = 2 \frac{\varepsilon_S}{\varepsilon_D + \varepsilon_S}
\]

and \(D\) and \(S\) represent cylindrical domain initially occupied by PMMA and PS matrix, respectively. \(\varepsilon_{\text{PMMA}}\) and \(\varepsilon_S\) values used in this study were 2.22 and 2.53, respectively, for the initial BCP film. The \(\varepsilon_S\) value of the crosslinked PS after UV exposure was ~2.58 measured by VASE and used for the prediction of the PS template.

The predicted patterns (green dotted curves) thus obtained, with the assumption that the PMMA microdomains were removed completely, are also shown in Figure 3. It was found that the coupling angles appear at lower angles than those of the experimentally observed patterns (green full curves), indicating that sections of the PMMA chains still remain after the etching condition used in this study. Comparison between the observed and predicted patterns reveals that pores of ~14 vol.-% were generated in the resulting film,\(^{[23]}\) which is smaller than that of the initial PMMA domains, 34 vol.-%. This large discrepancy may be attributed to several reasons:
i) UV exposure was insufficient to induce complete degradation of the PMMA block chains,[24] ii) the effective medium theory may describe composites possessing a lower contrast of the two dielectric constants, e.g., PMMA/PS: 2.22:2.53, better, but may not describe composites with higher contrast in dielectric constants, e.g., air/PS: 1.2.35 well,[25,26] iii) the number of guided modes in the s- and p- mode patterns was not enough to allow for precise evaluation; or iv) the beam spot did not probe the same area during the ex-situ experiment.

The porous templates were immersed in liquid media, a phosphate buffered saline (PBS) solution of pH 7.4, and ethanol (EtOH), in which most of the biological issues are explored. Filling of the pores with PBS followed by exchange with ethanol was detected as shifts of the resonance coupling modes to higher angles, illustrated in Figure 4.[26] The Δθ for TE1 and TM1 modes was 0.7° (from θ = 56.3° to θ = 57.0°) and 1.8° (from θ = 48.8° to θ = 50.6°), respectively. A large portion of this shift was induced by a change in the surroundings above the template film (waveguide cladding), which was independently detected as a change in the critical angle from θc-PBS to θc-ethanol as marked by arrows in Figure 4B.[27] This change in cladding alone should produce shifts of TE1 and TM1 modes to 56.7° and 50.4°, respectively, as predicted by the Fresnel calculation. However, the final angles obtained in ethanol were 0.3 and 0.2°, higher than the changes due to coupling alone, for TE1 and TM1 modes, respectively. This verifies the exchange between the PBS solution and the optically denser ethanol in the pores. Here one also sees a stronger shift in p-polarization (Fig. 4B) than in s-polarization (Fig. 4A) when the process occurs inside the film. However, the insufficient number of guided modes did not allow for a precise calculation of the volume fraction of the pores filled by each liquid media.

In summary, thin films of BCPs with cylindrical microdomains oriented normal to the substrate surface were successfully coupled in the Kretschmann configuration, and their functions as optical waveguiding layers were demonstrated. Pore generation and exchange of pore materials inside the BCP films were monitored by shifts in the coupling mode angles. Similar experimental strategies applied to porous aluminum oxide templates demonstrated the subangstrom sensitivity of OWS for such processes.[27] Fresnel fitting to the observed waveguide patterns allowed for precise evaluation of the dielectric constants and film thicknesses independently. Based on the methodology established by this work, fabrication of one-dimensional nanostructures (e.g., nanowires or nanotubes) inside, or functionalization/adsorption on top of BCP templates can be investigated by static or in-situ kinetic-mode OWS. Moreover, organic waveguide materials with controlled periodic nanostructures and tunable dielectric constants can be developed for integrated optical devices. The porous BCP templates, which were shown to be stable in PBS solution and ethanol, also show promise for use as a versatile model platform by which interactions with biological substances (e.g., proteins or cells) can be probed. This work has discussed initial steps towards the use of BCP templates in applications such as optical sensing and biosensing.

**Experimental**

3-MPS was purchased from Lancaster Synthesis GmbH. Tetramethoxysilane (TMOS), Fluka GmbH, was used as the sol-gel precursor for the silicon dioxide layers. A typical solution for the precursors was composed of a mixture of 163 μL H2O, 55 μL methanol, 81 μL 0.1 M HCl, and 20 μL TMOS. A commercial grade glass (n = 1.85; LaSFN9, Schott Glass GmbH) was used as a substrate onto which block copolymer thin films were fabricated.

PS-b-PMMA was prepared by using atom transfer radical polymerization as reported previously[28]. The volume fraction of the PMMA block was 0.30, and the weight-average molecular weight (Mw = 55500 with a polydispersity index of 1.19. Atatic PMMA homopolymers with Mn = 23600 were purchased from Polymer Sources, Inc. A hydroxy end-functionalized random copolymer of styrene and methyl methacrylate, denoted PS-r-PMMA, with a styrene fraction of 0.6, was synthesized in bulk via a TEMPO (2,2,6,6-tetramethylpiperidinyloxy) “living” free radical polymerization[19,20]. The molecular weight was determined to be Mw = 9600 with Mn/Mw = 1.80 (Mn = number-average molecular weight) determined by size exclusion chromatography.

AFM images were obtained using a Digital Instruments Dimension 3100 scanning force microscope in the tapping mode. Etched silicon tips on a cantilever (Nanoprobe) with spring constants ranging between 40.0 to 66.0 N m−1 were used.

A He–Ne laser with λ = 632.8 nm and 5 mW power was used to excite plasmon surface polaritons at the metal/dielectric interface. A polarizer for either s- or p-polarization was used for OWS. Both transverse electric (TE) and transverse magnetic (TM) modes (corresponding to s- and p-polarizations, respectively) were excited in the waveguide layers. To observe optical waveguide modes, reflectivity (R) was measured as a function of the incidence angle (θ). The chopped beam was reflected off the gold-coated sample, which was mounted on a θ/2θ goniometer, and the reflected intensity was then monitored by a photodiode that was read out by a lock-in amplifier.

Fresnel calculation was performed using the WINSPALL software (version 2.20) developed in the Max Planck Institute for Polymer Research in Mainz, Germany.

The block copolymer thin films were irradiated by deep UV light with a wavelength of 254 nm at a dose of 25 J cm−2 (XX-15S; UVP,
Inc.) under vacuum for 30 min to selectively degrade the PMMA and crosslink PS, leaving behind the porous PS template.

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Self-Organization of FePt Nanoparticles on Photochemically Modified Diblock Copolymer Templates**

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Phase-segregated block copolymers have received significant attention in the last decade as enabling materials for future technologies.\[^{[1–6]}\] Their value to nanotechnology derives from the expedient tunability of the size, shape, and periodicity of the self-assembled domains by means of manipulating molecular characteristics. Recently, the potential opportunities for block copolymer applications have been bolstered by new methods, which give fine control over long-range ordering of the microdomain structures.\[^{[4,5,7,9]}\] Thin polymer films, by themselves, have limited device applications, but myriad functions can be addressed with hybrid hard/soft matter systems in which the organic layer is used as a scaffold for nanoscale organization of inorganic materials. Of specific interest is the interaction of surfactant-mediated colloidal nanoparticles and diblock copolymer films because the nanoparticle-capping molecules can be tailored to exhibit preference for one of the polymer blocks.\[^{[10–13]}\] This hierarchical approach to create ordered nanostructures removes the linear correlation of size and patterning time associated with traditional lithographic techniques by self-assembling the entire surface in parallel. Also, the spatial limits of lithography can be transcended and the approach can potentially be adapted to industrial-scale processing. An alternative approach to the colloidal nanocrystal methodology is to use spherical- or cy-

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[21] A decrease of $\varepsilon_2$ from 2.220 to 2.048 was calculated from comparison of the dielectric constant of the initial film ($\varepsilon_2 = 2.402$, $\varepsilon_3 = 2.409$) and that of the porous template ($\varepsilon_2 = 2.377$, $\varepsilon_3 = 2.385$). Then, the best fractional combination of air and PMMA was deduced to fit the effective constant of 2.048, using Equation 2.
[22] The film was subjected to the exposure conditions used to etch away PMMA domains in a thin PS-b-PMMA film of thickness $\sim$3000 Å, which is about ten times thinner than the thickness used in this study.
[24] Note that the values of $\varepsilon_{PBS}$ and $\varepsilon_{BLOH}$ are 1.774 and 1.845, respectively.
[28] The film was subjected to the exposure conditions used to etch away PMMA domains in a thin PS-b-PMMA film of thickness $\sim$3000 Å, which is about ten times thinner than the thickness used in this study.
[30] Note that the values of $\varepsilon_{PBS}$ and $\varepsilon_{BLOH}$ are 1.774 and 1.845, respectively.
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[35] Note that the values of $\varepsilon_{PBS}$ and $\varepsilon_{BLOH}$ are 1.774 and 1.845, respectively.