

Research Article

Reversible Shaping of Microwells by Polarized Light Irradiation

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In the last years, stimuli-responsive polymeric materials have attracted great interest, due to their low cost and ease of structuration over large areas combined with the possibility to actively manipulate their properties. In this work, we propose a polymeric pattern of soft-imprinted microwells containing azobenzene molecules. The shape of individual elements of the pattern can be controlled after fabrication by irradiation with properly polarized light. By taking advantage of the light responsivity of the azobenzene compound, we demonstrate the possibility to reversibly modulate a contraction-expansion of wells from an initial round shape to very narrow slits. We also show that the initial shape of the microconcavities can be restored by flipping the polarization by 90°. The possibility to reversibly control the final shape of individual elements of structured surfaces offers the opportunity to engineer surface properties dynamically, thus opening new perspectives for several applications.

1. Introduction

In recent years, the ability to pattern large areas of stimuli-responsive materials at the micro- and nanoscale has opened the opportunity to engineer surface structures and trigger peculiar properties such as complex optical functionalities [1] or surface properties [2] by light-matter interactions. Over the last two decades, the progress on micro- and nanoscale research has enabled the development of several patterning techniques, including soft lithography [3], nanoimprint lithography [4], interference lithography, and advanced photo lithography [5]. Micro- and nanofabrication technology finds applications in a plethora of fields, such as optics and photonics [6], microfluidics [7], and even biology [8].

However, despite such impressive achievements, most of the conventional patterning techniques result in static structures, whose features cannot be modified after fabrication. To overcome such limitation, Karageorgiev et al. proposed exploiting the photoresponsivity of azopolymers in a technique named Directional Photofluidization Lithography [9–11], although the term “photofluidization” is controversial as the fundamental mechanism underlying such effect is still under debate [12]. Based on the same approach, here we propose a reversible manipulation of prefabricated azopolymermic structures driven by polarized light.

Azopolymers are well known to transduce a visible light stimulus into a mechanical modification due to cyclic isomerization of the azobenzene molecule triggered by photon absorption [13]. The cyclic isomerization results in a mechanical modification of the polymer [14] and sometimes in an effective mass migration [15] that allows structuring the polymeric surface via direct illumination [16]. As mass-migration occurs parallel to the polarization of the incident radiation, prefabricated microstructures can be anisotropically deformed depending on the illumination conditions [17, 18]. In detail, we start with an array of microwells and show that the shape of the microwells can be finely controlled by exposing the film to properly polarized light. If linearly polarized radiation is used, the initial shape of microwells can be restored by turning the polarization by 90°. The technique presented here represents a valuable approach to tune the features of prestructured surfaces.

2. Materials and Methods

Sample Preparation. The azopolymer considered is an azopolyelectrolyte (poly{1-[4-(3-carboxy-4-hydroxyphenylazo)benzenesulfonamido]-1,2-ethanediyl, sodiumsalt}) named as PAZO (molecular weight $[M_w] \approx 369,33$ g/mol), supplied by Sigma-Aldrich. The PAZO is mixed with

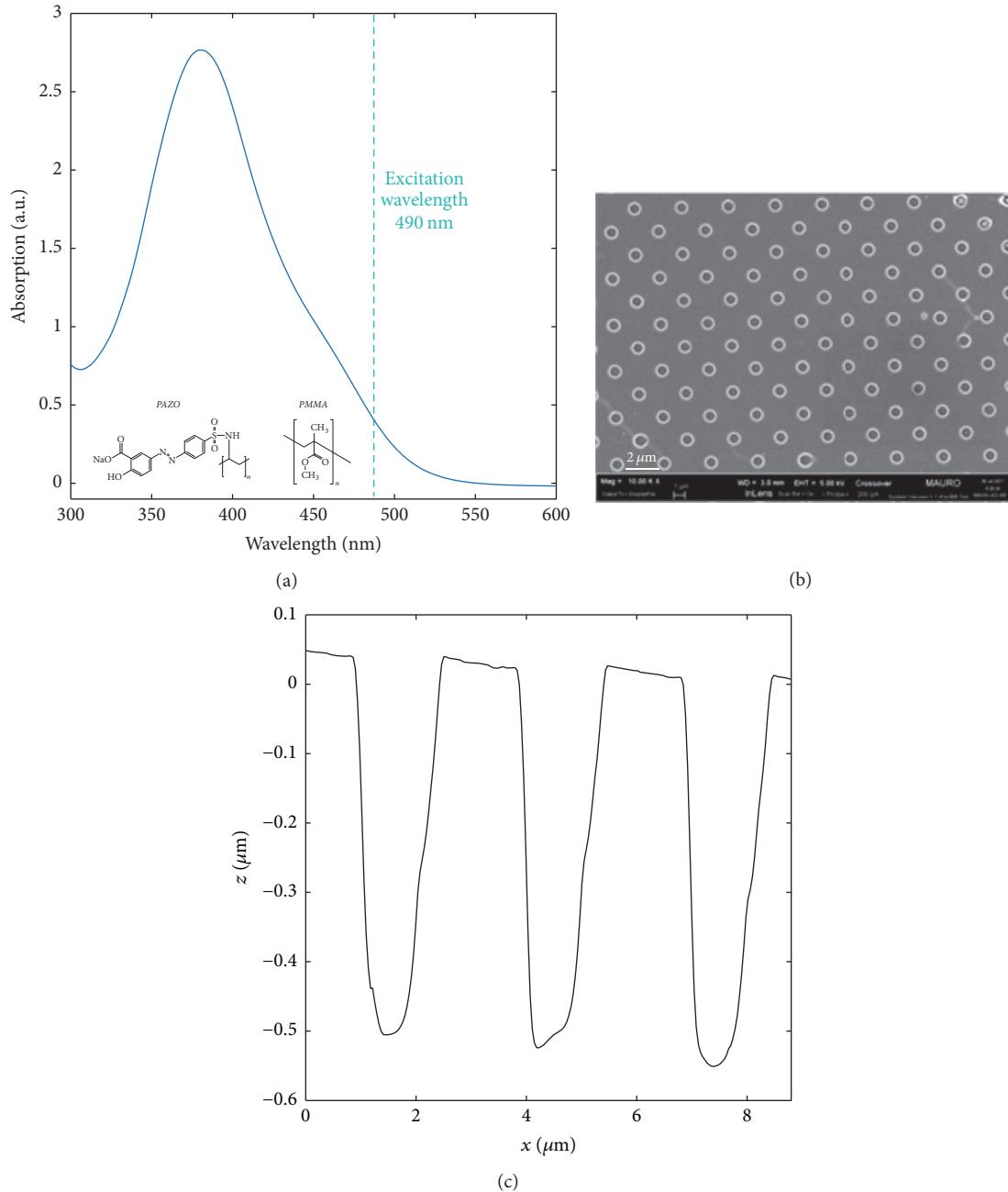


FIGURE 1: (a) Absorption spectrum of the PAZO-PMMA compound in the UV-Visible range. In the inset, sketch of the chemical structure of the two polymers. (b) SEM image of the array of microwells. (c) AFM profile of the microwells.

poly-(methyl methacrylate) (PMMA), resulting in a mixture prepared as follows: PMMA powder (molecular weight $[M_w] \approx 15000$ g/mol by GPC[®]) is dissolved in N,N-dimethylformamide at a concentration of 4 wt%. Nine units of PAZO ($M_w \approx 369,33$ g/mol), dissolved in methanol at concentration 25 mg/mL, are mixed to one unit of PMMA solution and then mechanically stirred and sonicated for few minutes until a homogeneous yellow blend is formed. Figure 1(a) shows the absorption spectrum of the PMMA-PAZO compound obtained and, in the inset, the molecular structure of PAZO and PMMA. We measured the glass

transition temperature (T_g) of the polymeric compound by means of Differential Scanning Calorimetry (DSC, data not shown) and we found $T_g = 53.9^\circ\text{C}$ as compared to $T_g = 60.7^\circ\text{C}$ of the PAZO without PMMA. An amount of 15 μl of PAZO-PMMA is spin-casted onto a glass substrate, previously washed in acetone, rinsed with isopropanol, and dried with a nitrogen flow. A PDMS stamp (10:1 ratio of the prepolymer and the curing agent, provided by Sylgard[®] 184, Dow Corning, cured at 60°C for 2 hours) is soaked in ethanol for few minutes and then gently pressed on the PAZO-PMMA film on glass. The sample is placed in a

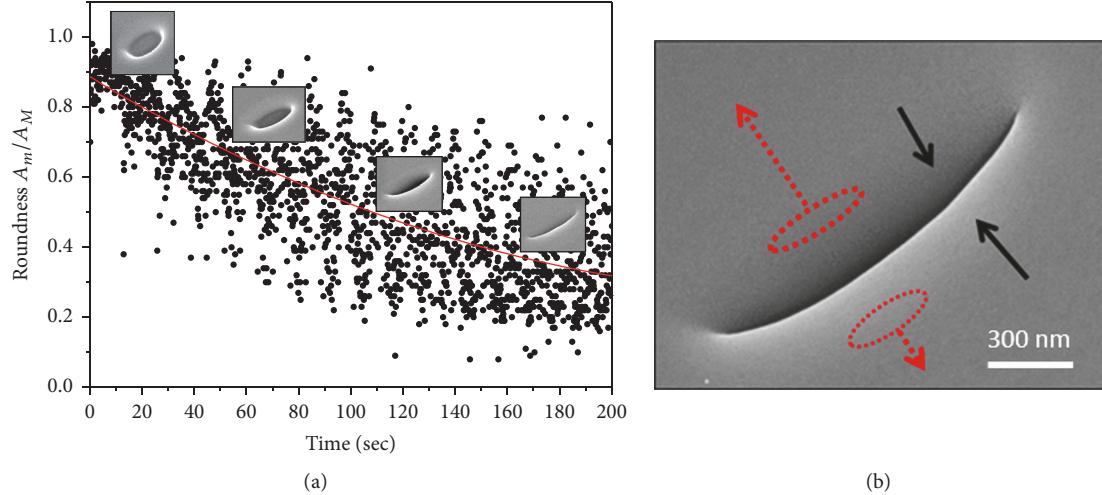


FIGURE 2: (a) Time-resolved roundness values for microwells during laser irradiation (black dots: roundness of individual wells, red line: mean roundness). In the insets, representative SEM images taken with sample tilted by 45° of four different time steps. (b) SEM image (sample tilted by 45°) of a single microwell after irradiation (black arrows: the direction of the pore reduction and red arrows: direction of the mass migration of the material).

vacuum desiccator for 2 hours to enhance patterning and let the polymer blend to fill in the PDMS template. As a next step, the sample is dried in oven, until the solvent is completely evaporated (3 hours at 60°C) before the final removal of the PDMS stamp. The SEM image in Figure 1(b) shows an array of azopolymeric microwells over a large area. The wells are circular with a diameter of 1 μm arranged as a squared lattice with periodicity $\Lambda = 3 \mu\text{m}$. Atomic Force Microscopy analysis reveals circular wells with a diameter of 1 μm arranged as a squared lattice with periodicity $\Lambda = 3 \mu\text{m}$. The depth, estimated by AFM analysis, is about 550 nm, as shown in Figure 1(c).

Optical Setup. The optical setup employed in this work is described in detail in [19]. Briefly, we employ a pulsed laser (laser temporal width ≈ 10 ps, wavelength 490 nm, and average intensity ≈ 50 mW) circularly polarized. The laser beam is slightly focused in such a way that the spot size on the sample surface is about 20 μm, thus resulting in a density of power of 1.6 kW/cm². We point out that the pulsed laser has been employed because of its tunable frequency that allows for a better matching with the absorption band of PAZO (see Figure 1(a)). Due to the slow thermal relaxation time of the isomerization transition (typically in the range of hundreds of ms [15]) as compared to the repetition rate of the pulsed laser (80 MHz), it seems reasonable to approximate the pulsed radiation to a CW light. The beam can be eventually filtered by a linear polarizer placed on a rotational stage to obtain linearly polarized light along arbitrary directions. A 100x objective (NA = 0.95) coupled with a tube lens allows imaging the sample surface on a CMOS camera (Thorlabs DCC3240M). In collection, the laser beam is filtered out by an edge filter (FEL550 from Thorlabs) along the imaging arm. In order to extract useful parameters, the images collected have been binarized by

setting a proper threshold and subsequently analyzed by employing an ImageJ tool able to compute roundness and the overall area of the pores.

3. Results and Discussion

First, we want to investigate the effect of irradiation with linearly polarized light on the azopolymeric microstructures. In agreement with our previous experimental findings [19] and results from literature [20, 21], we observe a directional mass migration of the polymer along the polarization direction. For a quantitative analysis, the mean roundness parameter is evaluated for each frame as the ratio between the minor (A_m) and the major axis (A_M) averaged over 20 microwells within a single field-of-view of the microscope. Initially, wells are almost circular, with a mean roundness of about 0.9 (Figure 2(a)). Upon linearly polarized irradiation, the wells deform into an elliptical shape (mean roundness down to 0.3). The SEM image in Figure 2(b), obtained keeping memory of the orientation of the sample during irradiation, reveals that the major axis of the ellipses is oriented perpendicular with respect to the polarization direction of light, confirming that the photoinduced mass migration of the material occurs along the electric field direction.

In order to show the reversibility of the deformation, we repeated the experiment with linearly polarized light. The microwells, initially showing circular shape (Figure 3(a)), turn into almost closed slits after about 220 seconds of irradiation (Figure 3(b)). Subsequently, we rotate the polarization of light by 90° and we observe a gradual restore of the circular shape (Figure 3(c)). The image analysis reveals that an average roundness of about 1 is achieved after further 200 seconds of irradiation. Such reversible modification is advantageously promoted by an improved stiffness of the polymeric mixture due to the presence of a stabilizing PMMA component, as

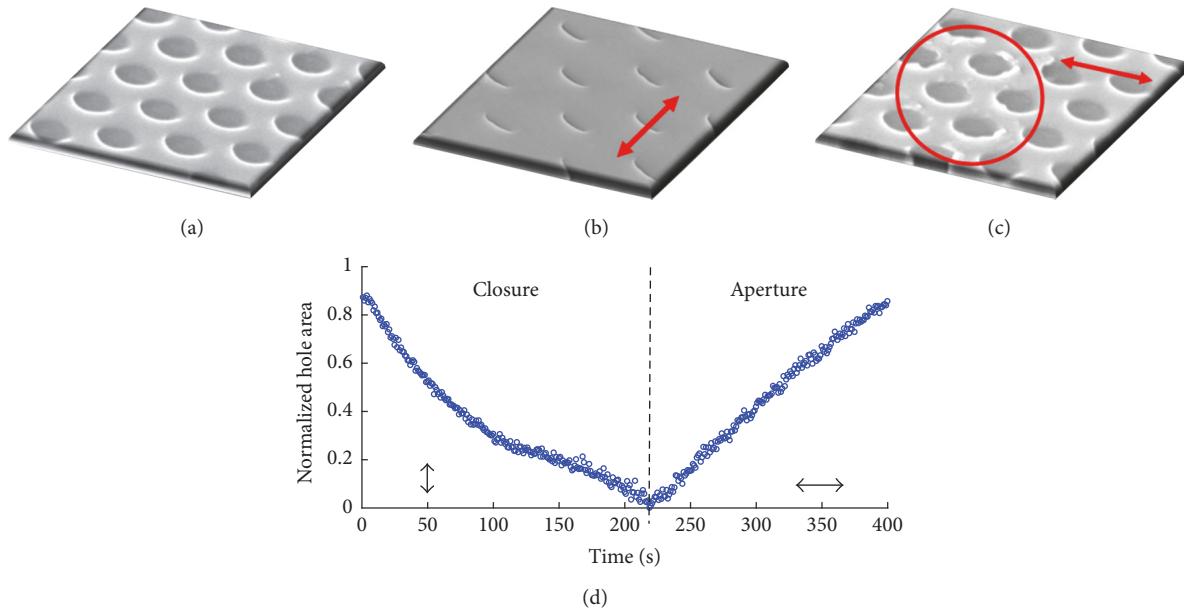


FIGURE 3: Photoreconfiguration of pristine microwells arrays by using linearly polarized light: (a) original concavity shape, (b) closed slits, and (c) restored shape. (d) Time-resolved hole area values for pores during laser irradiation with time-varying polarization states.

suggested elsewhere in other copolymer matrices containing azobenzenes [19–22]. We underline here that the role of PMMA seems to be crucial for reversibility, since with pure PAZO structures we were not able to restore the pristine shape.

In Figure 3(d) we plotted the average area of the holes versus time of irradiation. The graph shows that upon irradiation the holes area is actually reduced till a complete closing of the holes is obtained (black dashed line in the graph). Experimentally, we find that the polarization rotation by 90° triggers a recovery of the microwells area. Such effect is similar to photoinduced micropillar deformation reported elsewhere [17], but the fundamental mechanism underlying the observed phenomena is not straightforward and requires further investigation.

After probing the effect of linearly polarization on the holes shape, we aim to exploit circularly polarized light to reduce the pore size eventually until complete closing. Figure 4 shows the effect of circularly polarized light on an array of microwells. The graph indicates that irradiation with circular polarization does not affect the roundness of the microstructures that is kept constant around a mean value of 0.8. The inset in the graph shows optical images of a sequence of microwells individually addressed and irradiated with different doses (the position along the timeline corresponds to the irradiation time). The sequence shows that although the roundness is preserved, the wells tend to isotropically reduce their size until complete closure. In the case of circularly polarized light, we did not observe any reversibility related to the handedness of light polarization.

4. Conclusions

The presented work shows the possibility to exploit directional photofluidization to manipulate microstructures after fabrication. Starting from an array of circular microwells,

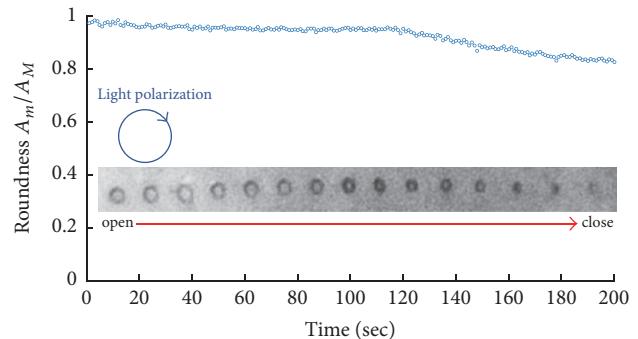


FIGURE 4: Time-resolved roundness evolution microwells during laser irradiation with circularly polarized light. In the inset, optical images of individual microwells are exposed to different light doses.

we employ linearly polarized light to change them into elliptically shaped wells and eventually into narrow slits, depending on the light dose. Alternatively, the microwells can be reduced in size without changing the roundness till an almost complete closing is reached. Tuning the morphology of surface microstructures might provide a smart engineered platform where surface properties can be engineered on demand after fabrication [19, 20]. The light-induced contraction and expansion based reshaping of polymeric microstructures shows exciting potential for several applications including microfluidics and lithography and, in the last years, is showing its potential application in biology, where cells behavior can be regulated in response to material manipulation.

Conflicts of Interest

The authors declare that there are no conflicts of interest regarding the publication of this paper.

Acknowledgments

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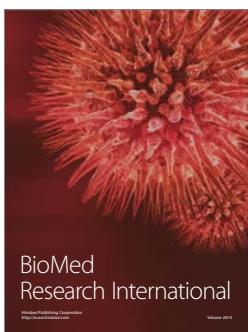
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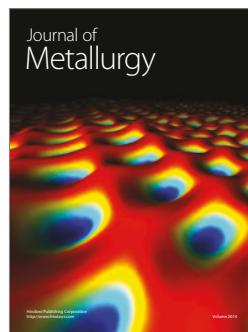
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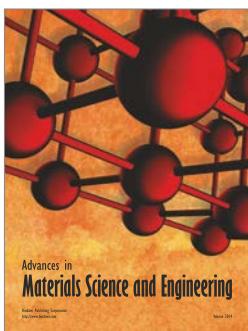
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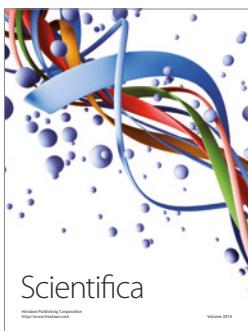
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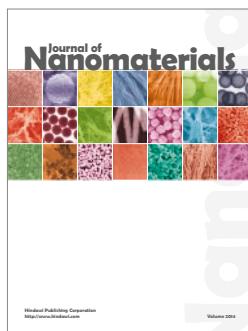
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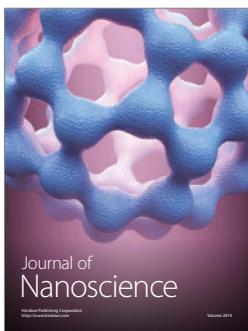
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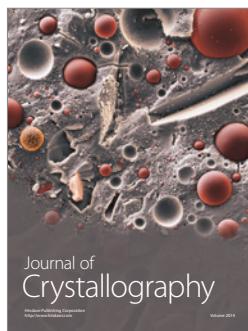
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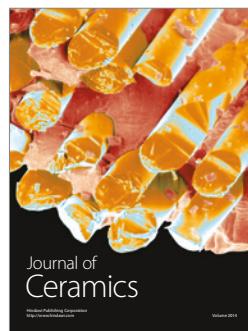
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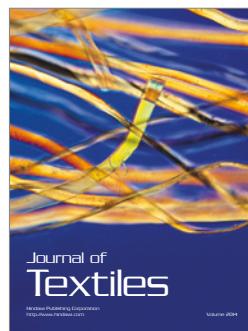
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