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Published in: Journal of Micromechanics and Microengineering

Link to article, DOI: 10.1088/0960-1317/21/11/115008

Publication date: 2011

Document Version Publisher's PDF, also known as Version of record

Link back to DTU Orbit


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Controlled deposition of sol–gel sensor material using hemiwickning

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Received 1 June 2011, in final form 16 August 2011
Published 5 October 2011
Online at stacks.iop.org/JMM/21/115008

Abstract
Optical sensors are fabricated by depositing liquid sol–gel sensor material on a polycarbonate surface, which has been decorated with arrays of periodic micropillars. Using the principle of hemiwickning, the liquid material is spread, guided by the surface structures, to homogeneously fill the volume between the surface structures and form a liquid film with a thickness determined by the height of the micropillars. After evaporation of solvents, a uniform layer of sensor material resides on the surface. This fabrication method enables easy and reproducible deposits of isolated spots of different sensor materials of precise thickness to be made on plastic surfaces, and it provides an improved method for fabricating cheap optical sensors integrated in disposable lab containers.

(Some figures in this article are in colour only in the electronic version)

1. Introduction

In reagent-based optical sensing systems, a change in the optical response of an intermediate agent, e.g., an analyte-sensitive dye molecule, is used to monitor the analyte concentration. Such reagent dyes can be immobilized in a solid matrix of sol–gel silica that is permeable to the analyte, thus making an optical sensor material. In a liquid form, sol–gel-based materials can be coated on a wide range of substrates, which makes this kind of optical sensor materials very versatile [1].

In biotech applications such as, e.g., cell culturing [2], constant monitoring of pH and dissolved oxygen is of great importance. Sol–gel-based sensor materials for measuring these analytes [3, 4] enable fabrication of cheap optical sensors, which can be integrated in disposable lab ware, such as cultivation bags, shake flasks and petri dishes. Using integrated sensors, invasive probes are avoided and sources of contamination are reduced, since the sensors are disposable and optically read from the outside of the container.

Coatings of sol–gel material can be produced using dip-, spray-, or spin-coating [5–7], and homogeneous thin films of sol–gel sensor material, which are required for obtaining fast and uniform response times of the optical sensors, have been demonstrated on dip-coated substrates [4]. However, for deposition of sensor material on the inside of containers, or when several different sensor materials are to be deposited in separate areas on the same surface, these coating methods are not feasible. In present commercial applications of this sensor technology4, 5 patches with sensor material are fabricated and manually glued to the inside of containers.

We present a method for deposition of liquid sensor material on a homogeneous layer in a well-defined region of a surface. By decorating the surface with periodic micropillars and using the principle of hemiwickning [8], a drop of liquid material that is deposited on the microstructured surface will spread, guided by the structures, to homogeneously fill the
volume between the structures. Spreading of the liquid is
governed by the geometry of the microstructures and surface
energy, and the thickness of the deposited film is determined
by the height of the micropillars and is thus independent of
the volume of the deposited drop. This enables easy and
reproducible deposits of spots of the sol–gel sensor material
of precise thickness to be made on plastic surfaces.
Microstructuring of the inside of blow-molded plastic
containers may be performed using step-and-stamp imprint
lithography [9], and for plastic components produced by
injection molding, microstructures can be integrated directly
in the mold [10]. Both of these fabrication methods are suited
for large-scale industrial production.
Spreading by hemiwicking thus enables direct, controlled
deposition of spots of the sol–gel sensor material inside
containers, and it simplifies the fabrication of optical sensors
in disposable lab ware.

2. Hemiwicking of sol–gel material

Hemiwicking is the phenomenon of liquid being imbibed
and spread by a surface texture, and it has previously been
studied in relation to wetting of surfaces [8, 11, 12] and
controlled spreading of liquids [13, 14]. In this paper, we use
hemiwicking to produce films of liquid sol–gel sensor material
with well-defined thickness. When deposited on a surface,
the sol–gel material is a liquid mixture of solvents, i.e. water
and ethanol, and solutes. After evaporation of the solvents,
the material is a liquid mixture of solvents, i.e. water
with well-defined thickness. When deposited on a surface,
the liquid will hemiwick to produce films of liquid sol–gel sensor material
other side. This will proceed until the drop, which was initially
on top of the pillars, has been drained completely and all the
liquid has been imbibed by the microstructure, provided that
the inter-pillar volume is large enough to contain the drop
volume.

The contact line at the top of the pillars is pinned by the
corner, so that the contact angle \( \phi \) is not limited to \( \theta_{eq} \) but can
take a wide range of values [8, 15]. For narrow structures,
surface energy forces \( \phi = 90^\circ \), producing a flat air–liquid
interface [14]. The pillar height \( h \) thus exclusively determines
the thickness of the deposited liquid layer. The volume of the
deposited drop controls the size of the wetted area, but it does
not affect the deposited material thickness. Hemiwicking can
thus be used for producing liquid films of precise thickness
[8].

In the case of a geometry with \( h/l < \tan \theta_{eq} \), as sketched
in figure 1(c), the liquid front gets pinned when the advancing
contact angle \( \theta_a \) equals \( \theta_{eq} \), and propagation stops before the
next row of pillars is reached [13]. The same situation is found
when a spreading liquid reaches the edge of the structured area.
However, propagation may continue in other directions on the
structured surface, and filling of the surface structure is hence
not sensitive to alignment as long as the liquid is deposited
inside the patterned area.

![Figure 1. (a) Cross-sectional sketch of a drop of liquid deposited on a two-dimensional array of micropillars of height \( h \), radius \( r \) and
period \( d = l + 2r \). Driven by surface energy, a liquid film
self-propagates within the microtexture, spreading the liquid and
draining the drop, which lies as a spherical cap on top of the pillars.
(b) Enlarged view of the dashed box in (a) showing the liquid front
in the case of \( h/l > \tan \theta_{eq} \). The contact angle of the advancing
liquid front is larger than the equilibrium contact angle \( \theta_{eq} \),
and the liquid front will reach the next row of pillars before attaining
\( \theta_{eq} \). The next row of pillars will thus be wetted, and the propagation
continues on the other side. In (c), the same situation is shown for a
different geometry with \( h/l < \tan \theta_{eq} \). In this case, the liquid front
is pinned when \( \theta_a = \theta_{eq} \) and the propagation is stopped. At the top
of the pillars, the contact line is pinned by the corner, and surface
energy forces \( \phi = 90^\circ \), producing a flat air–liquid interface.](image)
To create a uniform film of sensor material by hemiwicking, it is required that the solvent evaporation of the deposited sol–gel material occurs much slower than the spreading of the material. Otherwise, evaporation during spreading will result in inhomogeneous sensor spots. The propagation speed of the spreading liquid film increases with $h/l$ [13], and hence the geometry of the surface structures can be optimized to minimize the spreading time.

3. Optimization of surface structures

In order to find the optimal surface structures for fabricating sensor spots by uniform spreading of sol–gel sensor material, the material was deposited on a series of 7 mm diameter arrays of pillars of different dimensions and periods, which were fabricated by thermal imprint lithography [16] on a polycarbonate foil. Polycarbonate was chosen since it is a common material for laboratory bottles. Using a Krüss DSA 10 drop shape analyzer, the contact angle of the ETEOS-based sol–gel oxygen sensor material described in [4] was measured to be $\theta_{eq} = 12\,^\circ \pm 3^\circ$ on a flat polycarbonate surface. The solvents of the sol–gel material, i.e. water and ethanol, evaporate from a 0.5 μl drop in 10 s.

Four-inch silicon stamps with structure heights ranging between 10 and 50 μm, each containing different square lattice and hexagonal lattice geometries, were fabricated using UV lithography and dry etching. Pillar radii and periods were chosen to make ratios of $h/l$ ranging from 0.06 to 5. For all geometries, $\theta_{eq}$ is smaller than $\theta_c$, thus making hemiwicking possible. A fluorsilane coating was applied to the stamp surface by molecular vapor deposition to prevent sticking of the polymer. Imprints were made in 255 μm thick sheets of polycarbonate at 170 °C and 1.3 MPa for 5 min, before cooling the imprint stack to 120 °C and demolding.

Deposition of the sensor material on the structured polycarbonate foil was performed using an Eppendorf pipette, and best results were obtained by filtering the sensor material with a 0.1 μm PTFE syringe filter and deposition in a clean room to avoid dust defects in the coating. After deposition, the sol–gel material was fully dried and condensed by heating to 110 °C for 18 h.

Figure 2 shows examples of depositions of the sensor material on microstructures of different geometries. Like in [13] and [14], we find that the ability of a given geometry to homogeneously spread the sensor material mainly depends on the ratio of $h/l$, and for structures of equal $h/l$, it is relatively independent of pillar height and lattice pattern, i.e. rectangular or hexagonal. As shown in figure 2, the deposition results can therefore be grouped into four regimes of $h/l$ in relation to the equilibrium contact angle $\tan \theta_{eq} \approx 0.2$.

- For $h/l < \tan \theta_{eq}$, the liquid front gets pinned as sketched in figure 1(c), and propagation beyond the drop is not possible. Evaporation causes uneven material distribution with ring deposits as described in [17].
- For $\tan \theta_{eq} < h/l < 0.5$, metastable pinning by the pillars happens easily, and propagation is slow and very sensitive to the lattice pattern. This favors formation of polygonal shapes of the deposited material, as described in [13]. Unless the deposited volume is small, spreading is incomplete and material is left on top of the surface structures.
- For $0.5 < h/l < 2$, the deposited material spreads independently of the lattice pattern to form a round deposition area. This is in correspondence with [13] for increased $h/l$. However, the material distribution is uneven, with the deposit being homogeneous at the center but denser at the perimeter. The inhomogeneity diminishes as $h/l$ is increased.
- For $2 < h/l$, very homogeneous layers are produced.

Figure 2 shows that the homogeneity of the deposited film increases as the ratio of $h/l$ is increased. We explain this tendency by the increasing propagation speed for increasing $h/l$ [13]. For small $h/l$, the spreading time is large, and much of the solvent evaporates while the material is spreading. The solid content of the liquid material thus increases as the film progresses, resulting in an inhomogeneous deposition. Evaporation-induced redistribution of material, as described in [17], may also play a role. At high $h/l$, the spreading time is small, and less solvent evaporates during spreading. Most of the solvent thus evaporates from the material after a uniform

![Figure 2](image-url)
Figure 3. Spreading area as function of volume of oxygen sensor material deposited on a polycarbonate foil with the optimized surface microstructures. Deposition is performed using a 0.1 μl–2.5 μl-range Eppendorf pipette. The data are corrected for capillary filling of the pipette tip, which added approximately 0.09 μl to the volume reading on the pipette. Error bars show standard deviation. The dashed line shows the expected spreading area \( A = V/ (\beta [1 − \phi_S]) \), where \( V \) is the deposited volume, \( \phi_S = 2\pi r^2/ (\sqrt{3}d^2) \), \( h = 23 \mu m \), \( r = 9.5 \mu m \) and \( d = 30 \mu m \). Axis limits correspond to the volume/area of the microstructures of each sensor spot.

film has been made. Uniform evaporation rates over the film result in homogeneous material deposition, considering length scales larger than period \( d \) of the pillars. Between the pillars, on length scales smaller than \( d \), non-uniform evaporation may cause redistribution of material [17].

Evaporation time depends on the volatility of the solvents and the thickness of the liquid film, and spreading time depends on the distance, contact angle and liquid viscosity. The mentioned regimes of \( h/l \) are therefore only valid for the specific structures and materials.

Based on the uniformity of the depositions of sensor material on different surface structures and the sensing performance of the fabricated sensor spots, the best results were obtained with a hexagonal lattice with \( h = 23 \mu m \), \( r = 9.5 \mu m \) and \( d = 30 \mu m \) and \( h/l = 2.1 \). This structure is further characterized in the following sections.

4. Characterization of material deposition

In figure 3, the spreading area is plotted as a function of four different volumes of the sensor material deposited on the optimized microstructure. The linear relation between volume and area shows that spreading of the liquid only occurs as long as the deposited drop can be drained, and the filling stops once all liquid is inside the pillar volume. The excellent agreement between experimental and theoretical spreading areas indicates that the liquid film is uniformly distributed with a film thickness controlled by the pillar height.

If the volume of the deposited sensor material cannot be contained in the inter-pillar volume of a sensor area, the excess material will not spread but stay where deposited on top of the microstructures, thus producing a non-uniformity. Deposited volumes of the sensor material must, therefore, not exceed the 0.56 μl required to completely fill the 7 mm diameter microstructured area.

Figure 4 shows cross-sectional SEM images of the condensed sol–gel sensor material on the 23 μm high and 19 μm wide polycarbonate structures. The thickness of the sensor material is measured to be \( \ell = 5 \mu m \) at the center between the pillars. It is seen that the vertical sidewalls have been coated by a layer of sensor material with a thickness of the order of a few hundred nanometers, and at the bottom between the pillars, the surface is meniscus-shaped as sketched in figure 5. The material thickness is thus inhomogeneous within each unit cell of the pillar array. However, this inhomogeneity is repeated in every unit cell, so when averaging over length scales larger than the period \( d = 30 \mu m \), the material distribution can be considered homogeneous.

For thin films, fluorescence intensity is proportional to material thickness [18], and measurements of the fluorescence intensity of the sensor material were utilized to get a qualitative evaluation of the thickness homogeneity. Fluorescence images recorded with a CCD camera make it possible to evaluate the homogeneity on different length scales.

Fluorescence images were recorded using a Nikon Eclipse inverted fluorescence microscope, equipped with a metal halide light source and a Photometrics-cooled EMCCD camera, and a Nikon TRITC filter cube was used to isolate the fluorescence signal at 620 nm [19] of the Ru(II)-tris(4,7-diphenyl-1,10-phenanthroline) in the oxygen sensor material. A 4× objective with numerical aperture of 0.13 was used for
Figure 5. (a) During filling of the microstructure, the air–liquid interface is flat, as sketched in figure 1. When the solvent evaporates, the decreasing liquid level moves the contact line away from the top corner of the pillar and forces the contact angle to \( \theta_{eq} \), thus creating the meniscus sketched in (b). (c) As the meniscus descends, a thin film is deposited on the vertical sidewalls of the pillars like in a dip-coating process [5]. The solid content of the liquid increases, as indicated by the darker color. (d) The meniscus-shaped surface is maintained while the solid phase of the sensor material condensates and dries completely at the bottom of the pillars.

Figure 6. Top panel shows five fluorescence images (a)–(e) and the positions on the sensor spot where they were recorded. Lower panel shows intensity histograms of the five fluorescence images. Average peak centers are indicated by vertical dashed lines at intensities 9.20 au (polycarbonate pillars) and 13.70 au (sensor material).

Figure 7. Average histograms of ten different sensor spots. Vertical dashed lines indicate average peak positions at intensities 9.20 au and 13.70 au. The gray shaded area shows the average histogram of a sensor spot with a 4 \( \mu \)m layer of sensor material, which is produced by 20% dilution of the sensor material in ethanol. Horizontal dashed line at 800 pixels marks the level at which the peak widths in figure 8 are evaluated.

All histograms in figure 6, which were taken at different locations on the same sensor spot, are centered at intensity...
Figure 8. For fluorescence images of ten different sensor spots, histograms were produced for centered image sections of increasing area; 4 × 4 pixels, 6 × 6 pixels, 8 × 8 pixels, and so forth up to the full 512 × 512 pixels in each image. For every histogram, the peak center (blue), and the lower peak limit (red) and the upper peak limit (green) at the level of 0.3% of the pixels, corresponding to the 800 pixels level shown in the right panel, were calculated. In the left panel, the average positions of the peak center and peak limits are plotted as a function of the image section area, and the graph thus shows the evolution of the average position of the fluorescence intensity peak as larger and larger areas are evaluated. The right panel shows the average fluorescence peak based on the full images of ten different sensor spots. Dashed lines at $I_{\text{center}} = 13.70$ au, $I_1 = 12.49$ au and $I_2 = 14.64$ au show corresponding levels of the two panels. Shaded areas show the standard deviation.

13.70 ± 0.29 au. This implies that variations in the thickness distributions are equally small for all five areas within the same spot, and that the deposited material is uniformly spread over the structured surface.

Figure 7 shows average histograms of ten different sensor spots, each curve based on the average of five fluorescence images as in figure 6. The gray shaded profile shows the average histogram of a sensor spot with 80% sensor material in ethanol, producing a 4 μm layer thickness between the pillars. For comparison, it thus shows how much the intensity histogram changes when the layer thickness is reduced by 1 μm. The variation between the ten different sensor spots is much smaller and shows a good spot-to-spot reproducibility.

To investigate the influence of the image size on the histograms, for fluorescence images of ten different sensor spots, histograms were produced for centered image sections of increasing size: 4 × 4 pixels, 6 × 6 pixels, 8 × 8 pixels, and so forth up to the full 512 × 512 pixels in each image. For each histogram, the peak center and the width, at the level of 0.3% of the pixels, corresponding to the 800 pixels level shown in figure 7, were calculated. Figure 8 shows the average peak center and width for the ten spots as a function of the image size. The peak position and width are seen to be constant down to image areas of 0.5 mm². Below this area, the increasing trend of the peak center, but constant standard deviation, may be caused by light-source inhomogeneity increasing the excitation light intensity toward the image center. Large deviations from the average and increasing standard deviation are only seen for areas below 0.05 mm², corresponding to approximately 50 unit cells of the pillar geometry. Figure 8 thus shows that, at least above 0.5 mm², the same intensity histogram is obtained irrespective of the image size, and this self-similarity is another indication of good uniformity of the coating. Furthermore, it shows that the fluorescence intensity does not depend on the area, and that reproducible results can be obtained from small exposed areas. From a practical point of view it means that reproducible measurements of fluorescence can be obtained with a small probe, e.g., a laser, pointed randomly on the much larger sensor spot.

5. Sensor characterization

Sensor spots for pH and oxygen sensing were fabricated by depositing the two different sensor materials on a polycarbonate foil with the optimized 7 mm diameter pillar arrays of $h = 23 \mu m$, $r = 9.5 \mu m$ and $d = 30 \mu m$. The performance of the sensors was evaluated as described in [3] and [4].

Five pH sensor spots were characterized by measuring the intensity ratio between fluorescence at 460 and 408 nm ($I_{460 \text{ nm}}/I_{408 \text{ nm}}$) as a function of pH. The result is shown in figure 9 where a sigmoidal function has been fitted to calculate the pKₐ of 6.43 ± 0.08. This value corresponds well to the pKₐ of...
6.28 ± 0.02 given in [3], which was made using five different films produced by dip-coating. The small error bars in figure 9 indicate good spot-to-spot reproducibility.

Equally, on the oxygen sensor spots, excited-state life times τ were measured on 38 different spots in oxygen-free atmosphere τ₀ and ambient air of 20.9% oxygen. The Stern–Volmer constant \( K_{SV} = (\tau_0/\tau - 1)/[O_2\%] \), where [O₂%] is the concentration of O₂ in percentages, was calculated.

In table 1, the results of the sensor characterization are tabulated together with the results obtained in [3] and [4] on dip-coated films. The excited-state life time in oxygen-free atmosphere τ₀ is seen to be larger than for the dip-coated sensors, which is most likely caused by the ten-fold thicker layer of the sensor material deposited on the microstructured surface. The thicker material layer may also explain the slightly larger value of \( K_{SV} \). However, it is worth noting that the standard deviation of τ₀ is only 0.40% compared to the much larger 10% stated in [4]. The standard deviations of \( K_{SV} \) are of equal magnitude for the two deposition methods.

6. Conclusion

In conclusion, we have shown that hemiwicking can be used to spread a liquid sol–gel sensor material on a microstructured polycarbonate surface, and that, after evaporation of the solvents, the condensed material is uniformly distributed over the structured surface. The sensing performance of sensor spots fabricated by hemiwicking was characterized and found to be in good correspondence with the performance of sensors fabricated by dip-coating of the same sensor material. We have thus demonstrated an improved method for fabricating cheap optical sensors that can be integrated in disposable lab ware.

Table 1. Comparison of sensor characteristics for pH and oxygen sensors fabricated by hemiwicking (this work) and dip-coating.

<table>
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<tr>
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<th>This work</th>
<th>Dip-coating [3, 4]</th>
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<tbody>
<tr>
<td>pK_a</td>
<td>6.43 ± 0.08</td>
<td>6.28 ± 0.02</td>
</tr>
<tr>
<td>( \tau_0[\mu s] )</td>
<td>6.061 ± 0.024</td>
<td>4.91 ± 10%</td>
</tr>
<tr>
<td>( K_{SV}[O_2%]^{-1} )</td>
<td>0.080 ± 0.002</td>
<td>0.072 ± 0.002</td>
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References