Patterning of Dendrimer-Like DNA

Jon Swaim

NNIN REU Site: Cornell NanoScale Science & Technology Facility, Cornell University
NNIN REU Principal Investigator: Dan Luo, Biological and Environmental Engineering, Cornell
University

NNIN REU Mentor: Wenlong Cheng, Biological and Environmental Engineering, Cornell

University

Contact: hgval@uab.edu, <u>dl79@cornell.edu</u>

(The full report can be found in the 2006 REU Research Accomplishments. http://www.idex-hs.com/products/product_index.aspx) Here is an abbreviated report for users interested in patterning PDMS by etching...Beth

Abstract

Microwell arrays with a depth and diameter of 5 μ m and lattice distance of 15 μ m were patterned into PDMS using photolithography and reactive ion etching. The patterned PDMS acts as a template, guiding DNA combing in microscale dimensions. The patterned DNA can then be transferred to glass or mica by placing and holding the PDMS stamp against a glass slide.

Experimental Procedure:

Photolithography

PDMS was spun onto silicon wafers to a thickness of 100 μ m, and cured for 90 min at 60°C. To improve the adhesion of the PDMS, silicon wafers were initially treated with air plasma (29 W) at 250 mTorr for 2 min. After curing, SPR 220-7 photoresist was spun onto the PDMS to a thickness of 7 μ m and allowed to soft-bake for 30 min at 90°C. After being exposed on an EV620 contact aligner, the substrates were post-baked at 90°C for 30 min and developed.

Etching

Cured PDMS substrates were treated with O_2 plasma (100 W) at 170 mTorr for 60 sec to improve the adhesion of the photoresist. PDMS samples were then dry etched at 43 mTorr for 30 min using a 1:3 ratio of O_2 to CF_4 . Etch rates were optimized by varying the pressure and ratio of gases in the REI process. Etch rates were determined by attaching Kapton tape to the PDMS before etching, and

then measuring the step height using profilometry. After PDMS stamps were characterized with SEM and AFM. The depth of the microwell array was measured using optical profilometry.

Stamping

Molecular combing experiments were performed as previously described [2]. However, an additional force of 250 mN was exerted on some of the samples as the PDMS stamps were peeled off. Fluorescence microscope images were taken during all molecular combing experiments.

Results and Conclusions:

The fluorine-based REI process developed in this study etched PDMS with a stable and directional etch rate (Figure 1). A 1:3 mixture of O_2 to CF_4 at 43 mTorr was found to anisotropically dry etch PDMS at a rate of approximately 10 μ m per hour. Optical profilometry results revealed well depths of 4.5 μ m.

Molecular combing of λ -DNA produced uniform one-dimensional arrangements of DNA. As previously reported [2], linear strands of λ -DNA were observed to attach at the edges of the wells. However, the arrangements did not possess long-range order. It is hypothesized that the treatment of PDMS with O_2 plasma and/or exposure to n-methyl-pyrrolidine within Shipley 1165 remover altered the surface roughness and perhaps even the hydrophobicity of the stamp, compromising the molecular combing process.

Acknowledgements & References:

I would like to thank CNF, National Nanotechnology Infrastructure Network Research Experience for Undergraduates Program, Nanobiotechnology Center, NSF, Molbel, Wenlong Cheng, Suraj Kabadi, Bert Lannon, Steve Jones and Tom Wester.

- [1] Niemeyer, Christof M. Current Opinion in Chemical Biology (2000) 4:609-618.
- [2] Lee et al. PNAS (2005) 102:51; 18321-18325.
- [3] Paranjape et al. JVST (2002) 20:3 975-982.