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Citation: Analyst, 140(19): 6493-6499

Issue Date: 2015

DOI:

Doc URL: http://hdl.handle.net/2115/60219

Type: article

Additional Information: There are other files related to this item in HUSCAP. Check the above URL.

File Information: c5an00909j.pdf

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An instrument-free, screen-printed paper microfluidic device that enables bio and chemical sensing
An instrument-free, screen-printed paper microfluidic device that enables bio and chemical sensing†

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This paper describes a simple and instrument-free screen-printing method to fabricate hydrophilic channels by patterning polydimethylsiloxane (PDMS) onto chromatography paper. Clearly recognizable border lines were formed between hydrophilic and hydrophobic areas. The minimum width of the printed channel to deliver an aqueous sample was 600 µm, as obtained by this method. Fabricated microfluidic paper-based analytical devices (µPADs) were tested for several colorimetric assays of pH, glucose, and protein in both buffer and artificial urine samples and results were obtained in less than 30 min. The limits of detection (LODs) for glucose and bovine serum albumin (BSA) were 5 mM and 8 µM, respectively. Furthermore, the pH values of different solutions were visually recognised with the naked eye by using a sensitive ink. Ultimately, it is expected that this PDMS-screen-printing (PSP) methodology for µPADs can be readily translated to other colorimetric detection and hydrophilic channels surrounded by a hydrophobic polymer can be formed to transport fluids toward target zones.

Introduction

Microfluidic paper-based analytical devices (µPADs) have gained great attention in many fields such as point of care diagnosis,1 environmental testing,2,3 and food analysis.4 These devices have numerous advantages, including low-cost fabrication, facile application, portability, and environmental compatibility.5 µPAD systems have been applied for multiplex analysis in lab-on-a-chip devices.6 µPADs also do not require external pumps and, by taking advantage of the wicking properties of the paper, a complex flow design for various applications is possible.7 Several low-cost methods for the fabrication of µPADs have been reported including photolithography,5 wax printing,8,9 plasma treating,10 and laser etching.11 Various materials such as SU-8, poly(o-nitrobenzylmethacrylate) (PoNBMA), and octadecyltrichlorosilane (OTS) have been used to pattern hydrophobic barriers and form hydrophilic channels as µPADs on filter paper by photolithography. However, they can be easily damaged because of the flexibility of the support paper. Also, the photolithography method requires lithographic equipment and a rigid mask.12 To reduce costs, several non-lithographic methods such as wax printing, plasma treating and laser etching have been reported for rapid, easy, and high resolution fabrication of µPADs. These methods generally need expensive equipment such as wax printers, plasma oxidizers and CO₂ lasers. This restricts their use for fundamental research and for applications in ordinary laboratories, especially in less industrialized and resource-limited regions. Thus, cost-effective and simple methods to fabricate the µPADs without expensive equipment are highly desirable. An inkjet printing method as a simple and cost-effective alternative to expensive methods for patterning microstructures on filter paper has been developed.13 Although this method is simpler, it is still limited by the requirement for the customized cartridges. Other fabrication methods such as silanization of filter cellulose14 and printing of polymer solutions15 have also been developed which efficiently form hydrophilic channels surrounded by hydrophobic barriers.

In this study, we propose a low-cost, instrument free and rapid fabrication method for µPADs; the method is suitable...
for employment in developing countries and resource-limited settings. We use a screen-printing method to pattern PDMS onto chromatography paper which produces hydrophilic channels with clear hydrophobic barriers. Screen-printing that we use in this paper is also a low-cost and widely available printing technique in which a thick paste ink is forced through a stencil attached to a woven mesh screen.16 We have designed and fabricated several patterns for investigating the performance of the fabrication method. We have also performed several colorimetric tests on fabricated µPADs for quantifying pH, glucose, and protein in both buffers and artificial urine samples.

Experimental

Fabrication of the µPADs

A WHT desktop printing table was purchased from Mino International Co., Ltd (Tokyo, Japan). The WHT desktop printing table has three setting screws to allow movement of substrates in x and y directions. The printing table also has a vacuum pump to fix substrates on a board. Hydrophobic barriers as black zones on a white background were designed using Adobe Illustrator software (Adobe Systems, Inc.). A screen stencil (T-420 nylon mesh with ∼35 µm pore size on an aluminium frame) was ordered from Unno Giken Co., Ltd (Tokyo, Japan). Whatman chromatography paper 1# (200 × 200 mm) was purchased from GE Healthcare Life Sciences Whatman™ (Tokyo, Japan). First, the patterned screen stencil was placed directly on a piece of chromatography paper, and PDMS was rubbed onto the surface of the screen stencil using a squeegee, forcing PDMS past the pores of the woven mesh to form PDMS patterns in the paper (Fig. 1). After rubbing, PDMS can slowly penetrate into the cellulose structures. Therefore, the printed-paper was immediately put in an oven after rubbing. Afterwards, the patterned paper was cured in the oven set at 120 °C for 30 min. The PDMS-penetrated paper was ready for use after removing the paper from the oven and allowing it to cool quickly to room temperature.

Preparation of artificial urine solution

Lactic acid, calcium chloride, magnesium sulphate, ammonium chloride, sodium sulphate, sodium chloride and dipotassium hydrogen phosphate were purchased from Wako Pure Chemical Industries, Ltd (Osaka, Japan). Potassium dihydrogen phosphate, urea and sodium bicarbonate were obtained from Kanto Chemical Co., Inc. (Tokyo, Japan). Citric acid was purchased from Kishida Chemical Co., Ltd (Osaka, Japan). Ultrapure water was obtained from a Millipore water purification system (18 MΩ cm, Milli-Q, Millipore) and used for preparing all solutions and in all assays.

An artificial urine solution was prepared according to the literature.17 In brief, 1.1 mM lactic acid, 2.0 mM citric acid, 25 mM sodium bicarbonate, 170 mM urea, 2.5 mM calcium chloride, 90 mM sodium chloride, 2.0 mM magnesium sulphate, 10 mM sodium sulphate, 7.0 mM potassium dihydrogen phosphate, 7.0 mM dipotassium hydrogen phosphate, and 25 mM ammonium chloride were dissolved in ultrapure water. The pH of the solution was adjusted to 6.0 using HCl (0.1 M).

Visualisation of different pH stock solutions

Thymol blue (TB), methyl red (MR), and sodium hydroxide (NaOH) were purchased from Wako Pure Chemical Industries, Ltd. Bromothymol blue (BTB), and phenolphthalein were purchased from Kanto Chemical Co. HEPES buffer was purchased from Dojindo Laboratories, Ltd (Kumamoto, Japan). For visualisation of the pH assay, a pH-responsive ink was prepared according to the literature.13 Briefly, 0.5 mg of TB, 6 mg of

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Fig. 1 Schematic representation of PDMS-screen-printing for fabrication of the µPADs. (1), (2) Putting the screen directly on the chromatography paper surface; (3), (4) covering the screen with PDMS using a squeegee; (5) penetrating the PDMS into the paper; (6) curing the PDMS-screen-printed paper in an oven set at 120 °C for 30 min.
BTB, 1.2 mg of MR, and 10 mg of phenolphthalein were dissolved in 10 mL of 95:5 (v/v) ethanol/water. Then, 0.01 M NaOH solution was added dropwise into the mixed indicator solution until the colour turned to light green. HEPES buffer (0.1 M) was used to make stock solutions and the pH of stock solutions were adjusted (2–9) by HCl or NaOH addition.

**Glucose assay**

Glucose and glucose oxidase were purchased from Wako Pure Chemical Industries, Ltd, and Sigma-Aldrich Co., Inc. (Tokyo, Japan), respectively. Potassium iodide was purchased from Kanto Chemical Co. The glucose stock solution (1 M) was diluted with the artificial urine solution and adjusted to concentrations of 0, 2.5, 5, 10, 20, 50, 100 and 500 mM. For the glucose assay, a 0.6 M solution of potassium iodide (15 µL) was first introduced into the auxiliary zone, followed by 1:5 horseradish peroxidase/glucose oxidase solution (15 µL; 15 units of protein per mL of solution). After exposing to air for 10 min at room temperature, 0.5 µL of different concentrations of glucose solutions were spotted onto eight separate sample zones.

**Protein assay**

BSA standard solution was purchased from Takara-Bio Co., Inc. (Shiga, Japan). Tetrabromophenol blue (TBPB) was purchased from Sigma-Aldrich Co., Inc. Citric acid was purchased from Hidex Co, Inc. (Osaka, Japan) and trisodium citrate was purchased from Wako Pure Chemical Industries, Ltd. BSA standard solution was diluted with ultrapure water to achieve the desired concentrations (0, 2, 4, 6, 8, 10, and 20 µM). For the protein assay, 15 µL of a 250 mM citrate buffer solution (pH 1.8) was introduced into the auxiliary zone and exposed to air at room temperature for 10 min. Then, a 9 mM solution (15 µL) of TBPB in 95% ethanol was introduced into the citrate buffer solution residue followed by exposing to air for another 10 min. Finally, 0.5 µL of the different concentrations of BSA solutions were separately spotted onto eight sample zones.

**Results and discussion**

**Evaluation of the appropriate channel width**

To determine the minimum resolution of PDMS-Screen-Printing (PSP), we designed a pattern including different channel widths (Fig. 2A). After fabrication, 7 µL of a 0.01 M fluorescein solution was dropped onto the paper to allow the observation of the hydrophobic, hydrophilic, and wicking properties. Then fluorescence images were recorded by using a fluorescence microscope (Keyence BZ-9000, Japan) (Fig. 2B and C). In Fig. 2B, hydrophilic channels (300, 400, and 500 µm) smaller than 600 µm were observed but solvent could not flow through them. The minimum width of the hydrophilic channel surrounded by printed PDMS barriers to deliver an aqueous sample was 600 µm but considering the wicking properties, we recommend designing hydrophilic channels wider than 800 µm (Fig. 2B). Furthermore, as shown in Fig. 2B and Table 1, the printed channels were smaller than the pattern because after

![Fig. 2](image-url) Evaluation of different channel widths. (A) Patterned screen mesh for printing of different channel widths, (B, C) comparison of the printed feature with the patterned screen and tracing the wicking properties of them, (D) SEM image of the bare (left) and PDMS printed paper (right).
for the top side of the device was better than single rubbing but leakage of the indicator solution was still observed for the top and back sides. Furthermore, excess PDMS remained on the stencil. We increased the frequency of rubbing to three times (Fig. 3C-III). This led to PDMS leaking into the hydrophilic areas in the back side. On the other hand, because the total size of the hydrophilic areas was decreased, cross contamination of the sample zones was observed in the top side. So to prevent the leakage, we decreased the amount of PDMS to 10 g and two rubbing times. This result is shown in Fig. 3C-IV. For the back side, there was no leakage of the indicator solution but there was leakage from the top side. Excess PDMS still appeared on the stencil, so we decided to decrease the amount of PDMS to 7 g and use three rubbing times (Fig. 3C-V). Fig. 3C-V shows good penetration of PDMS solution deep into the cellulose structures with no leakage of the indicator solution from the printed channels. We concluded that the optimum conditions for screen-printing of PDMS for this pattern were: 7 g PDMS, three rubbing times, and curing at 120 °C for 30 min.

In the current study, production of 36 μPADs by one screen-printing of PDMS solution on a piece of chromatography paper was possible. The cost for the paper and an aluminium frame is ~$8 (US) per 100 cm², so mass production of the μPADs is possible at a reasonable cost. Moreover, our fabrication method using thermo-curable PDMS does not require an organic solvent for adjusting viscosity and controlling the penetration properties.21

In order to investigate the performance of the μPADs, results of different pH solutions were obtained (Fig. 3D). First, 0.5 µL aliquots of the different pH solutions (2–9) were separately spotted in the sample zones, and were allowed to dry at room temperature for 10 min. Then, 15 µL of the pH-responsive ink was spotted in the auxiliary zone. From Fig. 3D, we concluded that it was possible to detect the pH of an unknown solution as a strip test, visually. Significantly, using the auxiliary zone in this pattern allowed the pH of samples from alkaline to acidic conditions, to be seen simultaneously. Furthermore, the cured PDMS was compatible with alkaline and acidic conditions because no leakage of solution was observed. This result showed the capability of the μPAD for assays in a pH range from 2 to 9.
Glucose and protein assays

µPADs, as mentioned earlier, have great potential for applications in various biochemical assays. Here we applied our method to two important biochemical assays: glucose and protein assays (Fig. 4). We prepared solutions with known concentrations of glucose in artificial urine and BSA standard solutions, and performed the colorimetric assays. The results showed that the µPADs fabricated using the current method were applicable for the determination of 5 mM glucose in artificial urine which is adequate for detecting the critical concentration of glucose in diseases such as glucosuria. This concentration was easily detectable by observation and could also be quantified using a hand held camera and a simple image processing step. The assay was repeated several times and reproducible results were achieved.

We also tested a simple colorimetric assay for measuring the protein concentration by our µPADs. Similar to the glucose assay, intensity of the colour was checked by observation or by capturing an image and quantification of the signal using open source imaging software (ImageJ) (Fig. 4C and D). Limit of detection for BSA was 8 µM. The test can be applied to quantify protein in urine in nephrotic syndrome where, the concentration of protein is higher than 35 µM. In the current setting, detecting different concentrations of protein ranging from 5 to 100 µM is possible.

Conclusion

We used a simple, low-cost, and widely available screen-printing method to fabricate µPADs and we investigated the per-
formance of this method using typical colorimetric detection for glucose and protein. We used PDMS to form clear hydrophobic borders on conventional chromatography paper. High resolution micro channels were fabricated without using any printing machine such as jet injection printers. We tested the fabricated µPADs for different chemical and biochemical sensing assays.

Acknowledgements

We acknowledge helpful discussions with Prof. Daniel Citterio concerning pH assay. Saeed Mohammadi thanks the international education program (Advanced Graduate School of Chemistry and Materials Science: AGS) of Graduate School of Chemical Sciences and Engineering, Hokkaido University and the Japanese Government (MONBUKAGAKUSHO: MEXT) scholarship.

References


Fig. 4 Quantification and visualisation of glucose and protein assays. (A, B) Quantification results (0–100 mM) and a µPAD used to visualise a positive test for glucose in artificial urine (0–500 mM), respectively. (C, D) Quantification results (0–20 µM) and a µPAD used to visualise a positive test for BSA standard solution (0–30 µM), respectively. Each datum for the quantification results is the mean of three values for glucose and four for BSA; error bars represent the relative standard deviation of the measurements.