A Biomimetic Approach for the Creation of Two-Dimensional Microscale Surface Patterns: Creation of Isolated Immunological Synapses

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Accessibility
Current efforts in surface functionalization have not produced a robust technique capable of creating specific two-dimensional microscale geometrical arrays composed of multiple proteins. Such a capability is desirable for engineering substrates in sensing and cell patterning applications where at least two different protein functionalities in a specific configuration are required. Here we introduce a new approach for the creation of arrays of microscale geometries. We demonstrate our approach with a biomimetic structure inspired by the immunological synapse, a cell-cell interfacial structure characterized by two concentric rings of proteins: an outer adhesion protein structure and an inner recognition ligand core. The power of the technique lies in its ability to pattern any protein in any defined geometry as well as to create arrays in parallel.

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1. Introduction

Bioactive surfaces consisting of patterned proteins in specific defined geometries are important for screening applications [1], sensing [1–5], the study of cell-cell interactions [2, 6, 7], and the creation of cellular networks [2, 5, 8–10] or cellular aggregates for tissue engineering applications [11, 12]. A long sought after capability is the ability to fabricate patterned surfaces with cellular-like interfacial features (e.g., cell recognition, adhesion, and stimulatory ligands) to enhance cell-substrate interactions. Such biomimetic approaches require surface engineering with multiple chemical functionalities to facilitate attachment of a variety of proteins and to date, methodologies have been lacking in the robust production of arrays with multiple functionalities. Although surfaces capable of binding proteins at high densities are commercially available [13], these slides are designed for protein microarray screening and, in turn, require method development if specific patterns are required.

Current molecular patterning techniques such as poly(dimethylsiloxane) (PDMS) molding [9, 10, 14] are harsh in their application, potentially compromising surface functionality. Several groups have developed photo- and chemically-sensitive materials for protein patterning [5, 7, 15–17]; however, the techniques all require noncommercially available molecules and most have not yet been demonstrated to be capable of patterning multiple bioactive proteins [15–17]. Novel patterning approaches have also been developed, including microfluidic-directed patterning [18] and dip-pen lithography (DPL) [3]. Despite their promise in a number of applications, serial techniques such as DPL are not currently suited for the creation of large patterns and arrays and microfluidic patterning places strict geometric limitations on patterns.

Here we present a novel, photolithographic-based approach to create surfaces consisting of arrays of functional patterned proteins that uses only commercially available materials, as opposed to previously proposed methods [7, 15–17, 19–21]. We demonstrate our approach with a biomimetic structure inspired by the immunological synapse, an immunologically complex structure that occurs after lymphocytes encounter antigen-presenting cells [22].
The interfacial geometry of this structure is characterized by two concentric rings of proteins: an outer adhesion protein structure and an inner recognition ligand core.

The spatial reorganization of molecules on the surface of immune system cells such as T cells is a hallmark of early T-cell responses to foreign invasion [22]. Following interaction with antigen-presenting cells (APCs) this spatial reorganization takes a specific form, termed an immunological synapse (IS), which facilitates the encoding of information dictating the magnitude of the immune response. The hallmark of (IS), which facilitates the encoding of information dictating the magnitude of the immune response. The hallmark of the mature IS is a “bulls-eye” pattern, which consists of regions termed supramolecular activation clusters (SMACs) [23–25]. The central SMAC (cSMAC) consists of interactions with antigen-presenting cells (TCRs) on the T-cell and peptide-loaded major histocompatibility complexes (pMHCs) on the APC. The peripheral SMAC or pSMAC contains adhesion and costimulatory ligands. Sensitivity is achieved because clustering promotes a high avidity interaction in the cSMAC [23–25]. Thus, creation of APC mimics on a surface would not only facilitate the study of T cell-APC signaling but may enable a recapitulation of sensitive sensing mechanisms employed by T cells in recognition of antigen [6, 7].

Recently, the application of a specially synthesized photoresist was demonstrated for the successful patterning of a surface with IS-like domains [7]. Despite the utility of the approach, this technique is not applicable for the formation of specific and precise patterned two-dimensional IS structures as observed in biological interactions. A cellular-repellent moiety such as poly(ethylene glycol) (PEG) cannot be bound at the periphery of these structures, thus cells adhere across the entire substrate rather than specifically at the ISs, compromising the utility of the approach for specific detection [26, 27].

We demonstrate here a method capable of patterning functional antibodies and PEG in any defined geometry using only commercially available products. We use this technique to create a patterned IS with one antibody conjugated in the central region and another in the peripheral region of the IS, surrounded by poly(ethylene glycol) (PEG) [26, 27]. Our method thus produces a surface with three chemical functionalities, created by two successive steps of traditional photolithographic patterning [28] followed by functionalization with chemical cross-linkers. An attractive feature of this method is that subsequent conjugation to these groups with specific molecules after photoresist removal was performed entirely in aqueous buffers, thus protein functionality is not compromised.

2. Materials and Methods

2.1. Materials. Amine-functionalized microscope slides were purchased from VWR (cat. no. 16001-008) and were cut to 1 in² pieces using a diamond scribe. Photomasks were purchased from Benchmark Technologies and Photo Sciences, Inc and transparency photomasks were purchased from CAD Art Services. All photoprocessing steps were performed in a Class 1000 cleanroom (Harvard Center for Nanoscale Systems). All chemicals used for cleanroom processing were of cleanroom grade. The photoresist used for all processing was AZ 5214-E (MicroChem), an inverting resist. Glutaraldehyde, hydroxylamine, dimethyl sulfoxide (DMSO), phosphate buffered saline (PBS), ethylenedinitrilotetraacetic acid (EDTA), sodium cyanoborohydride, the fluorescein-5-isothiocyanate (FITC) conjugate of bovine serum albumin (BSA-FITC), and mouse IgG were purchased from Sigma-Aldrich; N-succinimidyl-S-acetylthioacetate (SATP), N-(6-(biotinamido)hexyl)-3’-(2’- pyridyldithio)-propionamide (biotin-HPDP), sulfosuccinimidyl acetate (NHS-acetate), N-hydroxysuccinimide (NHS), and 1-ethyl-3-(3-dimethylaminopropyl) carbodiimide (EDC) were purchased from Pierce Biotechnology; amine-terminated poly(ethylene glycol) (NH₂-PEG; 10 kDa) and aldehyde-terminated PEG (CHO-PEG; 10 kDa) were purchased from Polysciences, Inc; FITC, AlexaFluor 568 hydrazide sodium salt (AlexaFluor 568), AlexaFluor 568 C₅ maleimide (AlexaFluor 568 maleimide), streptavidin, and atto565-biotin were purchased from Invitrogen. Mouse IgG was obtained from Echitect-Bio. Antimouse-IgG-FITC was purchased from Zymed, Inc. All other antibodies were purchased from BD Biosciences.

2.2. Alignment Mark Deposition. Lift-off processing was used to pattern metal alignment marks on the slides for subsequent photolithographic steps. Amine-functionalized slides were cleaned with acetone and isopropanol (Baker Chemical Co.) and dried with nitrogen. Photoresist (AZ 5214-E) was spun on the slide at 3000 rpm using a Headway spinner. The slide was then baked at 90°C, exposed with the photomask using an AB-M maskaligner, baked at 110°C, and finally flood exposed. The slide was then developed in MF319 (MicroChem), washed with deionized water (DI; 18 MΩ), and blown dry with nitrogen. The slides were then loaded into a Sharon Systems Electron-Beam Evaporator, where a 100 nm titanium (99.9%, Kurt J. Lesker Co.) deposition was performed. After unloading, the slide was sonicated in acetone to remove the remaining photoresist, leaving only the metal alignment marks. Subsequent patterning steps followed the same procedure (without metal evaporation) with the exceptions that the photoresist was spun at 5000 rpm and that acetone was used to remove the photoresist without sonication.

2.3. Glutaraldehyde Conjugation. The slide containing the first layer of photoresist was treated for 1 hour at room temperature (RT) with a 0.01 M solution of glutaraldehyde in 0.1 M sodium acetate buffer, pH 5.5, with 1 mM sodium cyanoborohydride. Sodium cyanoborohydride was required for reduction of the unstable imine (C=N) bond. For all conjugation steps, the slides were placed face-up at the bottom of a 100 mL beaker with ~10 mL solution (enough to completely immerse the slide in the solution) and the beaker was covered with parafilm and placed on a shaker table. After each conjugation, the slide was removed, rinsed with DI, and blown dry with nitrogen (note after protein conjugation steps the slides were rinsed with PBS and not allowed to dry.
Figure 1: Sequence of photopatterning and chemical treatment steps for binding antibodies and PEG to create surface APC mimics. The steps are thoroughly discussed in the text. Briefly, Step 1 shows the first layer of photopatterning followed by subsequent glutaraldehyde conjugation to exposed amine (NH$_2$) groups. Step 2 is the second photopatterning (following removal of the first layer) with subsequent thiol (SH) conjugation to exposed NH$_2$ groups. In Step 3, PEG is bound to aldehyde (CHO) groups and free SH groups are biotinylated. Step 4A demonstrates IgG conjugation to exposed NH$_2$ groups and biotinylated-IgA binding to surface-bound streptavidin-Ac. Antibody recognition of specific antigens enables fluorescently conjugated antibodies to specifically bind active IgG and IgA. Step 4B shows a similar binding of two fluorescently-labeled IgGs, which create an artificial IS in this geometry.

2.4. SATP Conjugation and Deprotection. N-Succinimidyl-S-acetylthiopropionate (SATP; 2 mg) was dissolved in 0.5 mL DMSO, which was added to 9.5 mL PBS and the slide containing the second layer of photoresist was treated for 30 minutes at RT with this solution. After washing, the slide was treated with a solution of 0.5 M hydroxylamine and 25 mM EDTA in PBS, pH 7.4, for 2 hours at RT to deprotect the sulfhydryl groups.

2.5. NH$_2$-PEG Conjugation. Slides were treated with a 1% (w/v) solution of NH$_2$-PEG in PBS, pH 7.4, with 1 mM sodium cyanoborohydrate for 4–6 hours at RT.

2.6. Biotin-HPDP Conjugation. N-(6-(Biotinamido)hexyl)-3-(2-pyridylthio)-propionamide (Biotin-HPDP; 1 mg) was dissolved in 0.5 mL of DMSO, which was added to 9.5 mL of a 1 mM solution of EDTA in PBS, pH 7.4. The slide was treated with this solution for 1 hour at RT. We obtained more repeatable results when the biotin-HPDP conjugation followed that with NH$_2$-PEG.

2.7. Fluorescent Antibody Conjugate Preparation. Mouse IgG, biotinylated-mouse IgA, antimouse IgG-FITC, and anti-LFA-1-FITC were used as received. Antimouse-IgA was dissolved to 1 mg/mL in PBS, pH 7.4, AlexaFluor 568 hydrazide was added to achieve a final concentration of 0.1 mg/mL, and 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide hydrochloride (EDC) was added at 10 mg/mL. The reaction proceeded for 1 hour at RT with shaking and protected from light. A 50 mL
Amicon Ultra (Millipore, cat. #UFC905024) centrifugal filter tube with a 50000 MW cutoff was used to remove unconjugated dye (4 30-minute spins at 3000 rpm at 4°C) and the resulting antimouse-IgA-AlexaFluor568 was diluted to 1 mg/mL. The same procedure was used for AlexaFluor 568 conjugation to biotinylated-anti-CD3ε. Conjugates were stored at 4°C.

2.8. NHS/EDC Coupling. Mouse IgG or anti-LFA-1-FITC was dissolved to 10 μg/mL in PBS with 10 mg/mL EDC and 5 mg/mL N-hydroxysulfosuccinimide (sulfo-NHS) and reacted with the slide for 1 hour at RT. Washes with protein bound on the chip took place in PBS and the slides were not dried.

2.9. Streptavidin-Ac Preparation. Acetylated streptavidin was used because we found this protein to minimize nonspecific binding on the PEG surface (as compared with pure streptavidin, avidin, and neutravidin). To prepare the acetylated streptavidin, 5 mg/mL (sulfosuccinimidyl acetate) NHS-acetate was added to 1 mg/mL streptavidin in PBS, pH 7.4. The reaction proceeded for 1 hour at RT and the streptavidin conjugate was purified using a 50 mL Amicon Ultra centrifugal filter tube with a 10000 MW cutoff (4 30-minute spins at 3000 rpm at 4°C). The conjugate (streptavidin-Ac) was stored at 4°C.

2.10. Streptavidin-Ac Conjugation. The slides were treated with 10 μg/mL streptavidin-Ac in PBS, pH 7.4, for 1 hour at RT. This conjugation does not affect biotin binding at the concentrations used in this work.

2.11. Biotinylated Antibody Conjugation. Biotinylated-mouse IgG or biotinylated anti-CD3ε-AlexaFluor568 were dissolved to 50 μg/mL in PBS and reacted with the slide for 1 hour at RT (the latter was protected from light). When this reaction was completed, slides treated with anti-LFA-1-FITC and biotinylated-anti-CD3ε-AlexaFluor 568 were dried (gently) and viewed under a fluorescent microscope (Olympus IX81, equipped with the Carv-II unit from BD Biosciences). The low-intensity areas in the bottom of some images (notably Figure 3(d)) are due to the directional nature of the stream of air during this drying step.

2.12. Immunoactivity Study. Slides containing mouse IgG and biotinylated-mouse IgA were treated with a PBS, pH 7.4, solution containing 50 μg/mL antimouse-IgG FITC and 50 μg/mL antimouse-IgA-AlexaFluor 568 for 1 hour at RT. These slides were subsequently dried (gently) and viewed under a fluorescent microscope.

3. Preparation of Isolated ISs

3.1. Surface Chemistry Demonstrations. In Figure 1 (Step 1), photoresist spun on amine-functionalized microscope slides was patterned and the exposed amine groups were converted to aldehydes with glutaraldehyde. Subsequent removal of the photoresist left a slide with two functionalities, with the aldehydes defining areas between the ISs. Successful
glutaraldehyde conjugation was first demonstrated using a transparency photomask consisting of an array of 25 × 25 μm² squares (clear on a dark background), followed by glutaraldehyde treatment. Aldehyde functionality was thereby conferred to all areas of the chip but these squares. The photoresist was subsequently removed and the slide was treated with aldehyde-reactive PEG (NH₂-PEG). After subsequent treatment with fluorescein-5-isothiocyanate (FITC), a fluorescent amine-reactive probe, the slides were fluorescently imaged under a GFP filter (Figure 2(a)). The square fluoresces due to the conjugation of FITC to the amine surface but the formation of an amide bond between the surface aldehyde and free amine of the NH₂-PEG prevents FITC binding in areas surrounding the square.

In Step 2, a second photoresist layer was applied and patterned and the available amine groups were reacted with SATP, resulting in the conversion of exposed amine groups to protected thiols [29]. Photoresist removal with acetone followed by thiol deacetylation with hydroxylamine thus resulted in a slide with three chemical functionalities, where the thiols defined the cSMAC and the remaining amines defined the pSMAC. In order to demonstrate successful SATP conjugation, slides were photopatterned with a transparency mask consisting of an array of 25 × 25 μm² squares (dark on a clear background) followed by SATP conjugation and subsequent deprotection. This resulted in the presence of sulfhydryl groups only in the squares. The photoresist was then removed and slides were treated with amine-reactive PEG (CHO-PEG). A sulfhydryl-reactive fluorescent probe (AlexaFluor 568 maleimide) was then conjugated to the slides and subsequent fluorescent imaging using a TRITC filter showed that the high fluorescence intensity was confined to the squares, demonstrating that sulfhydryl groups are present in these areas (Figure 2(b)).

As shown in Step 3, in order to eliminate nonspecific adsorption of cells to nonsynaptic regions of the slide, NH₂-PEG (10 kDa) was conjugated to the exposed aldehydes. Next, biotin-HPDP was conjugated to the thiols in the cSMAC region, thereby biontinylating this area and readying the slide for protein conjugation [29]. Biotin-HPDP conjugation was demonstrated by processing slides with a transparency mask containing an array of 25 × 25 μm² squares (dark on a clear background), followed by SATP conjugation and deprotection. After photoresist removal slides were treated with CHO-PEG, followed by biotin-HPDP. The slides were then treated with acetylated streptavidin (streptavidin-Ac), followed by treatment with atto565-biotin, a fluorescently conjugated biotin molecule. Slides were fluorescently imaged under a TRITC filter (Figure 2(c)). As in Figure 2(b), the high fluorescence intensity is confined to the square, demonstrating the successful conjugation of biotin-HPDP to the sulfhydryl groups, as well as subsequent streptavidin-Ac and atto565-biotin binding.

3.2. Protein and Antibody Conjugation. Protein conjugation to slides using EDC/sulfo-NHS coupling [29] (Step 4A) was demonstrated by processing slides with a transparency mask containing an array of 25 × 25 μm² squares (dark on a clear background), followed by SATP conjugation and deprotection. After photoresist removal, slides were treated with biotin-HPDP, as described above. BSA-FITC was then
added to the chip with 10 mg/mL EDC and 5 mg/mL sulfo-NHS. The slides were then treated first with streptavidin-Ac and next with atto565-biotin. Fluorescence imaging was performed with both GFP (Figure 2(d)) and TRITC (Figure 2(e)) filters (images are from the same array point). The high fluorescence intensity in the region surrounding the square in Figure 2(d) indicates that BSA-FITC binding was confined to this region, while the high fluorescence intensity in the square in Figure 2(e) shows that atto565-biotin binding was confined to this region.

We next demonstrated the ability of this approach to pattern the surface with functional antibodies (Step 4A). We first conjugated mouse-immunoglobulin G (IgG) [30] to the exposed amine groups through standard EDC/sulfo-NHS chemistry. The slide was then treated with acetylated streptavidin and subsequently with biotinylated mouse-immunoglobulin A (IgA). Unlike other patterning techniques [15–17], our approach does not require the presence of blocking agents during the final conjugation steps. The slide was next incubated with an FITC conjugate of antimouse-IgG and an Alexa Fluor 568 conjugate of antimouse-IgA, washed, and imaged. The fluorescent micrographs in Figure 3 demonstrate the functionality of the conjugated antibodies as well as their segregation into different regions dictated by the surface functional groups. This also enables the creation of arrays of patterns with varying absolute and relative dimensions.

This chemistry can be utilized to functionalize slides with physiologically relevant antibodies to mimic the IS. For this purpose we chose antibodies to two ligands known to be sequestered in the IS: an antibody to leukocyte function-associated antigen 1 (anti-LFA-1) as the adhesion promoter for the pSMAC and an antibody to the ε subunit of cluster of differentiation 3 (anti-CD3ε), a well-established T-cell stimulus, for the cSMAC [7, 25]. As shown in Step 4B of Figure 1, antimouse-LFA-1 (FITC conjugate) was coupled to the free amines in the pSMAC region, followed by treatment of the slide with biotinylated antimouse-CD3ε (AlexaFluor 568 conjugate), which bound the streptavidin in the cSMAC (Figure 4). We chose to use streptavidin as the linker chemistry in the cSMAC because biotinylated T-cell stimuli (pMHCs and antibodies) are readily available. The dimensions of the SMAC mimics shown in Figure 4 are similar to those of T cells [23–25]. The protein patterns were stable for >2 months when slides were stored in PBS at 4°C.

4. Conclusions

The power of this technique is its simplicity of implementation and ability to create massive arrays in parallel of specifically-defined two-dimensional protein patterns without the need for blocking agents during conjugation steps, enabling cell patterning and the study of cellular interactions. Although we focused on the formation of surface SMAC mimics, any geometry is easily realized—much like integrated circuits—by designing and using a specific photomask set. Additionally, by combining this approach with that of microarray printing [13], arrays can be created in which each element contains a different functionality, enabling high-throughput screening.

Supporting Information Available. An alternative route to surface IS functionalization is given in addition to two figures, one a schematic describing the method and a second...
showing fluorescent optical micrographs of representative ISs patterned using the approach.

Acknowledgments

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References


