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Capillary flow in microchannel circuitry of scleral lenses

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Continuous monitoring of biomarkers in a quantitative manner at point-of-care settings can advance early diagnosis in medicine. Contact lenses offer a minimally-invasive platform to continuously detect biomarkers in tear fluid. Microfluidic components as lab-on-a-chip technology have the potential to transform contact lenses into fully-integrated multiplexed sensing devices. Here, simple and complex microchannels are created in scleral lenses that perform microfluidic operations *via* capillary action. The engraving of microchannels in scleral lenses were performed by laser micromilling, where a predictive computational model was developed to simulate the effect of laser power and exposure time on polymer behavior. Experimentally varying the CO₂ laser power (1.2–3.6 W) and speed (38–100 mm s^{−1}) allowed the micromilling of concave microchannels with groove depths of 10–240 μm and widths of 35–245 μm on polymetric substrates. The demonstrated laser micromilled circuitry in scleral lenses included linear channels, T/Y junctions, multiplexed arrays, mixers, and spiral channels, as well as serially organized multicomponent channels. Capillary forces acting in the microchannels allowed flowing rhodamine dye within the microfluidic components, which was visualized by optical microscopy in reflection and transmission modes simultaneously. The developed microfluidic components in scleral lenses may enable tear sampling, storage, analysis, and multiplexed detection capabilities for continuous monitoring applications.

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Introduction

Contact lenses have emerged as a minimally-invasive sensing platform for point-of-care diagnostics.¹ Contact lens sensors have been developed for analyzing the composition of tears as a surrogate medium for blood biomarker monitoring for potential application in the diagnosis of ocular or metabolic disorders. For minimally-invasive continuous monitoring applications, electrochemical sensors have been incorporated into the hydrogel film of contact lenses.² In the contact lens matrix, glucose oxidase (GOx) allowed for creating an electrical charge due to the formation of an enzyme–glucose complex.³ The magnitude of the charge created was proportional to the concentration of glucose oxidase. In another study, graphene with silver nanowires was used to create an electronic contact lens to measure glucose concentration.^{4,5} This contact lens featured resistance, inductance and capacitance circuits operating at radio frequency to enable real-time glucose detection. A sensing probe consisting of glucose oxidase was incorporated in

polyethylene as a substrate. The device detected glucose concentration of as low as 1 μM. Amperometric glucose sensors were also integrated in polyethylene terephthalate (PET) contact lenses by forming microstructures on a polymer substrate using photolithography.⁶ This enzyme-based electrochemical sensor was coated by a Nafion layer and allowed detecting glucose concentration as low as 0.1 mM.⁶ Hydrophilic poly(2-hydroxyethyl methacrylate) (p(HEMA)) contact lenses have been also utilized to create glucose oxidase based quantitative sensors.⁷

Optical components have been also embedded in a contact lens.⁸ A light-emitting diode (LED) pixel, a rectifier circuit and a stretchable antenna were integrated onto a substrate with high optical transparency and low haze. This substrate was then molded to fit the shape of a contact lens. Transparent nanomaterials were used to prevent obstruction of the wearer's view. The detection of high glucose concentrations turned the LED pixel off to enable qualitative readouts. In another study, an optical contact lens sensor was created by incorporating a boronic acid derivative in the hydrogel film of the contact lens.⁹ The contact lens had a photonic microstructure printed with a specific periodicity constant, which acted as a wavelength filter and diffracted specific wavelengths according to Bragg's law. Thereby, volumetric changes due to variation in glucose concentration (<50 mM) were detected by Bragg peak shifts

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within 20 min. In another study, holographic diffusing microstructures as an optical transducer were imprinted on glucose-responsive hydrogel contact lenses through lithography.¹⁰ In this contact lens, the scattering efficiency of the phenylboronic acid functionalized sensor was modulated by volumetric changes to determine glucose concentrations quantitatively.

Laser micromilling has emerged as a robust fabrication method for micropatterning plastics.¹¹ In laser micromilling, photoablation has been utilized for fabricating polymeric microfluidic channels.¹² Laser emission wavelength affects the machining efficiency in ablation depending on the absorption spectra of the substrates.¹³ Micromachining *via* laser ablation can be achieved without a mask in a direct-write process, where the substrate material is selectively patterned by a focused laser beam controlled by a scanning head. The micromilled regions are usually square or rectangular shaped with straight walls, although, depending on the depth of the channel, the walls can become slanted because of successive pass and laser defocusing. A wide variety of patterns have been fabricated *via* laser ablation offering low-cost and high speed as compared to conventional photolithography.¹⁴ We have proposed direct laser micromilling as a viable fabrication method for patterning linear channels and microconcavities in hydrogels.¹⁵ By utilizing direct laser micromilling, microfluidic components can be created in contact lenses to expand the capabilities in sampling, mixing, processing, and multiplexing.

Here, we create simple and complex microfluidic components in p(HEMA) scleral lenses *via* direct CO₂ laser micromilling. We have developed a computational model to predict the power and exposure time of the laser beam to form concave microchannels in scleral lenses. We experimentally demonstrate the fabrication of a wide range of microfluidic components including Y/T junctions, spiral channels, mixers, and multiplexed channels, as well as serially-organized multifunctional components in scleral lenses. Rhodamine B solution was utilized to optically visualize the capillary flow within the microfluidic channels of scleral lenses in reflection and transmission imaging modes.

Experimental section

Materials

p(HEMA) mini scleral lenses (water content 45 vol%; SPH: −0.00; diameter: 14.00 mm; base curve: 8.60 mm) were purchased from Meralens GmbH (Erding, Germany). Poly(methyl methacrylate) (PMMA) (2 mm thickness; 30 × 1000 mm cut) was purchased from B&T Metall- und Kunststoffhandel GmbH (Kirchheim bei München, Germany). Rhodamine B was purchased from Sigma Aldrich.

Equipment

A CO₂ laser (VLS 4.60, 10.64 μm, 60 W) was purchased from Universal Laser Systems (Vienna, Austria). Leica DM750 with Leica ICC50 HD camera was used to image the microfluidic channels. Leica Acquire (3.4.1) was used to capture and process the microscopic images. Multiple microscopic images were

merged and assembled to create wide-view microchannel images using Image J (1.52e, National Institutes of Health).

Laser micromilling of PMMA substrates and scleral lenses

Microfluidic patterns were designed as lines and surfaces in Adobe Illustrator CC (2018). A CO₂ laser cutter (10.64 μm, 60 W) was used to pattern microchannels on a PMMA sheet and p(HEMA) scleral lenses. The power of laser beam was varied from 1.2 W to 3.6 W at beam speeds ranging from 38 to 100 mm s^{−1}.

Microfluidic flow in scleral lenses

To visualize the fluid flow in the microchannels on scleral lenses, rhodamine B dye solution was prepared in deionized water (wt/vol, 1 : 10). A syringe with 23G needle was used to inoculate the rhodamine B solution (10 μl) at the inlet of the microchannels on the scleral lens. Capillary forces acting within the laser patterned microchannels wicked the rhodamine B solution and allowed the flow within a single channel leading to an outlet. The rhodamine B fluid flow was contained within the microchannels and no significant leakage was noted. The microfluidic channels were simultaneously imaged in reflection and transmission modes.

Results and discussion

CO₂ laser micromilling of polymers is a photothermal process. Excited CO₂ molecules in a resonant cavity amplify any electromagnetic wave associated with the bending energy transition of the CO₂ molecules, thus producing a photon. The photons then produce a laser beam until the gain medium is saturated and the beam is focused on the scleral lens (Fig. 1a). Laser ablation involves exposure of a highly-intense laser beam to the polymer surface leading to a shock wave, polymer vaporization, and ejecting particles from the substrate. This ablation process creates a microconcavity in the polymeric substrate (Fig. 1b). The beam spot size of the laser as well as the thermal capacity and density of the polymer affect the ablation efficiency. The polymer substrate (*e.g.* PMMA) transitions from a glassy state to a rubbery state above 115 °C.¹⁶ As the laser beam induced temperature increases from 170 °C to 210 °C, molecule chains begin deforming by releasing methyl methacrylate. At 360 °C, depolymerization initiated by the vinyl end groups, saturated C–C bonds, and –CH end groups.

The ablation process has been simulated in MATLAB to rationally design the geometrical characteristics of the concavity of the microchannel in the PMMA substrate. The concept behind the fabrication of the concavity consisted of applying thermal energy to increase the temperature at the surface of the polymer. When the temperature at the surface of the polymer increases to a value higher than the ablation threshold, the polymer is removed leaving a concavity at the surface, where the ablation temperature of PMMA was estimated as 700 K.¹⁷ In the simulation, this process was repeated iteratively while the beam moved through the polymer surface. The increment in temperature ΔT was found with the increment



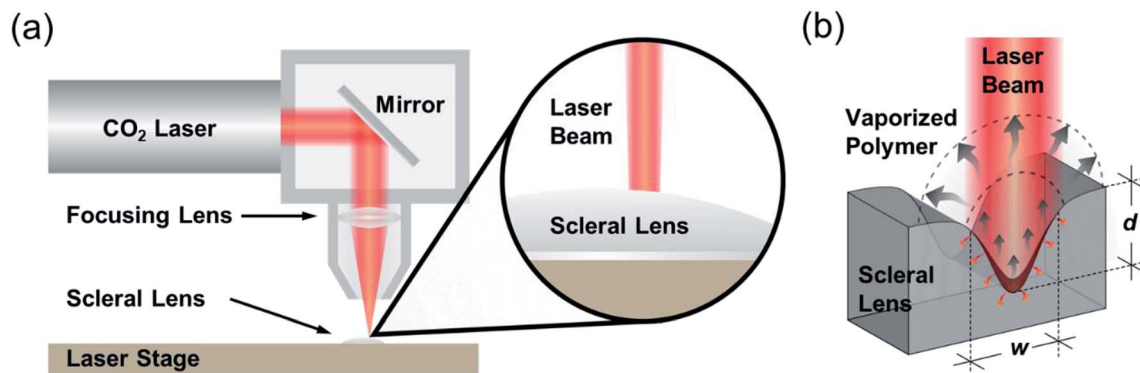


Fig. 1 The principle of laser micromilling of microchannels in polymers. (a) CO₂ laser setup for patterning scleral lenses. (b) Schematic of the interaction between the laser beam and polymer substrate. As the laser beam propagates toward the substrate, a shock wave is created and particles are ejected from the substrate by vaporization.

in thermal energy ΔU , the density ρ , and the volumetric heat capacity c_p of PMMA:

$$\Delta T = \Delta U \rho c_p \quad (1)$$

The thermal energy was retrieved during each iteration of the simulation through the absorption of the laser light. In this case, the conversion of optical energy to thermal energy was calculated with the loss produced in the propagation path of the laser beam. Simplifying the analysis for a perfectly collimated laser beam with an intensity normalized Gaussian profile:¹⁸

$$N(x, y) = \frac{2}{\pi w^2} e^{-\frac{2(x^2 + y^2)}{w^2}} \quad (2)$$

where w represents the radius of the beam profile, the radiant flux per unit of area I decreases exponentially along the propagation axis according to Beer–Lambert law:

$$I(x, y, z) = I_0 N(x, y) e^{-z\alpha} \quad (3)$$

where total I_0 is the intensity of the laser beam and α the absorption coefficient.¹⁹ When all the optical power is completely converted as thermal energy, then the difference of the optical power between two points of the propagation path is equivalent to the thermal energy dissipated. Therefore, in infinitesimal terms, the heating source is equivalent to the derivative of Beer–Lambert law equation. In this manner, the heating rate per unit of volume q for a collimated Gaussian beam was obtained as:

$$q(x, y, z) = I_0 N(x, y) \alpha e^{-z\alpha} \quad (4)$$

As in the case of the optical power, the heating rate per unit of volume was integrated to obtain the total energy introduced to the system. The total increment of energy was obtained by adding heating rate per unit of volume in each iteration while the beam was moving. Furthermore, the dissipation of the heat produced by the source can also be retrieved with the numerical solution of the diffusion equation:

$$q = \rho c_p \frac{\partial U}{\partial t} - k \nabla^2 U \quad (5)$$

where k is the thermal diffusivity of the polymer. In this work, the heat diffusion has been calculated with the forward Euler approach.²⁰ This is a first order method in which each point of the grid transfers the heat to adjacent points in an iterative manner. The model was defined based on the chemical and optical properties of PMMA, and 10.64 μm was used as the emission wavelength of a CO₂ laser beam. Fig. 2a illustrates the simulated 2D top view temperature distribution as a function of laser beam power. The temperature increase up to 400 °C resulted in the vaporization and subsequent ejection of material from the polymeric substrate. No ablation was present when the laser power was 1.2 W at a constant beam speed of 75 mm s^{-1} . However, changing beam power from 1.2 W to 3.6 W while holding the beam speed constant increased the ablation depth to 111 μm and width to 255 μm (Fig. 2b). Fig. 2c illustrates the simulated 2D cross-section temperature distribution and profiles of ablated concavities as a function of beam power. The resulting ablated pattern was concave due to the Gaussian distribution of the laser beam power.

In experimental studies, a PMMA sheet was engraved with a CO₂ laser to form concave microchannels. The beam power was varied from 1.2 W to 3.6 W when the beam speed was set constant (75 mm s^{-1}). Fig. 2d and e shows the optical microscopic images of the top views and cross-section of the laser ablated channels as the laser power increased. As the laser power increased, the ablation depth and width also increased. Increasing the power in range of 1.2 W to 1.8 W produced subtle microconcavities. However, increasing the beam power above 2.4 W produced deeper microconcavities while maintaining its concave geometry. Fig. 2f depicts the computational and experimental results of the ablation depths and widths as a function of the applied beam power. At a constant beam speed of 75 mm s^{-1} , 3.6 W of beam power produced an ablation depth of 90 μm while the ablation width reached 225 μm . The computational and experimental results showed that both curves follow a similar pattern, where the ablation width had larger margins of increase as the power increased. The experimental studies agreed with the simulated model, showing that the concavity depth could be accurately predicted based on the beam power variation.



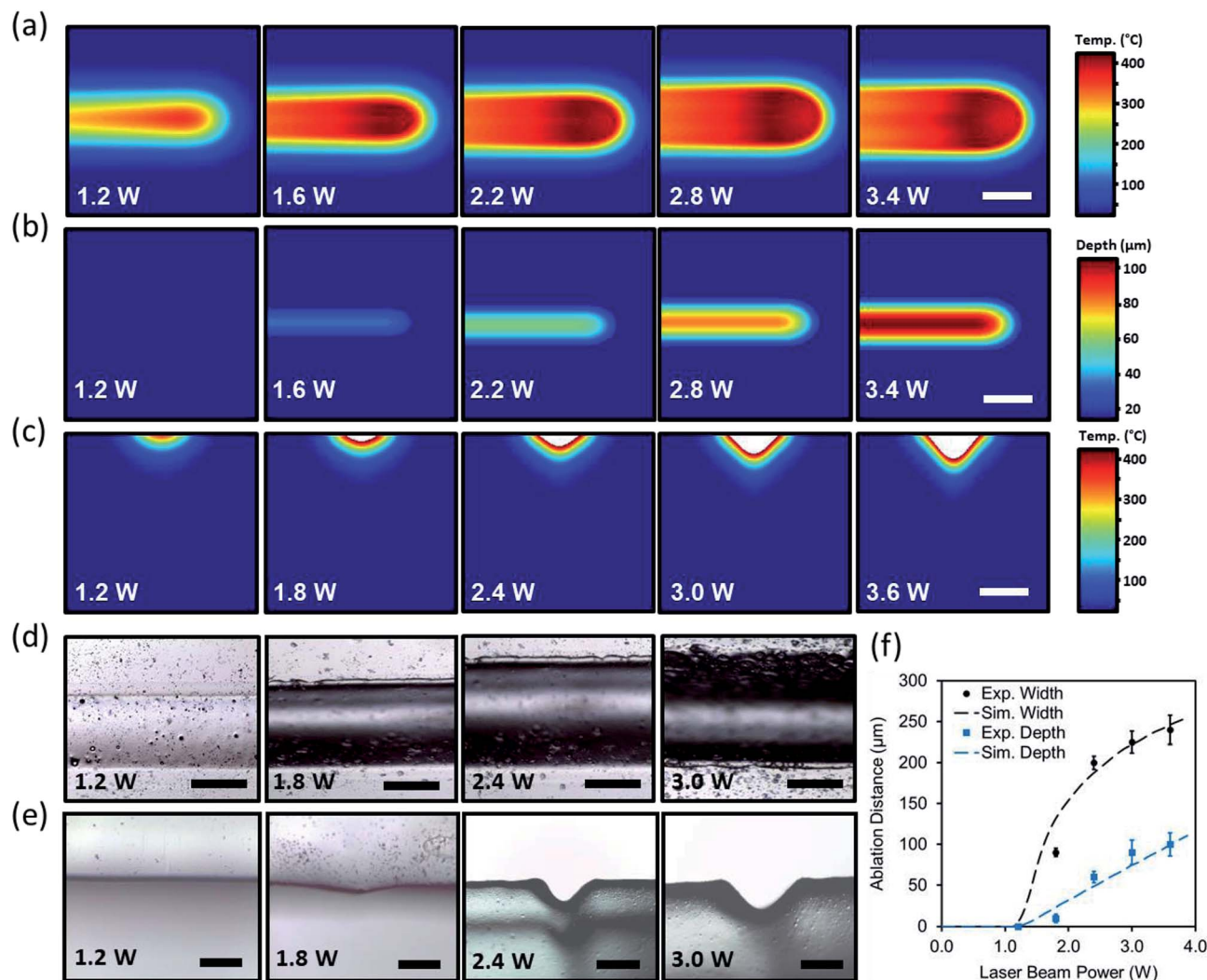


Fig. 2 The effect of laser beam power on the ablation depth and width at a constant beam speed of 75 mm s^{-1} . Simulated top view 2D (a) temperature and (b) ablation depth profile variations with beam power changes. Scale bar = $200 \text{ } \mu\text{m}$. (c) Simulated cross-section 2D temperature and ablated concavity profile variations with beam power changes. Scale bar = $200 \text{ } \mu\text{m}$. (d) Experimental top view and (e) cross-section images of the engraved microconcavities on a PMMA sheet as a function of power variation. Scale bars = $100 \text{ } \mu\text{m}$. (f) Ablation depth and width as the beam power was varied. Error bars represent the standard error ($n = 3$).

To understand the effect of beam exposure time on the laser micromilling process, the beam speed was computationally varied from 38 mm s^{-1} to 100 mm s^{-1} while the beam power was held constant at 3.6 W . Fig. 3a illustrates the simulated 2D top view temperature distribution as a function of laser beam speed. Increasing beam speed from 38 mm s^{-1} to 100 mm s^{-1} while having a constant beam power of 3.6 W decreased the ablation depth from $212 \text{ } \mu\text{m}$ to $26 \text{ } \mu\text{m}$ and the width from $274 \text{ } \mu\text{m}$ to $134 \text{ } \mu\text{m}$ (Fig. 3b). Fig. 3c illustrates the computational temperature distribution and 2D profiles of the ablated concavities as a function of laser beam exposure time.

In experimental studies, the influence of the beam speed on the ablation depth and width of microconcavities was investigated. Fig. 3d displays the microscopic cross-section view images of the laser ablated microconcavities as the beam speed increased from 38 mm s^{-1} to 100 mm s^{-1} as the beam power

was set to 3.6 W . Fig. 3d displays top view images of the cross-section cuts of the channels with varying beam speed. Using higher speed values (100 mm s^{-1}) produced a subtle microconcavity (Fig. 3e). The ablation depth and width were $120 \text{ } \mu\text{m}$ and $244 \text{ } \mu\text{m}$ at 38 mm s^{-1} beam speed and decreased to $35 \text{ } \mu\text{m}$ and $125 \text{ } \mu\text{m}$ at 100 mm s^{-1} beam speed respectively (Fig. 3f). The experimental studies agreed with the simulated model, showing that the concavity depth could be accurately predicted based on the beam speed variation.

Fabrication of microchannels on scleral lenses

Microchannels were engraved on p(HEMA) scleral lenses to form microfluidic components that may enable tear sample collection and multiplexed analyses in continuous monitoring applications. Fig. 4a–f shows the basic and complex microchannel patterns on p(HEMA) scleral lenses, and the insets



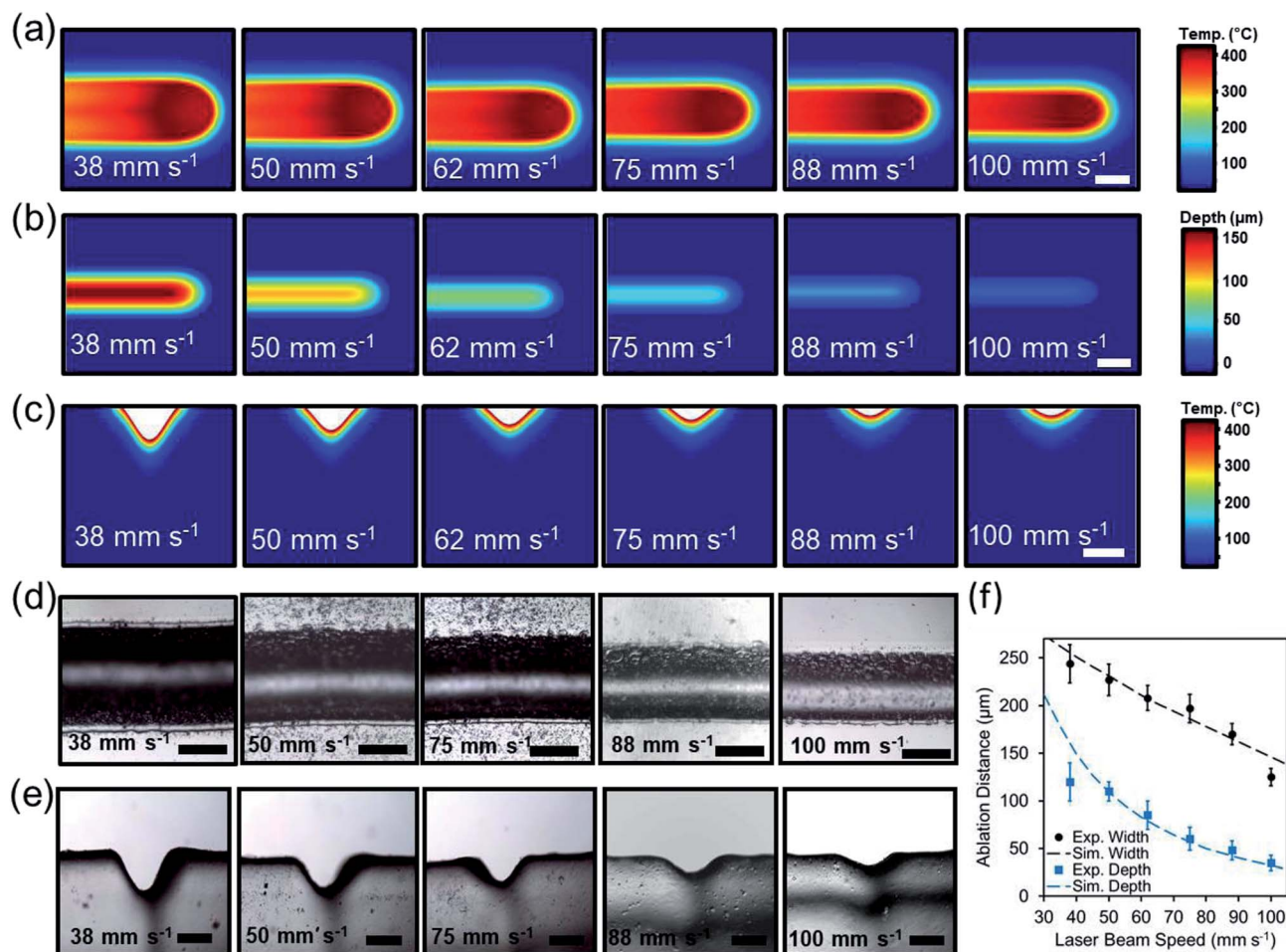


Fig. 3 The effect of laser beam speed on the ablation depth and width at a constant beam power of 3.6 W. Simulated 2D top view (a) temperature and (b) ablation depth profile variations with beam speed changes. Scale bar = 200 μm. (c) The simulated CO₂ ablation of a PMMA sheet. Temperature and 2D ablated concavity profile variations with beam speed. Scale bars = 200 μm. (d) Cross-section and (e) top view images of engraved microconcavities on a PMMA sheet with beam speed variation. Scale bars = 100 μm. (f) Ablation depth and width as a function of beam speed variation. Error bars represent the standard error ($n = 3$).

display optical microscopic top view images of the microchannels. Fig. 4a illustrates a basic line microchannel with one inlet and one outlet on the scleral lens. Fig. 4b shows a T-junction microchannel as a combination of two perpendicular line components. The T-junction microchannel can be utilized as 2 inlet 1 outlet component or as 1 inlet 2 outlet schemes. A Y-junction microchannel was also created as a basic component on a scleral lens for multiplexing the fluid from one branch to the other (Fig. 4c). Combining Y components together could achieve complex tree like microfluidic systems for highly multiplexed operations. Fig. 4d depicts a four-plexed pattern on the scleral lens, where the inlet branches out to four outlets. Microchannels featuring curves can also be used to achieve mixing functionality as the micropattern curves toward the inside and then outside in a cyclic motion that allows the mixing the fluid in the microchannel (Fig. 4e). A spiral microchannel could achieve separation of fluids or microscale solids with different densities (Fig. 4f). The inlet shown in outer circle swirls into smaller circles and ultimately to into the outlet.

Fluids with different densities could be separated as the circles get smaller to pass the least dense fluid or filter out a microscale solid. For example, microcontaminants could be extracted from the tear fluid using the spiral microchannel.

To provide complex microfluidic functions, wider available area over the scleral lenses was utilized. Fig. 4g shows a mixer system with a number of crown-patterned channels running around the whole scleral lens surface. Fig. 4h illustrates a channel with one the inlet and outlet expanding over the scleral lens surface. This system could be used to separate microparticles from the fluid as some solids will be filtered out in the microchannel. Fig. 4i shows a microchannel system with two inlets that could mix different fluids at the overlapping region of the microchannels leading to an outlet. Fig. 4j shows a multiplexer system with four outlets over the scleral lens. Fig. 4k shows three reaction chambers connected with one inlet and outlet, where tear fluid can be stored or reactions can occur on the scleral lens. Fig. 4l shows a microchannel system having one inlet connected to a mixer component which leads to



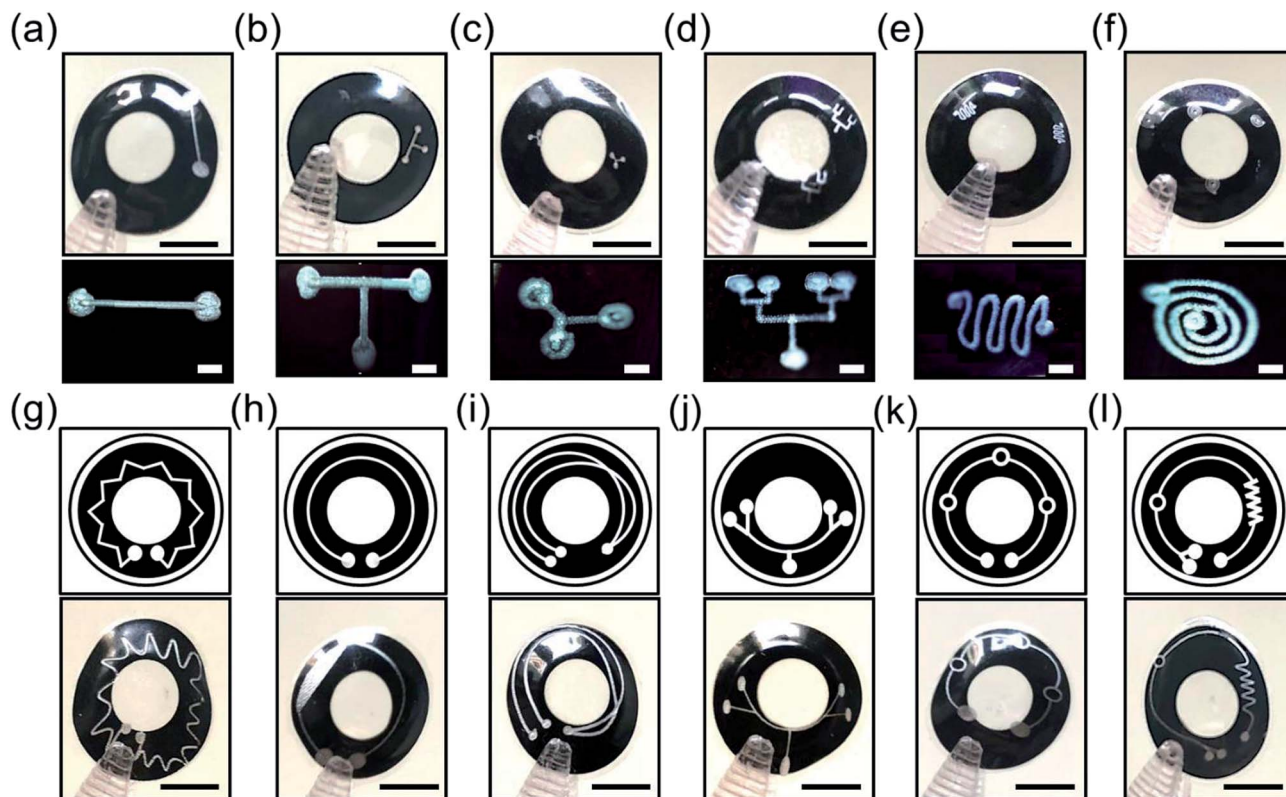


Fig. 4 Microchannels engraved on p(HEMA) scleral lenses. The insets show microscopic top view images of the engraved micropatterns. (a) Basic line component with 1 inlet 1 outlet. (b) T-junction component with 1 inlet and 2 outlets. (c) Y-junction component with 1 inlet and 2 outlets. (d) Multiplexed array with 1 inlet and 4 outlets. (e) Mixer component with 1 inlet and 1 outlet. (f) Spiral component with decreasing radius having 1 inlet and 1 outlet. (g) Mixer microchannel with a crown structure. (h) A microchannel system with 1 inlet and 1 outlet, where the channel widens around the scleral lens. (i) A system having intersected microchannels with 1 inlet and 2 outlets. (j) Multiplexer system with 1 inlet and 4 outlets. (k) A microchannel system with three reaction chambers, in which fluid could be stored or reactions could occur. (l) A system comprising of a mixer component connected to a reaction chamber that branches out to two outlets. (a–f) Scale bars = 5 mm and inset scale bars = 200 μm . (g–l) Scale bars = 5 mm.

a reaction chamber subsequently directed to 2 outlets. Such microchannels systems can carry out multistep reactions, processes, and multiplexed analyses on the scleral lens. Based on the Rayleigh length for a CO_2 laser, the Gaussian distribution of beam width was consistent within ± 2.0 mm from the focal point. The base curve radius of the contact lens used in this work was less than 2.0 mm; hence, the widths of the channels formed along the surface of the contact lenses were precise.

Fluid flow in microfluidic scleral lens components

To visualize the fluid flow in the microchannels on scleral lenses, rhodamine B dye solution in deionized water was utilized (wt/vol, 1 : 10). A syringe with 23G needle was used to introduce the rhodamine B solution (10 μL) at the inlet of the microchannels on the scleral lens. Fig. 5 illustrates rhodamine B solution (in pink) flow in basic line component, T/Y-junction components, multiplexed microchannel, mixer component, and spiral microchannel, and the insets show the magnified images of the fluid flow. The liquid flow was attained along the microchannels due to the capillary action of the water on the scleral lens. Fig. 5a shows the flow of the rhodamine dye into

a microconcavity. Such microconcavities could be used as reaction chambers in scleral lenses. Fig. 5b shows the dye flow into a T junction from two inlets merging to a single inlet; such devices can be used as microfluidic timers, in which different dye flow can be sequentially delivered to a reaction chamber. Fig. 5c shows a Y component that consists of three reaction chambers that may be utilized as a multiplexed assay. Fig. 5d illustrates a multiplexed array that distributes the dye into four reaction chambers for multiplexed analysis. Fig. 5e shows the flow of the rhodamine B dye in a microfluidic mixer system consisting of one inlet and outlet. Fig. 5f displays rhodamine dye flow in a spiral microfluidic component. Hence, complex fluid flows can be attained by capillary flow in scleral lens microfluidics. The flow of the fluids in the channels can be controlled by modifying the wettability characteristics of the channels. For example, hierarchical microcone arrays and pillars can be formed using water-assisted femtosecond laser irradiation^{21,22}

A continuous wave CO_2 laser was employed to engrave microchannels on scleral lenses. With the development of high-energy lasers with rapid scan head platforms, precise micro-milling operations can be carried out using ultrashort light



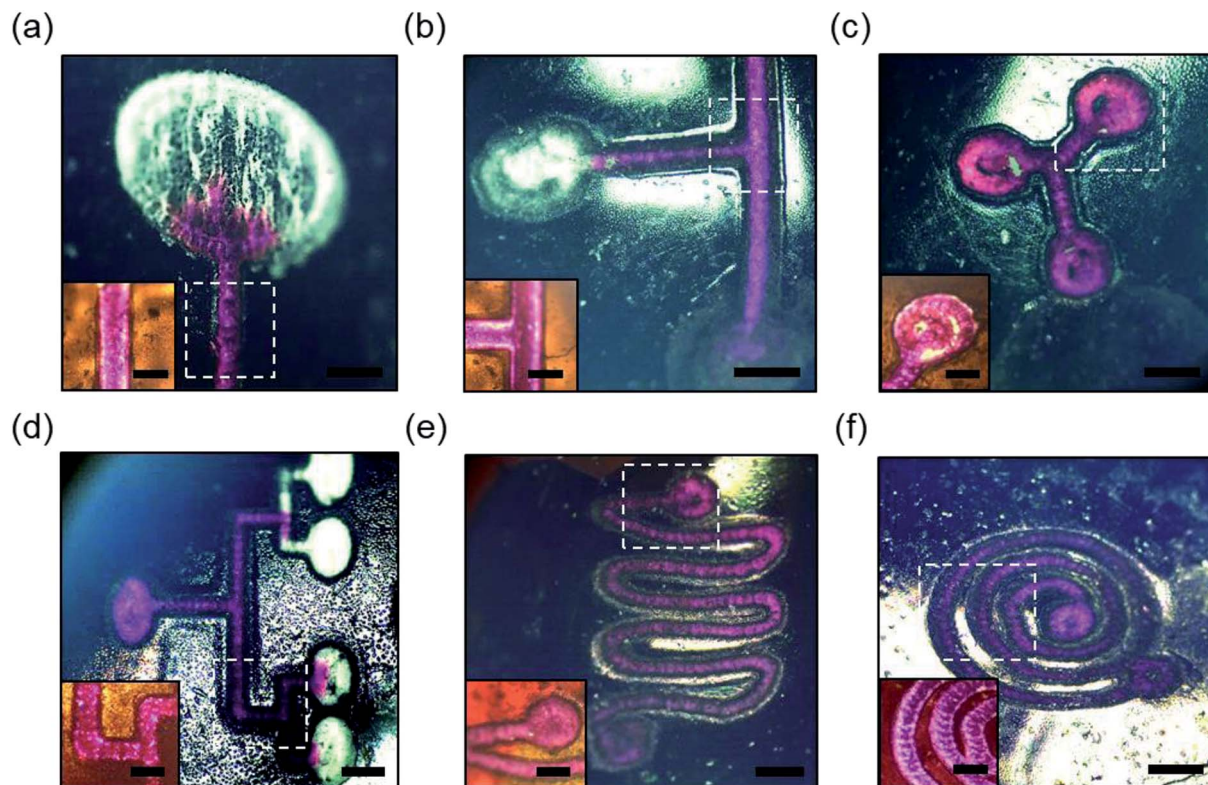


Fig. 5 Rhodamine B solution flow in microfluidic channels on scleral lenses. The insets show magnified images of the fluid flow. (a) Basic line component with 1 inlet 1 outlet. (b) T component with 1 inlet and 2 outlets. (c) Y component with 1 inlet and 2 outlets. (d) Multiplexed array with 1 inlet and 4 outlets. (e) Mixer component with 1 inlet and 1 outlet. (f) Spiral microchannel with decreasing radius having 1 inlet and 1 outlet. Scale bars = 200 μm and inset scale bars = 100 μm .

pulses. Femtosecond laser milling has emerged as a precise approach for laser ablation in microfluidic device fabrication.²³ An advantage of femtosecond lasers is the absence of thermal effects, where structural polymer characteristics do not significantly change as there is subtle thermal tension, shock waves and melting effects.²⁴ These ultrashort laser pulses have enough high peak power to drive nonlinear multiphoton absorption, leading to laser ablation at wavelengths where the polymer has low absorption. Ultrashort pulse laser ablation is preferred to that of longer pulse sources, where the photon energy must be greater than the bandgap to deposit energy *via* linear absorption.¹⁴ Laser ablation can also be used to generate 3D microfluidic channels inside bulk PMMA substrates.²⁵ This approach uses femtosecond pulses to produce nonlinear absorption inside the focal volume, triggering a polymer modification. Such laser fabrication approaches could be adopted to construct 3D channels in scleral lenses. Furthermore, in the present work, contact lenses made of pHEMA were demonstrated; however, other contact lens materials including silicone acrylates and other co-polymers can also be used to form channels with the laser ablation method. The intensity and the speed of laser beam can be controlled to pattern any polymer matrix depending on thermal conductivity, water content, absorbance, and density. Creating a microfluidic system is vital for continuous tear fluid sampling and multiplexed analysis at point-of-care settings. Furthermore, autonomous pumps

consisting of micropillars could be fabricated on the microfluidic channels to rapidly sample tear fluid.^{26,27}

Conclusions

Microfluidic components were developed in scleral lenses, where laser ablation has been chosen to machine microchannels on scleral lenses. A numerical model was developed to predict the depth and width of microscale channels in scleral lenses. Additionally, the physical heat phenomena behind laser ablation has been described. Beam power and speed was optimized to create microchannels in polymers. Upon optimizing the beam speed and power, basic and complex microstructures have been engraved on scleral lenses, including basic line components with inlets and outlets, multiplexed channels, Y/T junctions, spiral components, and microfluidic mixers. Rhodamine B solution was introduced to the microchannels where capillary flow guided the flow in the channels. Fabricated microfluidic channels will serve a platform role in the fabrication of lab-on-a-chip functionalities in scleral lenses for sample collection, storage, and analyses in real-time biomarker monitoring in tear fluid.

Conflicts of interest

The authors declare no competing financial interests.



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