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SIMULTANEOUS IMPROVEMENT OF SURFACE FINISH AND BONDING OF CENTRIFUGAL MICROFLUIDIC DEVICES IN CYCLO-OLEFIN POLYMERS

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ABSTRACT

Two key issues in the manufacturing of microfluidic devices are to obtain low surface roughness, i.e. in the range of nanometers (to facilitate optical detection and controlled flow through microfluidic networks), and to achieve robust bonding. Here we report that a chemical polishing step, used for smoothening the surface of cyclo-olefin polymer (COP) components manufactured by micromilling, reduces the surface roughness (to $R_a \sim 150$ nm) and facilitates a leak-tight COP-COP bond. We report new results on COP structuring and surface characterization by white light interferometry (WLI), infrared spectroscopy (FTIR-ATR), contact angle measurements and transmittance measurements, as well as demonstrations of a functional thermally bonded centrifugal microfluidic devices.

INTRODUCTION

Optical detection¹ is often required in microfluidic systems. Ideally, the sensitivity of the detection method should be solely limited by the biochemical recognition process, and not limited by the surface finish (light scattering at rough surfaces and alteration of the optical path) and the bonding interface.

Microfluidic devices in plastics with optical surface quality are typically manufactured by e.g. injection moulding^{2,3} using a polished mould. These technologies are well established. However, tooling and mould inserts required for such processes are expensive and many labs do not have access to an injection moulding machine. Surface finish requirements can also be an important cost-driving factor in the device development stage.

Sealing of microfluidic devices can be achieved using a variety of direct and indirect bonding techniques⁴. For example, adhesive bonding is one of the straightforward alternatives. However, when channel dimensions are narrower than about 800 μm , reproducible cutting and alignment of fluidic structures in adhesives becomes challenging.

While there are some studies of chemical polishing and vapor-assisted thermal bonding of surfaces of cyclo-olefin polymers (COP) and cyclic olefin copolymers (COC)⁵⁻⁷, most contributions address processes for other transparent polymeric substrates such as polymethyl-methacrylate and polycarbonate⁸.

Here, we provide the details of a process that can be used both to obtain acceptable optical surface quality (low roughness and acceptable transparency) and to assist in achieving a leak-tight COP-COP bonding. This study focuses on COP and the surface finish that can be achieved by micromilling⁹, and by micromilling in

combination with chemical polishing (i.e. controlled exposure to cyclohexane vapor⁴), and subsequent solvent-assisted thermal bonding^{10,11}.

MATERIALS AND METHODS

Disc design

A design with 5-7 equally spaced test structures on a $\text{Ø}120$ mm disc was used for developing the bonding process. Each structure contained both wide chambers and fluidic channels machined to several depths. Dimensions are provided in Table 1.

The most critical area for bonding was determined to be chamber B, therefore several different discs with depths varying between 100 and 500 μm were prepared.

Table 1: Dimensions of fluidic test structures.

| Structure | LxWxH (mm) | Position on disc* |
|------------------|----------------------|-------------------|
| Chamber A | 15 x 12 x 0.6 | R23 |
| Chamber B | 24 x 20 x (0.1–0.5) | R35 |
| Chamber C | 50 x 10 x 0.9 | R50 |
| Fluidic channels | 30 x 0.5 x (0.1–0.2) | R20-R50 |

*Radius values refer to the center of the disc.

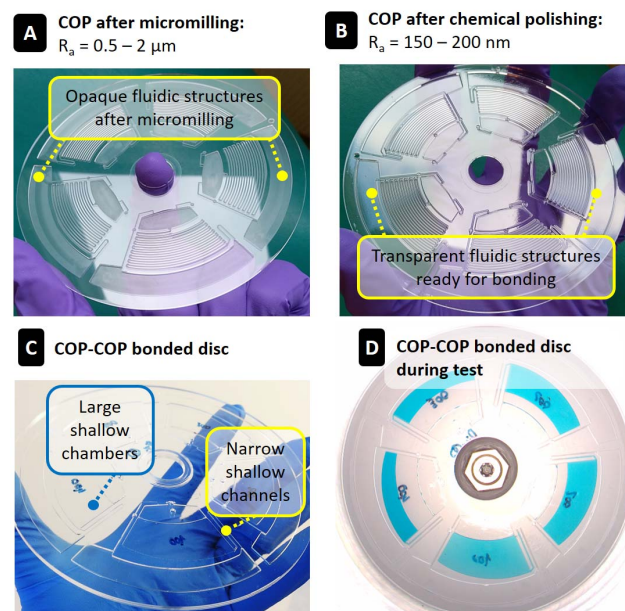


Figure 1: Images of COP discs showing opaque chambers after milling (A), transparency after chemical polishing (B), thermally bonded COP-COP disc (C), and bonded disc during testing at a spin rate of 4000 rpm and sample volume of 100 μL in chambers of various depths (D).

Machining of fluidic structures

Fluidic structures were directly milled in injection-molded blank COP discs (1.2 mm thick $\text{\O}120$ mm discs; COP grade: Zeonor 1060R from Zeon) using a 3-axis milling machine (DMG DMC 1035 V). The micromilling was optimized with respect to type of milling tool (single tooth end mill 0.5 and 1 mm diameter), spindle speed (37 000 rpm), feed rate and cut width, in order to achieve low surface roughness (with R_a in the range of 300-500 nm for optimized processes, but can be as high as 2 μm). The disc fixture during milling ensured minimal warpage of the disk and mechanical stability, specifically for machining of 0.9 mm deep structures.

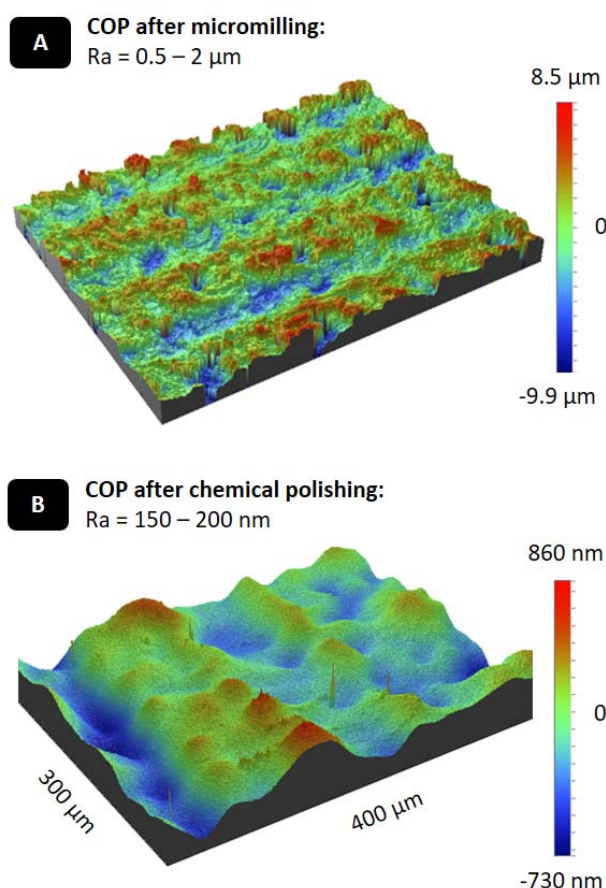


Figure 2: Roughness measurements after milling (A) and after milling followed by chemical polishing (B).

Surface polishing and bonding

A single exposure to cyclohexane vapor was used for both reducing the surface roughness (after micromilling) and as a surface treatment prior to thermal bonding (illustrated in Figure 1). The discs were sonicated in DI water for 10 min and dried with compressed air. Next, the discs were placed inside a petri dish in a Teflon holder with the bonding side facing a cyclohexane bath (disc surface ca. 4.5 mm above the level of cyclohexane) and kept for 4 to 6 min. Immediately after, the discs were aligned using a jig, pre-bonded manually and placed inside an automated hot press between two supporting glass plates.

This was followed by thermal bonding at 65°C for 30 min, followed by 15 min at 25°C before removing from

the press. The bonding force applied was varied between 1 and 15 kN in the bonding trials. Upon completion of the bonding process the disc was removed from the press and allowed to cool down. At least four discs were tested for each of the bonding conditions.

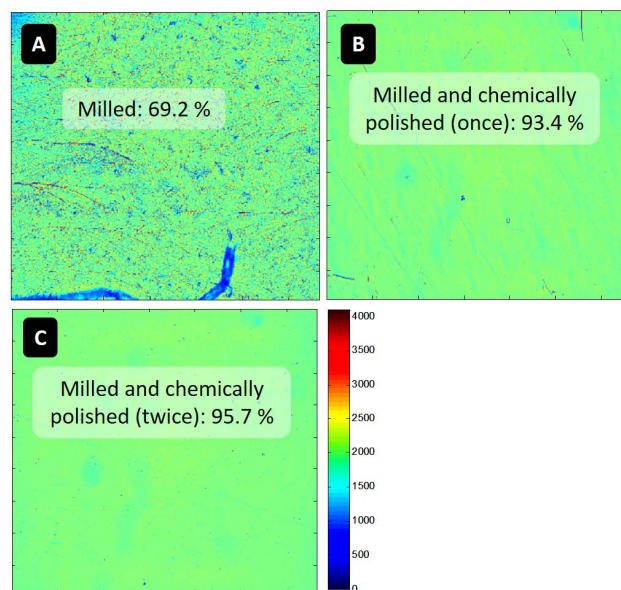


Figure 3: Comparison of microscope images of COP surfaces after milling (A) and after chemical exposure (B, C) with average transmittance values collected in the range 400 to 650 nm indicated. The transmittance value after chemical polishing is close to that of a part injection molded with a polished mold.

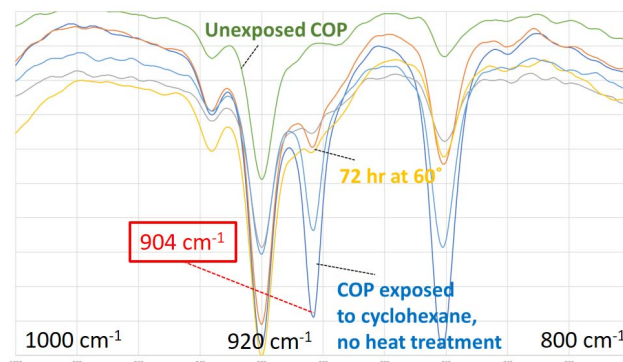


Figure 4: FTIR analysis of COP surface after milling, polishing and heat treatment to eliminate cyclohexane (corresponds to 904 cm^{-1}) from the COP surfaces: peak reduction can be observed for cyclohexane treated surface after heat treatment; no cyclohexane peak is found on unexposed COP surface.

Characterization

Surface roughness and dimensional integrity of fluidic structures were assessed before and after the chemical polishing step using WLI (Veeco® Wyko NT9800) and optical microscopy (Olympus® DSX100). FTIR-ATR measurements (PerkinElmer® Spectrum One with diamond crystal) were performed with 4 scans and 3 areas per sample. Water contact angles (Krüss® DSA 100 drop shape analyser) of the COP surfaces were

measured with drop volumes of 3 μL and 10 μL .

Optical transmittance was characterized using a custom-made setup. A plane monochromatic wave (bandwidth of ~ 4 nm) was focused onto the COP surface at normal incident. The transmitted light was collected by a 2x microscope lens and focused onto a CCD camera.

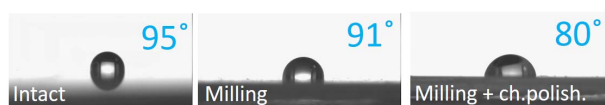


Figure 5: Contact angle measurements (3 μL drop volume) of COP surfaces before milling ("intact"), after milling, and after milling and chemical polishing.

The shear stress required to debond COP substrates was measured using a Dage 400Plus multipurpose bond tester. Test samples (5 x 5 mm²) were milled out of the COP-COP stack bonded at optimal conditions from three positions on the disc (R18, R30, R55). The force was applied to the centre of the top COP substrate in the bonded sample.

The fluidic tests were carried out using an automated custom-made centrifugal test stand equipped with a stroboscopic light source, a camera for visualization of liquid flow and a motor. Rotational speed (< 5000 rpm) and time (< 30 min) were controlled during leak tests.

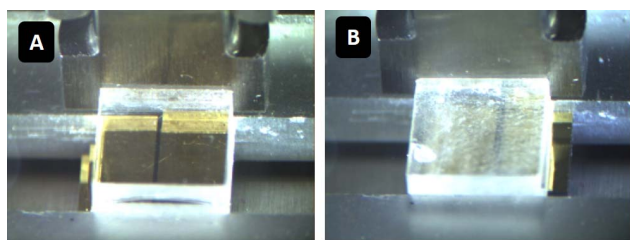


Figure 6: Shear test: COP-COP bonded sample before (A) and after (B) delamination (shear stress of 15.7 MPa).

RESULTS AND DISCUSSION

Reduction of surface roughness

It was demonstrated that a 6 min chemical polishing step is sufficient to reduce the surface roughness from 500 nm (R_a after micromilling is in the range of 300-500 nm for optimized milling processes, but can be as high as 2 μm) down to 100–150 nm, see Figure 2. Multiple exposures to cyclohexane yield further reduction in surface roughness by a few percent. However, maximum two subsequent exposures are recommended in order to avoid distortion of channel and chamber dimensions.

Surface transparency

It was shown that chemical polishing gave transmittance values in the range 92 – 95 % as measured with our setup. These values were comparable to the reference parts (prepared by injection molding with a polished mold). Transmittance data with corresponding images are summarized in Figure 3. Comparison of optical transmission of COC surfaces treated with various solvents, including cyclohexane, can be found elsewhere⁶.

Physical and chemical surface properties

Residues of cyclohexane were found on the COP surfaces even after 72 hours of drying at 60°C (Figure 4). However, this residue is expected to be negligible for typical surface-based chemistries. The wettability of the COP surface is slightly increased by milling and polishing (Figure 5), based on the static water contact angle measurements. This might eliminate the need for hydrophilic coatings¹² and surface treatment for some fluidic applications.

Bonding

The same polishing step was used to prepare COP discs for thermal bonding. Bonding conditions were optimized for both large and shallow detection chambers, and shallow micro channels, resulting in a transparent interface. It was found that 65 °C and 10 kN are optimum conditions for achieving a tight and transparent bond without the largest chambers collapsing.

Bond strength measurements on test structures (Figure 6) indicated that delamination occurred after applying a shear stress of about 6.8 MPa (mean value). The highest shear stress value measured was 15.7 MPa. The bond strength was found to vary depending on the area of the disc, with lowest value of 0.6 MPa measured for the samples taken from the outer edge of the disc. However, even the lowest value is still sufficient for leak-tight device operation and comparable to some previously reported data for COP-COP bonded microfluidic chips^{4,13}.

Successful testing of a centrifugal device with surface quality suitable for optical detection (e.g. in transmission mode) was demonstrated, see Figure 7.

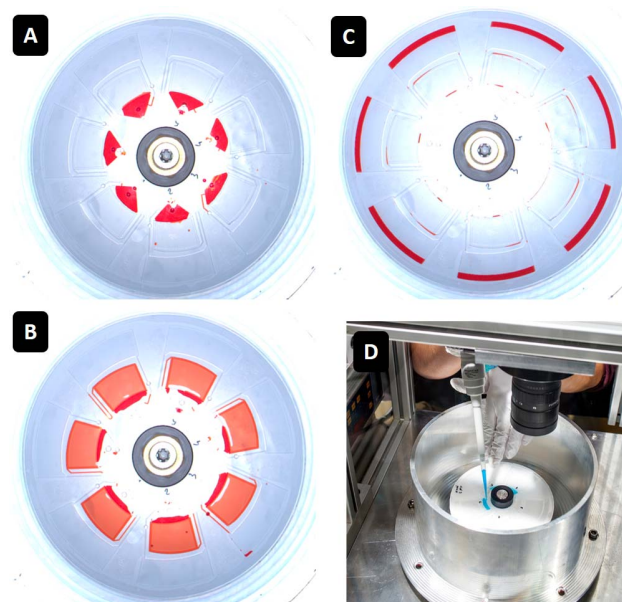


Figure 7: Example of a centrifugal device manufactured using the discussed prototyping method: loading of reagents – chamber A (A), transfer of reagents for incubation – chamber B (B), transfer of reagents to waste via a siphon valve – chamber C (C), disc loading (D).

CONCLUSIONS

We have demonstrated manufacturing, characterization and testing of COP-based centrifugal microfluidic devices. For COP-based microfluidic prototypes, micromilling in combination with chemical polishing is an attractive alternative to injection molding and hot embossing (expensive inserts and masters are required), and to 3D printing (3D printed COP parts are not available), when microfluidic structures with low surface roughness and high transparency are required.

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