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# Plug-and-play polymer microfluidic chips for hydrated, room temperature, fixed-target serial crystallography†

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The practice of serial X-ray crystallography (SX) depends on efficient, continuous delivery of hydrated protein crystals while minimizing background scattering. Of the two major types of sample delivery devices, fixed-target devices offer several advantages over widely adopted jet injectors, including: lower sample consumption, clog-free delivery, and the ability to control on-chip crystal density to improve hit rates. Here we present our development of versatile, inexpensive, and robust polymer microfluidic chips for routine and reliable room temperature serial measurements at both synchrotrons and X-ray free electron lasers (XFELs). Our design includes highly X-ray-transparent enclosing thin film layers tuned to minimize scatter background, adaptable sample flow layers tuned to match crystal size, and a large sample area compatible with both raster scanning and rotation based serial data collection. The optically transparent chips can be used both for *in situ* protein crystallization (to eliminate crystal handling) or crystal slurry loading, with prepared samples stable for weeks in a humidified environment and for several hours in ambient conditions. Serial oscillation crystallography, using a multi-crystal rotational data collection approach, at a microfocus synchrotron beamline (SSRL, beamline 12-1) was used to benchmark the performance of the chips. High-resolution structures (1.3–2.7 Å) were collected from five different proteins – hen egg white lysozyme, thaumatin, bovine liver catalase, concanavalin-A (type VI), and SARS-CoV-2 nonstructural protein NSP5. Overall, our modular fabrication approach enables precise control over the cross-section of materials in the X-ray beam path and facilitates chip adaption to different sample and beamline requirements for user-friendly, straightforward diffraction measurements at room temperature.

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## Introduction

As a result of continued development in the field of X-ray crystallography, the number of X-ray structures deposited annually in the Protein Data Bank ([rcsb.org](https://rcsb.org)) has continued to increase rapidly from 135 in 1990 to 11 234 in 2020. Alongside

the rise of cryo-crystallography,<sup>1</sup> these include improvements in crystallization methods,<sup>2</sup> approaches to high-throughput screening,<sup>3</sup> and the emergence of highly brilliant, microfocus synchrotron beamlines enabling the collection of data from ever smaller crystals.<sup>4,5</sup> Meanwhile, the development of femtosecond XFELs has ushered in a new era of structural biology, with radiation damage-free data collection at room temperature (RT) using serial femtosecond crystallography (SFX) due to the diffraction before destruction principle.<sup>6–8</sup> While single crystal cryogenic measurements continue to be the workhorse of macromolecular crystallography, technological development to support XFEL experiments has elicited a resurgence of interest in RT data collection at synchrotron facilities in recent years, with the parallel appearance of serial synchrotron crystallography (SSX) and serial oscillation crystallography (SOX) methods. In contrast with cryogenic measurements, RT synchrotron studies are advantageous because they eliminate the need to screen cryoprotectants and freeze samples.<sup>9</sup> Importantly, measurements at RT have

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opened avenues to study not just static structures but also protein dynamics over a broad time range, from picosecond to seconds, using time-resolved measurements. This new frontier in capability is paving the path for making molecular movies involving ligand binding, photoactivation, and catalysis to better understand dynamic structure–function relationships but requires facile and functional sample delivery.<sup>10–14</sup>

The high intensity of 3rd and 4th generation microfocus X-ray beamlines, fast onset of radiation damage at RT, and the necessity at times to use microcrystals limits the number of high-resolution diffraction frames that can be collected from a single crystal. To address this limitation, serial crystallography (SX) can be used, where data from multiple crystals is combined.<sup>6,15,16</sup> For SOX, in which small rotation wedges (1–20°) are collected from small crystals (~tens of  $\mu\text{m}$  in size), data from tens to hundreds of crystals is typically sufficient.<sup>17</sup> For SSX or SFX, each diffraction volume is exposed only once and diffraction data from thousands of crystals is often required to obtain a complete dataset with high redundancy and good signal-to-noise at high resolution.<sup>15,18</sup> In most cases, these techniques require a much larger amount of crystalline material than conventional crystallography, using crystals that may be too small to successfully mount using conventional cryo-loops. This necessitates the development of specialized crystal delivery methods which are continuous, robust, and keep the crystals hydrated over the course of the measurement. Therefore, an optimal sample delivery platform should: (1) maintain crystal quality and hydration by minimizing sample stresses during preparation, characterization, and delivery; (2) minimize scatter contribution from the delivery device and excess buffer surrounding the crystals; (3) optimize crystal density and crystal size to the beam and data collection requirements; (4) allow efficient use of beamtime by minimizing down time (*e.g.* from clogging or sample alignment for rastering) with fast/automated sample changes (*e.g.* sample preparation in advance); and (5) allow experiments to probe structural dynamics by allowing different triggering methods.<sup>19,20</sup> Various sample delivery approaches have been proposed and demonstrated including liquid-jets,<sup>21–24</sup> droplet-on-demand tape drives,<sup>25,26</sup> and fixed-target devices.<sup>27–32</sup> The work presented here focuses on a new fixed-target platform for SOX or SSX/SFX. Fixed-target devices are advantageous compared to other approaches because they require very low sample volumes. Our novel geometries are compatible with *in situ* crystal growth which eliminates crystal handling and enable tuning of crystal densities and sizes to achieve high hit rates and high-quality diffraction images. Furthermore, they allow for the use of non-jetable crystal morphologies, like needles or plates, or crystals of heterogeneous size. They can also be used to facilitate dynamic experiments using stimuli like electric fields gradients, temperature jumps, pH gradients, ligand exchange and photo-activation.

For fixed targets, the choice of construction materials and fabrication strategy are important considerations as they

significantly impact the cost, fragility, and stability of the devices. Several materials such as low-Z polymers (COC, PMMA, PDMS, Kapton, Mylar, epoxies), silicon, silicon nitride, glass, and quartz have been used to make devices that balance these attributes with the measurement requirements. The two most common design formats are (1) microgrids and (2) microfluidic chips. Microgrids are chips with a 2D array of micropatterned holes for crystal entrapment upon deposition, sometimes used with thin films supports (0.1–10  $\mu\text{m}$ ) to improve sample retention and/or protect against dehydration.<sup>17,27–30,33–46</sup> These chips can offer the advantages of high hit-rates and ultra-low background if wicking or vacuum application is used to localize crystals into pores and remove excess crystallization solution, but the exposed sample (even with sandwiching thin films) is sensitive to dehydration, requiring sample preparation and assembly shortly before measurements (<1–2 hours). Microfluidic chips, on the other hand, are enclosed devices offering precise control over sample volume and thickness, long-term stability against evaporation, and *in situ* crystal growth to eliminate crystal handling. But the thick flow channels (25–300  $\mu\text{m}$ ) and capping layers (25–600  $\mu\text{m}$ ) often used to construct these devices contribute significantly to background scatter.<sup>18,47–57</sup> While several such devices have demonstrated high-resolution data collection from large, well-ordered protein crystals, for many proteins obtaining large crystals often proves intractable. In the case of these small or weakly-diffracting microcrystals (desired for SX), the X-ray attenuation and scatter background from typical microfluidic chips can limit the resolution attained. Therefore, there is a need for the development of novel fixed targets with the stability and ease-of-use of microfluidic approaches, that are inexpensively fabricated and easily modified to match different sample and beamline requirements, while maintaining the advantages of high hit-rates and low background of microgrid approaches.

In this paper we describe the design of polymer fixed-target chips that address this need for routine and reliable RT SOX and SSX/SFX experiments. The polymer materials used were selected for low water permeability, high X-ray transparency, and high optical transparency (for on-chip imaging and light triggering of structural changes for future time-resolved structural studies). A modular, layered fabrication process enabled control over the cross-section of materials in the beam path and easy design modification for adaption to different sample and beamline requirements. The chip is compatible with *in situ* crystallization using micro-batch and vapor diffusion methods. Pilot X-ray data collection with the chips was performed using SOX. The robustness and versatility of the chips was demonstrated, showing that they provide long-term storage, stability, easy transportation, and straightforward on-chip crystallization to diffraction measurements. Preliminary results also indicate that the chip is amenable to SFX measurements without further alteration.

## Materials and methods

A layered assembly process was used to construct the polymer microfluidic chip from hot-embossed COC supports, spin-coated COC thin films, laser-cut PMMA frames, and an adhesive sample flow layer. Five proteins (hen egg white lysozyme, thaumatin, bovine liver catalase, concanavalin-A, and SARS-CoV-2 nonstructural protein NSP5) were crystallized *in situ* and diffraction data directly collected on-chip at RT at beamline 12-1 at SSRL. The minimum beam spot size used was  $55\ \mu\text{m} \times 5\ \mu\text{m}$  (X-Y, FWHM) and a SOX data collection approach was used to benchmark the performance of the chip.

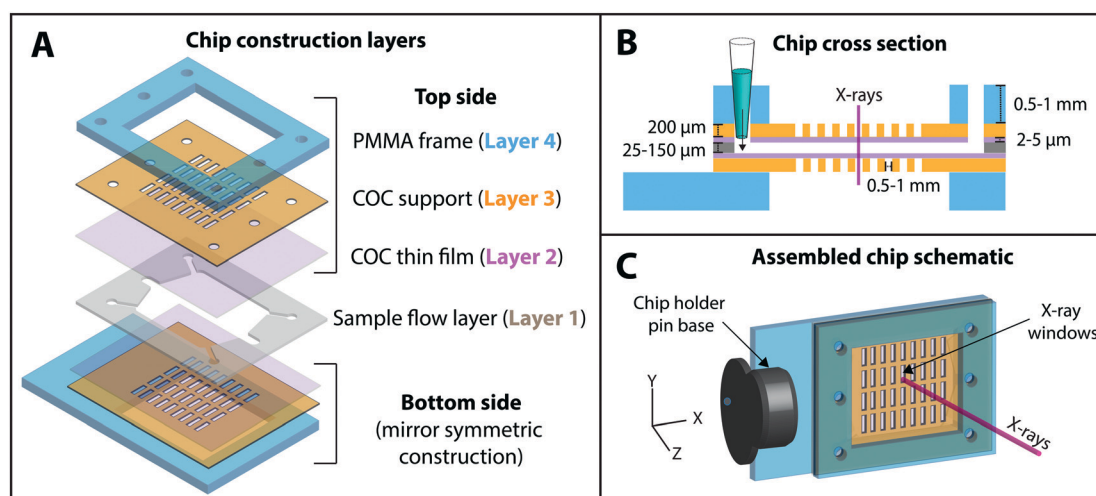
### Microfluidic chip fabrication and assembly

A schematic of the chip construction layers, and the final assembled chip are shown in Fig. 1. The X-ray imaging regions of the chip were made of cyclic olefin copolymer (TOPAS® COC, Grade 8007,  $T_g = 75\ ^\circ\text{C}$ ). The sample flow layer (layer 1) consisted of a CO<sub>2</sub> laser cut, tunable acrylic or silicone pressure-sensitive adhesive (25  $\mu\text{m}$  AR 92734 or 48  $\mu\text{m}$  AR 92712, Adhesives Research Inc., Glen Rock, PA, USA) used to bond the two microfabricated sides (top and bottom) together. 2–5  $\mu\text{m}$  COC thin films (layer 2) provided low-background sample enclosing layers. These were prepared by spin-coating solutions of 10–20 wt% COC dissolved in *sec*-butylbenzene on UV-ozone treated silicon wafers. Films ranging from 500 nm to 10  $\mu\text{m}$  in thickness could be easily produced by varying the COC concentration and spin speed (Fig. S1†). The 200  $\mu\text{m}$  thick COC support with windows (layer 3) was hot embossed using a mold made of elastic polydimethylsiloxane (PDMS, details of the mold fabrication and hot embossing below). And finally, 0.5 or 1 mm polymethyl methacrylate (PMMA) frames (layer 4) with an adhesive layer (3M™ F9460PC) were produced by CO<sub>2</sub> laser cutting.

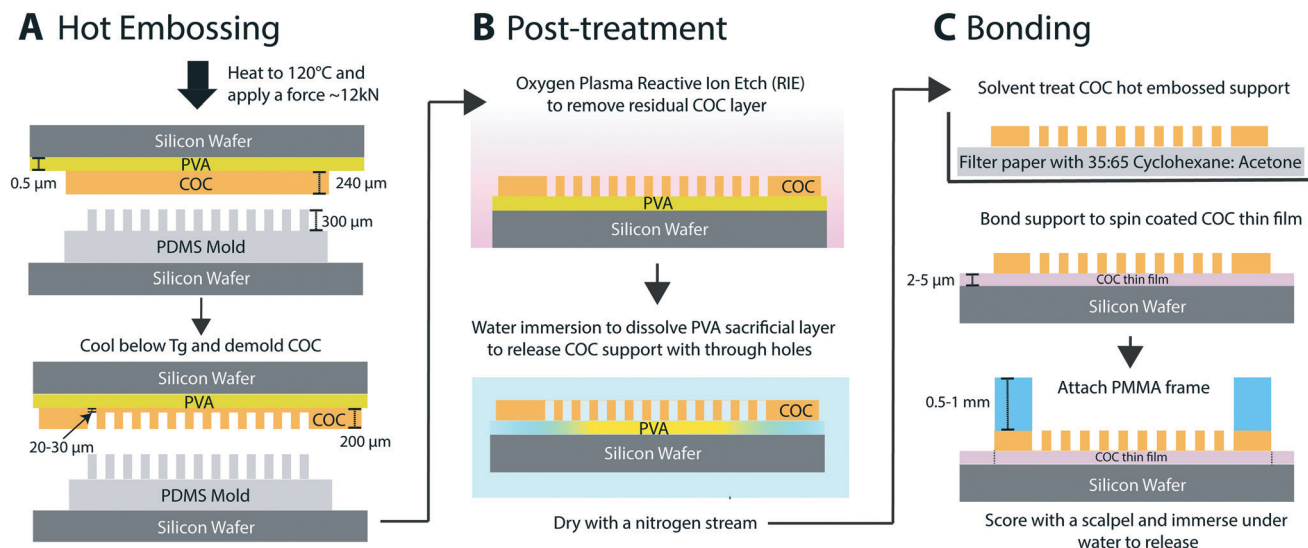
The use of hard mold materials like steel, silicon, or high-temperature epoxy used to emboss COC in previous works<sup>49,50,58</sup>

proved difficult due to feature entanglement and warping while demolding the COC sheets with 200  $\mu\text{m}$  deep through-holes from the rigid molds. PDMS molds proved successful due to their elasticity and low adhesion to COC. These molds were fabricated by casting a replica of a silicon master mold with an array of 300  $\mu\text{m}$  deep X-ray window features using a 5:1 monomer:curing agent mixture of Dow Sylgard™ 184 (fabrication of Si master and additional process details provided in ESI† Section S1). The PDMS replica was demolded and bonded to a silicon wafer using oxygen plasma treatment (50 W, 25 sccm O<sub>2</sub>, 0.79 Torr, 1 minute). Hot embossing was performed using a semi-automated EVG 501 wafer bonder. The PDMS mold was brought into contact with a 240  $\mu\text{m}$  COC sheet (Europlex 0F304, Roehm America LLC, Sanford, ME, USA) attached to a polyvinyl alcohol (9 wt% PVA, 1500 rpm, 60 s) coated silicon wafer. The assembly was heated to 120  $^\circ\text{C}$  and a force of 12 kN was applied for 15 minutes to emboss window in the supports. The embossed sheet was demolded after cooling below the glass transition temperature of COC. Obtaining perfect through holes with hot embossing was difficult due to the flexibility of the PDMS mold. Instead, the ~20–30  $\mu\text{m}$  residual thin film in the windows features was removed by an oxygen plasma reactive-ion etching process (500 W, 25 sccm O<sub>2</sub>, 330 mTorr, 30 minutes) to yield the COC window support with through-holes (layer 3). A schematic of the fabrication steps is shown in Fig. 2A and B.

After fabricating the various layers as described above, Fig. 2C shows a schematic of the processing steps involved in assembling the two symmetric sides of the chip (top and bottom, Fig. 1). First, a COC window support (layer 3) was solvent treated with 35:65 acetone:cyclohexane solution for 1 minute to render the surface tacky,<sup>59</sup> dried with a nitrogen stream, and brought into contact with a spin-coated COC thin film (layer 2) on a silicon wafer. This process created a strong room temperature bond between layers while maintaining the integrity of the thin film (layer 2). The assembly was further



**Fig. 1** A schematic of (A) the different construction layers used to assemble the polymer microfluidic chip; (B) the cross-section view showing the different layer thicknesses; and (C) the final assembled chip.



**Fig. 2** A schematic of the fabrication scheme used to produce the top and bottom of the microfluidic chip (as shown in Fig. 1). (A) Hot embossing was performed using an elastic PDMS mold to imprint an array of “X-ray window” features in COC sheets (240  $\mu\text{m}$ ). (B) The residual thin film (20–30  $\mu\text{m}$ ) from the embossing step was removed using oxygen plasma reactive ion etching (RIE) to create COC supports ( $\sim 200$   $\mu\text{m}$ ) with through-hole windows. The supports were detached from the silicon wafer by dissolving the PVA sacrificial layer in water. (C) To assemble each side (top and bottom), a COC support was solvent treated to facilitate bonding to COC thin films of desired thicknesses (2–5  $\mu\text{m}$ ), followed by attachment of a 0.5–1 mm thick PMMA frame using an adhesive layer to provide rigidity and flatness.

reinforced by attaching 0.5 or 1 mm thick PMMA frames (layer 4) on the other side to improve planarity, prevent bowing of the thinner layers, and make the chip easy to handle. At this stage, the assembled layers (layers 2–4) were robust and could be stored until needed. This was advantageous as the flow layer (layer 1) could be varied in thickness to tailor the chip for a particular protein crystal size or beamline spot size on demand. Both thin (25–50  $\mu\text{m}$ ) or thick (80–150  $\mu\text{m}$ ) flow layers could be used depending on sample requirements, *e.g.*, ultra-small protein crystals benefit from thinner flow layers to decrease background from excess crystallization solution, while efficient slurry loading of larger crystals necessitates thicker flow layers. The hydrophobic COC films needed to be rendered hydrophilic prior to final assembly to facilitate fast and complete solution loading into the chip. To do this, assembled top and bottom sides (layers 2–4) were exposed to atmospheric plasma treatment for 3 minutes in a Harrick PDC-32G Basic Plasma Cleaner. Advancing water contact angles using a Ramé-Hart goniometer were  $83 \pm 6^\circ$  for the native COC surface,  $22 \pm 5^\circ$  after plasma treatment, and recovered to  $\sim 60^\circ$  upon storage in ambient conditions for 2–4 weeks (Fig. S2†). To complete the chip fabrication process, the two chip sides were bonded using a laser-cut, double-sided adhesive sample flow layer of the desired thickness. The holes or “window” features (0.5 mm  $\times$  1.75 mm or 1 mm  $\times$  1 mm) in the two sides of the chip were aligned by maximizing light transmission through the features using a backlight. The aligned sides were manually pressed together to create an enclosed microfluidic chip. A contact dwell time of 24–72 hours was required to ensure strong adhesion between all the layers before using the chip. A more detailed protocol on the individual microfabrication steps is provided in the ESI† Section S1.

### Water permeability measurements

Water vapor transmission rate as a function of COC film thickness was measured using a slightly modified version wet-cup tests described elsewhere.<sup>60</sup> In brief, free standing COC thin films of different thickness were solvent bonded to a 240  $\mu\text{m}$  COC sheet with a 1 cm diameter hole in the center and affixed to the opening of a 3 mL clear glass vial containing Millipore water using Dow Corning® high vacuum grease. Sample weight loss was monitored over a period of one week from five replicates. Steady-state water vapor transmission rates (WVTR) were calculated from the weight loss measurements using  $\text{WVTR} = \frac{\Delta m}{A \times t}$ , where  $\Delta m$  was the weight loss,  $A$  is the area of the membrane, and  $t$  is the time. The water vapor permeability (WVP) of COC, which is a function of the solubility and diffusivity of water in the material, was calculated by fitting the experimental data to  $\text{WVTR} = \frac{\text{WVP} \times \Delta P}{L}$  where  $\Delta P$  was the differential pressure of water vapor across the membrane of thickness  $L$ . The measurements were carried out in a climate-controlled room with a relative humidity of  $20 \pm 2\%$  at  $23 \pm 2^\circ\text{C}$ . The relative humidity gradient ( $\Delta\text{RH}$ ) was approximately 80% assuming the relative humidity inside the sealed vial enclosure was close to 100%, resulting in  $\Delta P = P \times \Delta\text{RH} \cong 2.2$  kPa, using a water vapor saturation pressure of 2.8 kPa at  $23^\circ\text{C}$ .

### Protein expression, purification, and crystallization

SARS-CoV-2 main protease nonstructural protein NSP5 was expressed and purified using a modified protocol as previously described.<sup>61</sup> In short, NSP5 was expressed from PGEX-6p-1-NSP5 plasmid (kindly provided by R. Hilgenfeld, University of Lübeck, Germany), from *E. coli* BL21 DE3 Gold



cells in 2YT media overnight at 18 °C. Freshly streaked plates from transformed glycerol stocks were used for inoculation. The harvested cell pellets were stored at –80 °C until purification. The pellet was thawed in buffer A (20 mM Tris pH 7.8, 150 mM NaCl, 5 mM imidazole) and disrupted by sonication at 4 °C (2 min total) and two passes through a cell disruptor (12–14 kPa). The suspension was clarified by centrifugation (30 kg, 45 min), filtered (0.8 µm) and loaded onto a 5 mL HisTrap FF column (Cytiva). The column was washed with 5 CV of buffer A and the protein eluted in 5 mL fractions of buffer B (20 mM Tris pH 7.8, 150 mM NaCl, 300 mM imidazole). The protein was dialyzed against buffer C (20 mM Tris pH 7.8, 150 mM NaCl, 0.5 mM TCEP) concurrently with GST tag removal by overnight digestion with 10% w/w HRV3C protease. The protein was loaded into a 5 mL HisTrap FF column and the flow through collected in 12.5 mL fractions. The protein was further purified by size exclusion chromatography (Superdex 200 pg, Cytiva) in buffer D (20 mM Tris pH 7.8, 150 mM NaCl, 0.5 mM TCEP, 1 mM EDTA). The protein was concentrated to 10 mg mL<sup>–1</sup> and crystallized *via* vapor diffusion with 100 mM Bis-Tris pH 6.5, 17.5% w/v PEG 3350, 175 mM Li<sub>2</sub>SO<sub>4</sub>. Seed stock was generated from the resulting large plate clusters using Hampton Research's PTFE seed beads and diluted by a factor of one hundred with crystallization buffer. The vapor diffusion conditions were spiked with 10% 1:100 seed stock and this process was repeated to generate second generation seeds. To crystallize NSP5 on-chip, the same condition used to generate second generation seeds was loaded into a microfluidic chip with 10% second generation seeds (Table 1) and crystallized *via* vapor diffusion as described below.

Chicken egg-white lysozyme (#L6876), thaumatin from *Thaumatococcus daniellii* (#T7638), concanavalin A from *Canavalia ensiformis* (Jack bean) type VI (#L7647) and catalase from bovine liver (#C40) were purchased from Millipore Sigma (St. Louis, MO, USA) and dissolved in MilliQ water or low ionic strength buffers as listed in Table 1. The protein solutions were gently vortexed for a few seconds until the solution was well-mixed, centrifuged at 10 000 rpm for 5 minutes to remove any insoluble materials, and the supernatant was removed and stored at 4 °C. All buffers and precipitant solutions were filtered through a 0.22 µm syringe filter prior to use. The crystallization conditions used in this work were adapted from previous literature<sup>62–64</sup> and are also reported in Table 1.

Direct crystal slurry loading requires filtration to ensure removal of large crystals which can otherwise clog the inlet channel. While this was tested, this work mainly focusses on demonstration of on-chip crystallization. The microfluidic chips were loaded with a well-mixed 1:1 solution of protein and precipitant solution by pipetting ~8–10 µL into one of the inlet holes in the PMMA frame. Corner vents in the spacer flow layer ensured that the solution filled the wide fluid chamber (~10 mm × 10 mm × 25–50 µm) uniformly

while minimizing bubble entrapment. After filling, the inlets and outlets were sealed using Hampton crystal clear sealing tape for micro-batch crystallization or stored unsealed for vapor diffusion crystallization to allow for equilibration with the reservoir chamber. The filled chips were then placed in a Falcon 6-well tissue culture plate either on a microbridge-like pedestal or affixed to a small magnet on the wall of the well plate using magnetic chip holder pin bases (Crystal Positioning Systems, Jamestown, NY, USA). The well was filled with 1.5–2 mL of precipitant solution and the plate was sealed using crystal clear sealing tape to ensure that the chamber remained humidified to prevent sample desiccation during storage over several days to weeks. The optically transparent microfluidic chip and well sealing tape enabled on-chip sample monitoring without disturbing the equilibrated enclosure.

### On-chip X-ray diffraction

X-ray diffraction data were collected at RT on the 12-1 beamline at SSRL, equipped with an Eiger X 16M detector (Dectris AG). A magnetic chip holder pin base with a slot and set screw was used to securely hold the microfluidic chip before magnetically mounting it on the goniometer. Inline high-resolution cameras at ~0° and 90° orientation to the beam were used to position the chip in the beam path, scan through sample regions/windows, and to center the protein crystals along the rotational axis. The beamline allowed for a translation range of ±2.5 mm along the Y-direction (vertical) and greater than ±7.5 mm along the X-direction (horizontal). The close proximity of an upstream microcollimator limited the range of rotation about the axis (Z) to ±35°. The smallest beam size used was 55 µm × 5 µm (X–Y, FWHM), with Y varied between 5 to 50 µm based on crystal size and morphology to maximize the sample-beam interaction cross-section.

Data was collected remotely using the Blu-Ice package.<sup>65</sup> Individual single crystals were manually centered and 30° rotation wedges with 1° oscillation at 0.1 s exposure per frame were collected at 10–20% transmission (full photon flux 4 to 5.6 × 10<sup>12</sup> photons per second at 12.5 keV) with a 200 mm detector distance. A drop off in diffraction resolution was observed by the 20–25th frame due to cumulative radiation damage. Data from 15–30 crystals was collected from each chip with the final number of crystals merged for each protein reported in Table 2. Diffraction data from multiple crystals was processed in xia2<sup>66</sup> (multiplexing mode) running the CCP4<sup>67</sup> and DIALS<sup>68</sup> packages to perform indexing, merging, and scaling. Structures were solved with Phaser,<sup>69</sup> part of the PHENIX package,<sup>70</sup> using PDB entries 1VED, 1RQW, 8CAT, 1SCR, and 6XR3 as templates for phasing *via* molecular replacement for lysozyme, thaumatin, concanavalin-A, catalase and NSP5 respectively. Iterative refinement was performed with *phenix.refine*<sup>71</sup> alternating with molecular modeling performed with Coot.<sup>72</sup> Final data processing and structure refinement statistics are provided in Table 2.

**Table 1** A summary of crystallization conditions used in this work and a comparison of crystal size, morphology, density, and time to first crystals, obtained from on-chip crystallization and comparable crystallization in 24-well crystallization plates. For on-chip crystallization,  $\sim 8 \mu\text{L}$  of solution was loaded into the chip and the chip was stored either sealed (micro-batch) or unsealed (vapor diffusion) in an enclosed environment with the precipitant solution in the reservoir. Similar droplet volumes were used for crystallization in a 24-well plate format using micro-batch under oil (paraffin) or hanging drop vapor diffusion methods for comparison, as appropriate. Crystal density ( $n = \#$  of crystals per  $\text{mm}^2$ ) was measured using optical microscopy and qualitatively described as low for  $n \leq 10$ , medium for  $10 < n < 100$ , and high for  $n \geq 100$

Protein	Protein solution	Precipitant solution	Incubation temperature	Crystallization method	Crystal size, morphology, density, crystallization time	
					On-chip	24-Well crystallization plate
Lysozyme	30 $\text{mg mL}^{-1}$ in 20 mM sodium acetate buffer, pH 4.6 50 $\text{mg mL}^{-1}$ in DI water	1 M sodium chloride, 0.1 M sodium acetate buffer, pH 4.6 2 M sodium chloride, 0.1 M sodium acetate buffer, pH 4.6	4 °C	Micro-batch	25–35 $\mu\text{m}$ , cuboid, high, $\sim 2$ –3 hours	25–35 $\mu\text{m}$ , cuboid, high, $\sim 2$ –3 hours
			21 °C	Micro-batch	60–100 $\mu\text{m}$ , cuboid, low, 12–24 hours	30–70 $\mu\text{m}$ , cuboid, medium, 2–4 hours
Thaumatococcus	25 $\text{mg mL}^{-1}$ in DI water	1 M L-sodium potassium tartrate, 0.1 M ADA buffer, pH 6.5	4 °C	Micro-batch	$\sim 25 \mu\text{m}$ , tetragonal bipyramidal, low, $\sim 12$ hours. Crystals grow to 70–100 $\mu\text{m}$ in size over 2–3 days.	$\sim 20 \mu\text{m}$ , tetragonal bipyramidal, high, $\sim 12$ hours
			21 °C	Micro-batch	50–100 $\mu\text{m}$ , tetragonal bipyramidal, low, 3–4 days	Did not crystallize
			21 °C	Vapor diffusion	50–100 $\mu\text{m}$ , tetragonal bipyramidal, low, $\sim 24$ hours	50–150 $\mu\text{m}$ , tetragonal bipyramidal, low, $\sim 24$ hours
			21 °C	Micro-batch (with seeding using 1° seed stock 1 : 5 : 5 seed : protein : precipitant) Micro-batch	15–20 $\mu\text{m}$ , tetragonal bipyramidal, high, $\sim 24$ hours	15–20 $\mu\text{m}$ , tetragonal bipyramidal, high, $\sim 24$ hours
Concanavalin-A (type VI)	70 $\text{mg mL}^{-1}$ in 20 mM Tris buffer, pH 8.0	2.8 M ammonium sulfate in 0.1 M Tris buffer, pH 8.5	21 °C	Micro-batch	150–300 $\mu\text{m}$ , round base rhombic tetrahedron, low, $\sim 12$ hours	30–60 $\mu\text{m}$ , cubic, high, 0–2 hours initially; equilibrates to 150–300 $\mu\text{m}$ , round base rhombic tetrahedron, low, over the next 3–7 days
Bovine liver catalase	40 $\text{mg mL}^{-1}$ in 50 mM sodium phosphate buffer, pH 6.8	22.5% PEG 4000 in 0.1 M Tris buffer, pH 8.5	21 °C	Micro-batch	70–120 $\mu\text{m}$ , prism shaped, low, $< 12$ hours	70–120 $\mu\text{m}$ , prism shaped, low, 2–4 hours
			21 °C	Vapor diffusion	30–100 $\mu\text{m}$ , prism shaped, medium, 0–4 hours	30–100 $\mu\text{m}$ , prism shaped, medium, 0–2 hours
NSP5	5 $\text{mg mL}^{-1}$ in 20 mM Tris pH 7.8, 150 mM NaCl, 0.5 mM TCEP, 1 mM EDTA	100 mM bis-tris pH 6.5, 17.5% w/v PEG 3350, 175 mM; $\text{Li}_2\text{SO}_4$ ; 10% 1:100 2' seeds	21 °C	Vapor diffusion, with seeding	300 $\mu\text{m}$ , large thin plate clusters, low, $\sim 12$ hours	150–200 $\mu\text{m}$ large single plates or plate clusters, low, $\sim 12$ hours

## Results and discussion

### Microfluidic chip fabrication and device performance

The goal of this work was to develop robust, user-friendly, low-background microfluidic chips to support room temperature serial crystallography experiments at synchrotrons or XFEL facilities. A modular fabrication process was designed and used to construct large-area polymer chips that could deliver hundreds to thousands of hydrated protein microcrystals to the X-ray beam. An exemplary chip design

(overall dimensions 25 mm  $\times$  15 mm) with a 4  $\times$  9 array of rectangular X-ray windows (0.5 mm  $\times$  1.75 mm) is shown in Fig. 3A. The X-ray imaging regions of the chip were made of COC, a thermoplastic material with excellent water barrier properties, optical transmissivity, chemical compatibility with acids, bases and alcohols,<sup>73</sup> and low X-ray attenuation.<sup>58</sup> Microfluidic chips based on this material have been reported previously, with designs using either injection molding to make mm-thick devices, or hot embossing to produce few-hundred  $\mu\text{m}$ -thick devices.<sup>48,49,51,52,58</sup> The characterization of

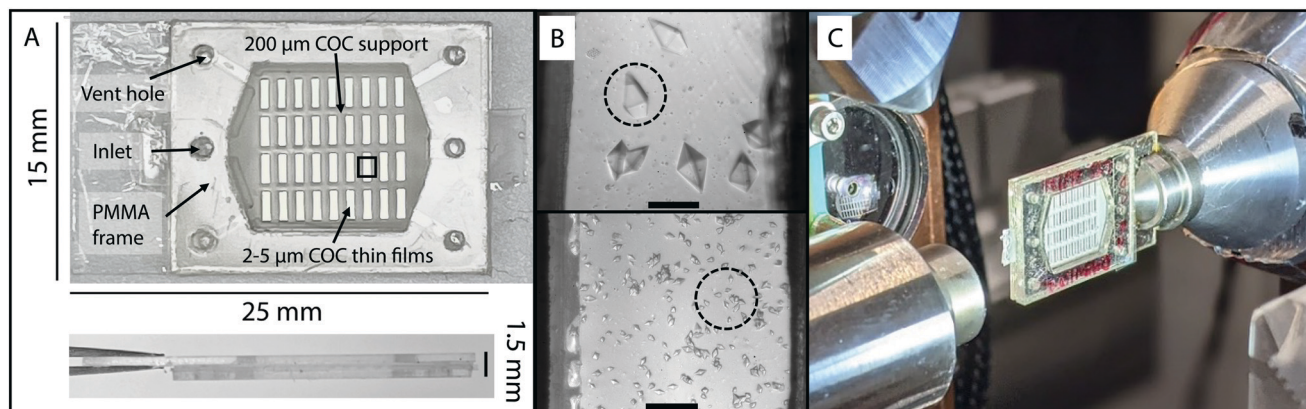
**Table 2** Crystallographic statistics obtained from lysozyme, thaumatin and concanavalin-A, catalase and NSP5 crystals grown in the microfluidic chip. Values in parentheses correspond to the highest resolution shell

Protein	Lysozyme (fresh)	Lysozyme (aged)	Thaumatin	Concanavalin-A	Catalase	NSP5
Number of crystals used	7	6	16	12	7	11
Average crystal size (largest dimension)	~30–35 $\mu\text{m}$	~70 $\mu\text{m}$	~100 $\mu\text{m}$	~200 $\mu\text{m}$	~90–100 $\mu\text{m}$	~200 $\mu\text{m}$
Resolution range ( $\text{\AA}$ )	56.05–1.87 (1.94–1.87)	35.25–1.50 (1.55–1.50)	58.61–1.45 (1.48–1.45)	51.18–1.30 (1.35–1.30)	52.8–2.274 (2.355–2.274)	31.64–2.7 (2.797–2.7)
Unit cell dimensions	$a = b = 79.26 \text{ \AA}$ , $c = 37.94 \text{ \AA}$ , $\alpha = \beta = \gamma = 90^\circ$	$a = b = 78.82 \text{ \AA}$ , $c = 38.21 \text{ \AA}$ , $\alpha = \beta = \gamma = 90^\circ$	$a = b = 58.60 \text{ \AA}$ , $c = 151.44 \text{ \AA}$ , $\alpha = \beta = \gamma = 90^\circ$	$a = 63.20 \text{ \AA}$ , $b = 87.22 \text{ \AA}$ , $c = 89.12 \text{ \AA}$ , $\alpha = \beta = \gamma = 90^\circ$	$a = b = 141.77 \text{ \AA}$ , $c = 103.47 \text{ \AA}$ , $\alpha = \beta = 90^\circ$ , $\gamma = 120^\circ$	$a = 114.29 \text{ \AA}$ , $b = 54.81 \text{ \AA}$ , $c = 45.33 \text{ \AA}$ , $\alpha = \gamma = 90^\circ$ , $\beta = 101.45^\circ$
Space group	$P4_32_12$	$P4_32_12$	$P4_12_12$	$I222$	$P3_221$	$C121$
<i>Data processing statistics</i>						
Total reflections	120 310 (12 080)	331 623 (33 168)	1 392 515 (65 410)	786 259 (24 955)	1 040 538 (97 971)	49 766 (5027)
Unique reflections	10 449 (1003)	19 849 (1949)	47 884 (2332)	60 428 (5732)	55 166 (5256)	7530 (747)
Multiplicity	11.5 (12.0)	16.7 (17.0)	29.1 (28.0)	13.0 (4.4)	18.9 (18.6)	6.6 (6.7)
Completeness (%)	99.82 (98.91)	99.97 (99.95)	100.0 (100.0)	98.8 (89.5)	99.56 (95.89)	96.30 (93.18)
$\langle I/\sigma(I) \rangle$	11.47 (2.02)	16.80 (1.59)	15.4 (0.9)	17.6 (0.6)	10.64 (1.05)	8.24 (1.78)
$R_{\text{merge}}$	0.159 (1.63)	0.09752 (1.953)	0.146 (3.606)	0.072 (2.107)	0.218 (2.587)	0.248 (2.01)
$R_{\text{meas}}$	0.167 (1.703)	0.1007 (2.013)	0.149 (3.671)	0.075 (2.384)	0.224 (2.658)	0.2691 (2.183)
$R_{\text{pim}}$	0.048 (0.481)	0.02467 (0.4847)	0.027 (0.677)	0.019 (1.055)	0.051 (0.606)	0.097 (0.795)
$\text{CC}_{1/2}$	0.994 (0.602)	0.999 (0.773)	0.999 (0.647)	0.993 (0.159)	0.997 (0.576)	0.968 (0.267)
Wilson $B$ -factor	28.48	22.45	19.9	17.8	48.65	55.59
<i>Refinement statistics</i>						
$R_{\text{work}}$ (%)	15.61	15.02	15.2	14.6	16.2	18.0
$R_{\text{free}}$ (%)	17.9	17.13	16.0	16.5	20.1	21.8
RMS (bonds, $\text{\AA}$ )	0.005	0.008	0.022	0.004	0.003	0.003
RMS (angles, $^\circ$ )	0.59	0.84	1.59	0.83	0.58	0.53
<i>Ramachandran statistics</i>						
Favored (%)	99.21	99.21	98.0	97.8	95.85	96.04
Allowed (%)	0.79	0.79	2.0	2.2	3.94	3.96
Outliers (%)	0	0	0	0	0.20	0
Average $B$ -factor						
Macromolecule	33.30	28.13	21.9	22.7	56.46	59.64
Ligands	—	—	—	14.6	55.17	—
Solvent	36.39	36.13	38.6	34.6	46.48	49.38

small or weakly-diffracting microcrystals in these devices can prove challenging due to significant background from thick enclosing and flow layers, limiting the diffraction resolution attained. Therefore, a key focus of our design was to minimize the cross sectional thickness to maximize signal to noise from crystals. This was achieved by using a layered assembly process to produce physically robust and easy-to-handle chips with a total thickness of ~1.5 mm but an effective cross sectional thickness of only 30–60  $\mu\text{m}$  in the X-ray window regions.

The chip consisted of two sides (Fig. 1, layers 2–4) mirror symmetric in construction, bonded using a tunable pressure-sensitive adhesive spacer layer that defined the flow layer thickness (layer 1, 25 or 48  $\mu\text{m}$  in this work). Windows in the 200  $\mu\text{m}$  hot-embossed COC support had ultra-thin 2–5  $\mu\text{m}$  COC films attached that served as the low-background X-ray interaction regions. The outermost 0.5–1 mm thick PMMA frame attached to the COC support imparted rigidity and included inlet, outlet, and vent holes for introduction of crystallization cocktails or crystal slurries into the chip by micro-pipetting. A major advantage of this approach was the

straightforward incorporation of tunable-background, wrinkle-free thin films in an enclosed flow-chip design. Solvent bonding eliminated the need to handle fragile films, as is typically required for alternatives like the single/few-layer graphene (<1–3 nm, expensive and laborious to produce),<sup>42,45,74</sup> freestanding commercially-available Mylar (2.5–3.5  $\mu\text{m}$ )<sup>38,40</sup> or Kapton (3–8  $\mu\text{m}$ )<sup>34</sup> films used as enclosing layers in open-format fixed-target chips. The modular construction of the chip allows rapid modifications to the chip design, *e.g.*, the sample area available for rastering, sample volume, and thickness of enclosing and films, depending on sample and experimental requirements. Sample fabrication and assembly of 12 chips took approximately 1 hour, but parallel processing could further reduce the fabrication time. For applications that do not require ultra-thin supports (<5  $\mu\text{m}$ ), the reactive-ion etch step could be omitted after hot-embossing COC sheets to retain ~20–30  $\mu\text{m}$  residual thin films over X-ray windows. This decreases the number of fabrication steps and manufacturing time, but at the cost of proportionally higher background scatter.



**Fig. 3** (A) A macroscopic view of the assembled polymer microfluidic chip with a  $4 \times 9$  array of rectangular X-ray windows in the COC supports, to which COC thin films are attached. Features for sample introduction are highlighted in the outermost layer of the chip, the PMMA frame. (B) Optical microscopy images of tetragonal thaumatin crystals grown directly on-chip using micro-batch crystallization as observed through the optically transparent X-ray windows in the chip. The top and bottom figures correspond to crystallization without and with seeding to control crystal size and density. The scale bar is  $100 \mu\text{m}$ . (C) The chip magnetically mounted on the SMB goniometer at SSRL using a magnetic chip holder pin base. Experiments were performed at room temperature without the need for any external humidification.

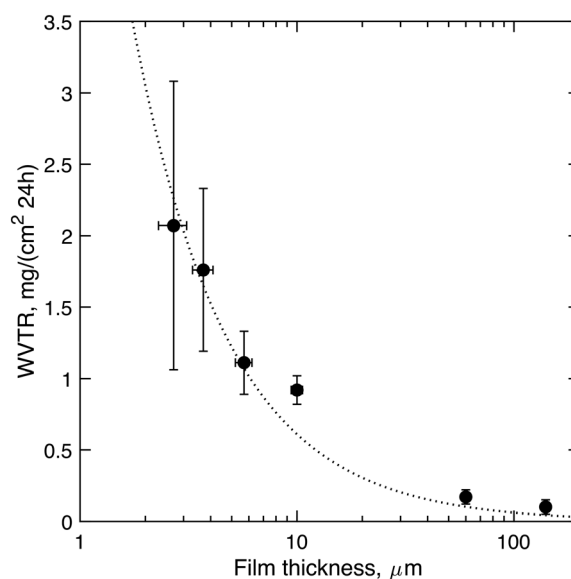
Hydrophilic oxygen plasma surface treatment provided facile loading of aqueous solutions into the chip as shown in Fig. S3.† The advancing contact angle of atmospheric plasma treated COC films was  $\sim 22^\circ$  for freshly treated films and increased to  $\sim 60^\circ$  over a period of 4 weeks as shown in Fig. S2.† Thus, fully assembled chips could be stored for at least one month before use without significantly impacting solution loading.

To identify the timescales over which diffraction measurements could be carried out on our chips without external humidification, measurements of water evaporative loss through thin COC films were performed using a modified wet-cup method. Fig. 4 shows the steady-state water vapor transmission rate (WVTR) through COC films as a function of film thickness. The WVTR measured was inversely proportional to the film thickness, as expected for diffusive transport. Film thicknesses ranging from 2.5 to  $140 \mu\text{m}$  were included to serve as a guideline for users to choose barrier properties as required for their application, keeping in mind that scatter background would scale linearly with film thickness. For the 2–5  $\mu\text{m}$  films used in this work, a WVTR of  $\sim 2 \frac{\text{mg}}{\text{cm}^2 \text{ day}}$  translated to a low evaporative loss of 0.5–1 wt% per hour, ensuring that crystals would remain hydrated without external humidity control for several hours during diffraction measurements. To extend sample stability to several weeks, we stored the chips in individually sealed wells with saturated salt slurries of  $\text{K}_2\text{SO}_4$  (RH set point  $\sim 97.3\%$  at  $25^\circ\text{C}^{75}$ ) or with the precipitant solution to reduce the relative humidity gradient against which the chips were equilibrating. We found that under these conditions, chips with crystals could be stored over a period of 2–4 weeks without visible or measurable loss of crystallinity despite the small sample volume in the chip ( $<10 \mu\text{L}$ ). In comparison, previous microfluidic chips used thick ( $>50$ – $200 \mu\text{m}$ ) sealing layers of COC, Kapton or Mylar and were stable for weeks in ambient conditions. Thin support films ( $\sim 2.5$ – $3.5 \mu\text{m}$  Mylar or  $3$ – $8 \mu\text{m}$  Kapton) have been used previously as enclosing

layers with microgrids, but these approaches are seldom designed or characterized for long term stability, requiring sample preparation at the beamline.<sup>34,38,40</sup>

### On-chip protein crystallization

To test the utility of the chips for *in situ* crystal growth and long-term storage, vapor diffusion and micro-batch crystallization trials were conducted for five different proteins



**Fig. 4** Steady-state water vapor transmission rates (WVTR) through COC films measured using a modified wet-cup method at  $23^\circ\text{C}$ , against a  $\Delta\text{RH}$  gradient of approximately 80%. COC films in the thickness range of 2–5  $\mu\text{m}$  that were used in this work as sample enclosing layers (layer 2) have a low WVTR of  $\sim 2 \frac{\text{mg}}{\text{cm}^2 \text{ 24h}}$  or  $\sim 0.5$ –1% sample wt loss per hour of storage in ambient room temperature conditions without external humidity control. The fitted line corresponds to a water vapor permeability (WVP) of  $\sim 10^{-14}$ – $10^{-13} \frac{\text{g m}}{\text{m}^2 \text{ s Pa}}$ .



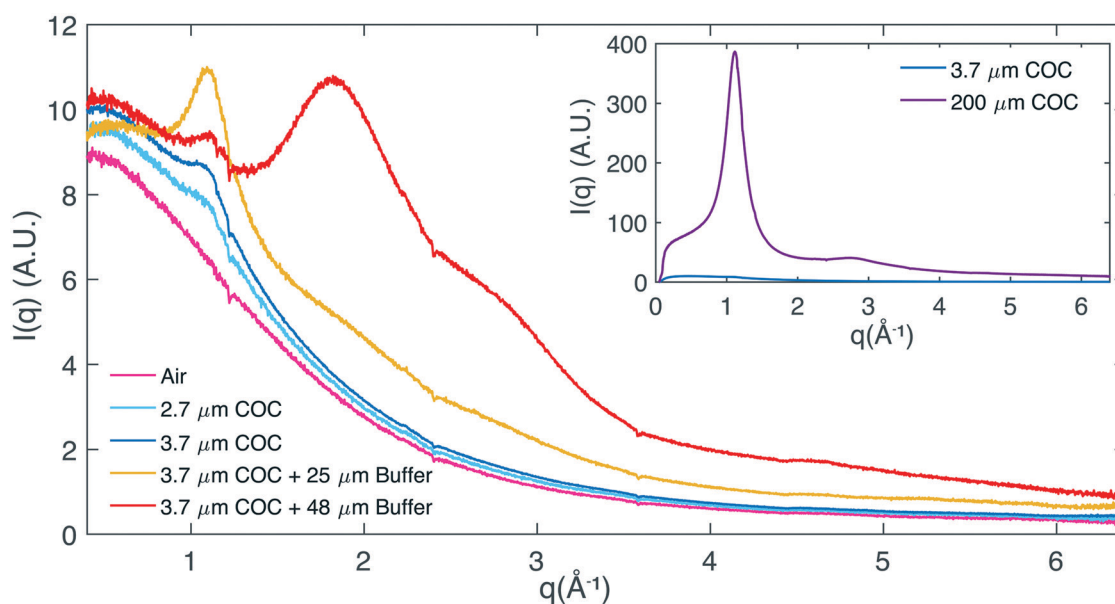
by adapting existing conditions from the literature.<sup>61–64</sup> Table 1 summarizes the crystallization conditions used and compares observations of crystal size, morphology, densities, and time-to-crystallization between on-chip and on-crystallization plate methods (hanging drop vapor diffusion or micro-batch under oil methods were used for comparison as appropriate). For most of the conditions tested, similar crystal sizes and morphologies were obtained on and off chip without the need to modify established crystallization conditions. On average, slightly lower nucleation rates and slower crystallization kinetics were observed on-chip leading to larger crystals with sparser densities. Compared to the large droplet solution geometry in plate-based methods, solution in a microfluidic chip is confined to the micro-scale in one or more dimensions. In the near-2D geometry of our chip, a large interfacial contact area between the solution and the polymer surface is expected to affect crystal nucleation. The slower crystallization kinetics are due to the smaller evaporation rate in the chips and that mass transport is primarily driven by molecular diffusion as buoyancy driven convection is suppressed in this geometry.<sup>76,77</sup>

The 30–200  $\mu\text{m}$  crystal sizes obtained were ideal for proof-of-principle SOX measurements due to the comparable minimum beam size ( $5 \times 55 \mu\text{m}$ ) used at the 12–1 microfocus beamline at SSRL. Seeding can be used to control the nucleation rates to produce a high density of smaller 10–20  $\mu\text{m}$  microcrystals. Fig. 3B shows fully hydrated, randomly oriented tetragonal thaumatin crystals grown on chip using micro-batch crystallization (without and with seeding, top and bottom respectively) and stored for three days. Fig. S4† shows exemplary images of the other protein crystals measured

in this work. It is worth noting that the thin spacer film (25 or 48  $\mu\text{m}$ ) not only reduced the sample volume requirements (4–8  $\mu\text{L}$ ) but also restricted crystal growth in the Z-direction to span the “set” spacer thickness. This minimized scatter contribution from the crystallization solution surrounding the microcrystals. Preferential alignment of 200–300  $\mu\text{m}$  long plate-like crystals of concanavalin-A and NSP5 was observed microscopically. The concanavalin-A crystals appeared to grow to span the entire spacer thickness  $\sim 50 \mu\text{m}$  whereas individual NSP5 plates in the crystal clusters were  $\sim 10$ –15  $\mu\text{m}$  thick. Lysozyme and catalase crystals, which were cuboid in appearance, did not have any visible preferential alignment.

### X-ray scatter background measurements

To quantify and compare the X-ray scatter background contributions from different materials in the beam interaction cross section, background measurements of ambient air scattering (5 mm path length), chips with different COC films thicknesses, and buffer-filled chips (0.1 M Na acetate buffer, 1 M NaCl) with two different spacer thicknesses were carried out. Radial averages of the scattered intensity for air and buffer filled chips (Fig. 5) are dominated at low angles by air-scatter background. Contributions to background scattering from the enclosing amorphous COC films represent an increase of approximately 20 percent over ambient air scattering at the peak of the COC-associated scattering at approximately  $1.1 \text{ \AA}^{-1}$ , corresponding to 5.8  $\text{\AA}$  in real space (seen as a broad “halo”). This is in agreement with previous observations of increased scattering  $\sim 5$ –6  $\text{\AA}$  for different COC grades<sup>58,74</sup> and indicates there is some degree



**Fig. 5** Radial averages of scattering associated with crystal-free chips (or instrument-associated air scattering) to quantify background scattering contributions. Air (cyan, blue) and buffer-filled (red, yellow) chips with nominal 2.7  $\mu\text{m}$  (cyan) or 3.7  $\mu\text{m}$  (yellow, blue, red) COC enclosing films and 48  $\mu\text{m}$  (red) or 25  $\mu\text{m}$  (yellow) spacer layers are shown. Note: the data represented by the yellow curve, collected during a separate beamtime using a different batch of COC material from Polysciences Inc., shows a slightly different scatter profile at low  $q$ . Inset: comparison of scattering intensity from the 200  $\mu\text{m}$  COC support frame material and a 3.7  $\mu\text{m}$  enclosing film.

of systematic packing between adjacent polymer backbone strands.<sup>78</sup> COC also contributes a broad, featureless scattering signal at low resolution ( $>6$  Å). At higher angles, the enclosed buffer layer was the most significant background contribution with buffer filled chips exhibiting a “water ring” at  $1.8$  Å<sup>-1</sup> corresponding to  $3.4$  Å in real space. This contribution can be decreased by using a thinner spacer layer as demonstrated by the yellow curve in Fig. 5, corresponding to a chip with a  $25$  µm spacer layer. Because of the inherent flexibility of the enclosing thin COC films, some variability (10–20 percent) in the buffer or flow layer thickness is expected. This explains why the scattering signal (around  $1.8$  Å<sup>-1</sup>) for the  $25$  and  $48$  µm spacer samples does not scale linearly. Minor changes in peak intensities at  $\sim 0.3$  and  $\sim 1.1$  Å<sup>-1</sup> were also observed between the  $25$  and  $48$  µm spacer samples, that cannot be explained sufficiently by COC film thickness variation between different batches ( $<0.2$  µm). Since the two samples were characterized during different beamtimes using different batches of COC from Polysciences, Inc., we suspect minor batch to batch compositional variation in the supplied COC material and subsequent spin-coating solution preparation and processing steps (filtration, postbake) could influence the degree of short-range chain packing, resulting in slightly different amorphous scatter signal at low  $q$ . Regardless, both the enclosing film thickness and the flow spacer thickness are parameters that can be changed to control background scatter to match sample requirements. While a direct background comparison with other microfluidic chips from literature is made challenging due to differences in material attenuation properties, film and flow layer thicknesses, and beam characteristics, the inset in Fig. 5 shows the drastic reduction in background that results from using thin films over X-ray windows ( $\sim 3.7$  µm per side) vs. shooting through thick supports ( $\sim 200$  µm per side) that are on the order of film thicknesses sometimes used as capping layers in microfluidic chips.

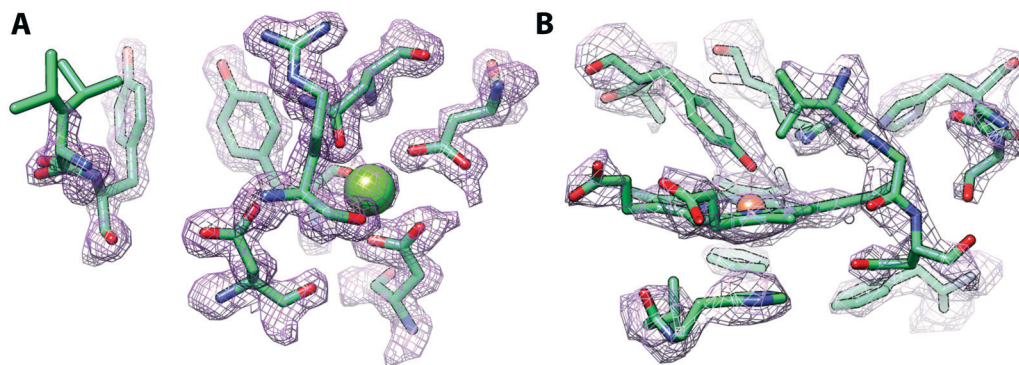
### *In situ* X-ray diffraction at room temperature

Fig. 3C shows the COC microfluidic chip mounted on a slotted magnetic pin base and attached to the goniometer for room temperature data collection at the 12–1 beamline at SSRL. The device's large measurable area and facile mounting allowed rapid collection of data in a semi-serial mode where  $30^\circ$  rotation wedges were collected on a series of manually centered crystals, eliminating the need for mounting of individual crystals. Individual wedges were collected in 15–20 seconds. The average data collection time for each chip was between 30 min and one hour, where the limiting step was manual centering of the crystals. Minimal settling or crystal movement was observed for most *in situ* grown crystals where the crystal size was well matched to the spacer layer thickness due to contact with the COC films. Some movement was observed when the crystal size was well below the spacer thickness ( $\sim 50$  µm spacer vs.  $10$ – $20$  µm lysozyme crystals).

Data from 5 to 16 individual crystals were merged to constitute complete data sets for the measured proteins. Data collection and refinement statistics for each protein are shown in Table 2. The lysozyme, thaumatin, and concanavalin-A crystals generated high resolution datasets, diffracting to  $1.5$  Å,  $1.45$  Å, and  $1.30$  Å respectively, with high multiplicity and excellent merging statistics including overall  $CC_{1/2}$  of 0.999, 0.999, and 0.993 respectively. The catalase dataset diffracted to a slightly lower resolution of  $2.27$  Å ( $CC_{1/2}$  0.997) but is comparable in quality to other room temperature structures reported for the protein.<sup>79</sup> Random crystal orientation for lysozyme, thaumatin and catalase was confirmed by a high degree of completeness, both in the total dataset and in the highest resolution shell. Concanavalin-A had only slightly less favorable completeness statistics because of its lower symmetry space group and apparent preferentially orientated crystal growth to span the spacer thickness (Fig. S4F†).

In cases where *in situ* growth limited crystal size to the full spacer thickness, background scattering around the water ring ( $1.8$  Å<sup>-1</sup>) was lower due to displacement of excess crystallization solution from the X-ray cross section (Fig. S5†). Moreover, when contact between crystals and the enclosing COC films was high, mother liquor background could also be reduced by physically removing excess buffer from the chip, leaving film-supported crystals surrounded by a minimal amount of mother liquor as observed in the case of catalase (Fig. S6†). The differences in the unit cell dimensions between the fully hydrated and excess solvent removed or partially “dehydrated” crystals were within the measurement error with  $\sim 0.1$  Å larger *a*- and *b*-unit cell dimensions for the fully hydrated crystals. The merging statistics for both these datasets is reported in Table S1.†

Plate-like or plate-cluster morphologies for the larger, thinner NSP5 crystals, however, proved more challenging as the chip geometry limited possible crystal orientations (Fig. S4C†). This made collection of an adequately redundant data set with sufficient completeness in all resolution shells difficult. Some of the crystals in the NSP5 dataset diffracted up to  $2.3$  Å which is comparable to other room temperature data sets for the protein that use similar crystallization conditions, PDB entries 6WQF (single crystal,  $2.3$  Å) and 7JVZ (SFX,  $2.5$  Å). But ultimately, a lower resolution cut-off of  $2.7$  Å was used for the final structure refinement due to constraints imposed by the crystal morphology and the low symmetry space group ( $C2$ ), although only one chip with this sample was measured. We expect that additional samples and tuning of on-chip crystal density and crystal size would improve these results. Overall, each of the solved structures were in excellent agreement with the PDB references used for molecular replacement except for a previously unreported carboxymethylation on the Cys376 residue observed in our catalase structure. The RMSD values ranged from  $0.121$ – $0.416$  Å, which includes minor movements of sidechains and loops and are shown in Table S2.† Representative  $2mF_o - DF_c$



**Fig. 6** Active sites of two structures determined from on-chip crystallization and synchrotron data collection. The  $2mF_o - DF_c$  density maps at  $1.5\sigma$  are shown in purple mesh. (A) The saccharide binding site and calcium coordination site of concanavalin A. (B) The heme active site of catalase bound to a proximal tyrosine residue.

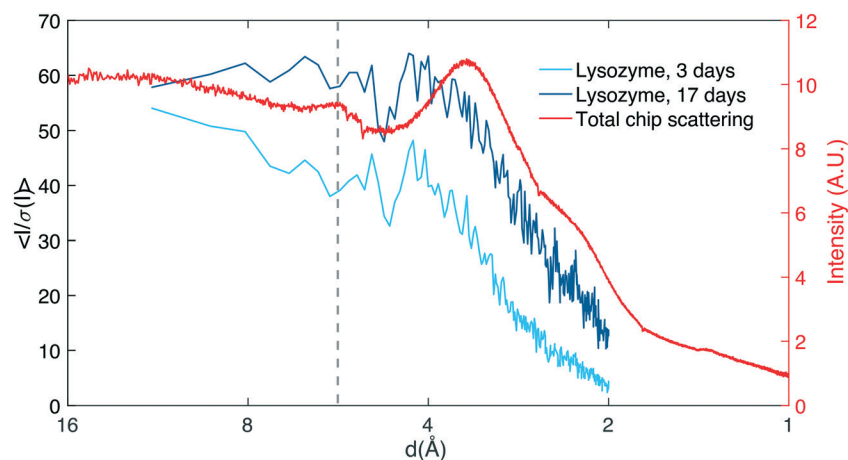
electron density maps of the concanavalin-A and catalase active sites are shown in Fig. 6.

To study the effects of long-term storage on-chip, data was also collected on lysozyme crystallized under similar conditions but stored in a humidified environment at 21 °C for 17 days ("aged"). These crystals showed minimal signs of dehydration based on unit cell shrinkage, and diffracted to similar resolutions, as much as 1.5 Å for the aged samples compared to 1.87 Å for freshly prepared crystals (stored for 3 days on-chip). This difference in resolution can be attributed to slight differences in crystal dimensions and normal variation in crystal quality, especially given that each of these datasets was obtained from multiple crystals which diffracted to differing resolutions. Thus, long term storage effects such as evaporative dehydration did not degrade crystal quality. The RMSD between the two structures was 0.094 Å, demonstrating that the two structures were identical. A detailed comparison of crystallographic statistics for the aged and fresh lysozyme sample is included in Table 2.

To assess the effect of both the polymer thin film and the mother liquor surrounding the crystals on the structures, we made a comparison between the  $I/\sigma(I)$  vs. resolution and the measured chip background (purely buffer-filled) vs. resolution for both fresh and aged lysozyme, as shown in Fig. 7. Both samples appear to have no features in  $I/\sigma(I)$  that correlate with background scattering and no deleterious effects of the COC scatter peak at 5.8 Å are apparent. While there is a small drop in  $I/\sigma(I)$  around the peak of COC scattering, in both cases it is within the level of noise in the data.

## Conclusion

In summary, this work demonstrates the development of robust, easy-to-use polymer microfluidic chips that can be used for routine and reliable room temperature diffraction measurements on fully hydrated protein microcrystals. Our large area, optically transparent chips are compatible with *in situ* crystallization, sample monitoring, crystal storage and



**Fig. 7** A comparison between the  $I/\sigma(I)$  for the two lysozyme structures solved in this work and the total scattering background from the chip. The fresh sample shown in light blue ( $\sim 30\ \mu\text{m}$  crystals) diffracted to 1.87 Å and the aged sample shown in dark blue ( $\sim 70\ \mu\text{m}$  crystals) diffracted to 1.5 Å confirming that long term storage of crystals on the chip does not affect crystal integrity. A small dip in signal to noise is observed around scattering peak for COC around 5.8 Å (dashed line). The larger dip around 5 Å corresponds to the location of a gap in the detector panel (masked out during analysis) and is confirmed by a decrease in the number of reflections in this resolution bin.

transport, and diffraction measurements directly on the chip, eliminating the need to handle small or fragile crystals. The chips are stable over several weeks in a humidified environment, and for several hours in ambient conditions, removing the stress of on-the-fly sample preparation. The modular device construction allows design flexibility to tune sample or flow layer thickness, enclosing film thickness, chip volume, or rastering area to match sample and experimental requirements. In this work, the chips were designed to be directly compatible with the standard goniometer setup at SSRL. High resolution structures (1.3–2.7 Å) for five different proteins (including one “non-model” protein, NSP5) were collected on chip using SOX. Looking forward, we envision these low-cost chips could be made available to users to allow crystallization and sample screening well ahead of beam-time, followed by streamlined plug-and-play experiments with minimal sample handling or mounting requirements. The platform can be expanded to add functionalities to study not just static structures but dynamics by leveraging microfluidic capabilities to allow ligand introduction, electric field application, temperature-jumps, or pump-probe time-resolved experiments.

## Outlook: XFEL measurements

We are currently expanding this work and are applying the design and fabrication principles to produce and demonstrate chips that are tailored for XFEL SFX applications. The diffraction before destruction principle of operation at XFEL sources demands fresh sample at every beam shot. Based on the crystal symmetry and the data analysis techniques used,<sup>80,81</sup> hundreds to tens of thousands of crystals must be delivered to the beam in random orientations. Since XFEL beamtime is limited and precious, our goal is to develop a “shelf-stable”, plug-and-play microfluidic chip for XFEL sample introduction that maintains crystal hydration for up to 24 hours without the need for external humidity control, while still contributing minimally to background. This would address some of the limitations of existing approaches like the need for on-the-fly sample preparation (open-format chips) and high background (microfluidic chips) to enable routine, high-resolution SFX with minimal downtime. Validation of the first XFEL chip prototypes in preliminary experiments is promising. The current design allows collection of  $\sim 10^4$  useful, low background shots per chip when continuously rastering through the thick and thin regions of the chip support with minimal alignment requirements due to the diffuse scattering of the amorphous target material. Our ongoing efforts focus on (1) maximizing the X-ray window area of the chip to increase the fraction of area useful for sample characterization (beyond current limit of 50% due to thick supports), (2) controlling crystal nucleation and crystal densities on the chip leveraging surface chemistry modifications, and (3) demonstrating the use of this platform for more non-model proteins of interest.

## Author contributions

Conceptualization, D. G., T. L. K., M. L. S., M. F., M. A. C.; sample preparation and crystallization, D. G., M. L. S., D. C. F. M.; polymer microfluidic chip design and preparation, D. G., T. L. K.; data evaluation and analysis, D. G., M. L. S., A. Y. L.; instrument operation for SOX experiments at SSRL, J. L. W., A. E. C., S. R.; writing, D. G., M. L. S., A. Y. L., D. C. F. M., T. L. K., M. A. C., M. F. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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