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Characterizing close-focus lenses for microendoscopy

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Abstract

Microendoscopes are commonly used in small lumens in the body, for which a focus near to the distal tip and ability to operate in an aqueous environment are paramount for navigation and disease detection. Commercially available distal optic systems below 1mm in diameter are severely limited, and custom micro lenses are generally very expensive. Gradient index of refraction (GRIN) singlets are available in small diameters but have limited optical performance adjustability. Three-dimensional (3D) printed monolithic optical systems are an emerging option that may be suitable for enabling high performance, close-focus imaging. In this manuscript, we compared the optical performance of three custom distal optic systems; a custom-pitch GRIN singlet, 3D-printed monolithic doublet, and 3D-printed monolithic triplet, with a nominal working distance (WD) of 1.5mm, 0.5mm and 0.4mm in 0.9% saline. These short WDs are ideal for microendoscopy in collapsed or flushed lumens such as pancreatic duct or fallopian tube. The GRIN singlet had performance limited only by the fiber bundle relay over 0.9mm to 1.6 mm depth of field (DOF). The 3D printed doublet was able to achieve a comparable DOF of 0.71mm, while the 3D printed triplet suffered the most limited DOF of 0.55mm. 3D printing enables flexible design of monolithic multi-element systems with aspheric surfaces of very short WDs and relative ease of integration.

Keywords

Endoscopy; Lens design; 3D printing; microendoscope; multi-modal imaging

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Disclosures

The authors declare that there are no conflicts of interest related to this article.

1 Introduction

Minimally invasive endoscopic imaging offers the opportunity for early detection of cancer and other disease states by bringing powerful optical techniques to the interior of the body. For example, in one study colonoscopy was associated with a 61% reduction in colon cancer mortality among Veterans.¹ However, the colon has a large lumen, accommodating endoscopes of around 12mm in diameter which typically include imaging lenses in the 2–3mm diameter range. The desire to extend minimally invasive clinical imaging to organs with smaller lumens, such as the pancreas², fallopian tube³, bronchioles⁴, or guiding needle biopsies of the lungs⁵ or during neurosurgery⁶, has necessitated the development of microendoscopes that are sub-mm in diameter. These endoscopes may contain one or more imaging modalities, and possibly other functionalities such as cell/tissue collection, laser therapy, or drug delivery. In these microendoscopes, optical channel diameters are limited to the 100–500 μ m diameter range. The small lumens themselves define a new set of challenges that drive specialized lens requirements. Typical endoscope optics for large cavities or lumens have moderate to large angular fields of view (AFOV), are designed for operation in an air or insufflation gas environment, and are focused at the hyperfocal distance, from a few millimeters to infinity. However, small lumens may be collapsed and filled with mucus, cilia, or plicae. Therefore, it is more likely that microendoscopes will need moderate AFOV, depth of field (DOF) from less than a millimeter to a few millimeters, and ability to operate in an environment where 0.9% saline is used for flushing and irrigation as an immersion objective. We are particularly interested in distal optic systems of 0.5mm or less in diameter that can operate in these cramped, aqueous environments.

There are two main detection options for microendoscopic imaging, a distal miniature sensor (recently dubbed the “chip-on-tip”), or a fiber image guide, which directs light to a remote, possibly high-sensitivity and relatively large, CMOS or CCD camera. The chip-on-tip technology powered by a new generation of micro-CMOS detectors is applicable to sub-mm diameter endoscopes, but these sensors are still too large for multi-modality or multifunctional microendoscopes. Fiber image guides with 1,000–10,000 cores are appropriate for situations where the imaging channel should be 500 μ m or less in diameter and/or where a highly sensitive camera is required.

With either of these options, corresponding 100–500 μ m diameter optical elements must be used to focus the image onto the fiber guide or miniature detector. Optical elements manufactured in a variety of ways can be used. Glass catalog optics with diameters smaller than 1mm are rare, but specialty optics companies can use conventional lens grinding procedures to create custom miniature optics. The advantage of this approach is that a large variety of materials can be used together with well-established optical design methods. For example, we previously demonstrated a custom 300 μ m diameter triplet distal optic system consisting of two plano-convex and a meniscus lens made from glasses and sapphire, with excellent optical performance.⁷ The disadvantages of this method are the high cost and challenging assembly of multi-element systems.

Gradient index of refraction (GRIN) rod lenses are readily available in 0.5mm or smaller diameters to be used as a singlet. Their ease of assembly, a cylindrical rod shape abutting

the end of a fiber bundle or miniature sensor, makes them an attractive choice for miniature endoscopes. Commercial off-the-shelf (COTS) GRIN lenses used as a singlet objective generally have too long of a working distance (WD) for a microendoscope, but custom pitch (length) GRIN singlets can be obtained at a relatively low cost, or a GRIN microendoscope system for a higher cost. However, GRIN singlet systems have limited optical parameters that can be varied (gradient parameter and pitch) and typically suffer from strong chromatic aberration as a lone element. In regards to microendoscope biocompatibility, care must be taken that the GRIN material and dopants are either non-toxic or the GRIN lens is coated/covered with a biocompatible material.⁸ Use of a GRIN singlet system with fiber bundles is common and our group has previously demonstrated the use of a 0.25mm diameter GRIN singlet with 3000 element fiber bundles for imaging of the fallopian tubes.^{9, 10} Optical performance limitations of a GRIN singlet has been mitigated through additional optical elements placed in proximity to, or manufactured directly on, the GRIN element. For example, an epoxy window was created with soft lithography¹¹ to correct a GRIN-fiber-based optical coherence tomography (OCT) probe, and a three dimensional (3D) printed correction surface for the proximal end of a GRIN-rod based two-photon microscopy system¹² has been demonstrated. Nanopatterned metasurfaces¹³ and 3D-printed off-axis freeform total internal reflection mirrors¹⁴ are further methods for creating miniature optical systems, and have been combined with fiber optics to create OCT fiber probes with high resolution over extended DOF. The flexibility afforded by 3D-printing is beginning to be more widely utilized in endoscope optics. For example, a monolithic optical design for OCT and fluorescence microendoscopy has been demonstrated.¹⁵ Relevant to *en face* imaging with fiber bundles or miniature sensors, we previously demonstrated a 0.5mm diameter 3D-printed singlet lens that showed large DOF and excellent surface quality,¹⁶ however bulk scattering of the resin material degraded optical performance.

Recent activity has focused on expanding 3D printed lens material beyond the resins typically used. Liquid silica resin (LSR) has recently been introduced and is an exciting option due to the potential for biocompatibility and higher ultraviolet/blue transmission where other typically used resins may absorb.^{17,18} However, it is important to note that the 3D printing fidelity of precision glass optics for imaging applications is still limited by its shrinkage and highly controlled surface accuracy during manufacturing. 3D printing allows for the design of monolithic optical systems, simplifying the alignment process of coupling to a fiber bundle or sensor, and an optical stop can be incorporated directly into the lens system. Most importantly, the proposed printing strategy significantly increases the flexibility of the design and fabrication of miniature aspheric optics, which are currently unachievable with conventional grinding and polishing processes.

We report the performance of three close-focus microendoscopic distal optical systems. Each system utilized the same aperture stop placed in front of the distal optical element, and the same fiber bundle, proximal optical system, and camera. First, a custom-pitch GRIN singlet was designed, modestly decreasing the usual several-mm WD to 1.5mm. Then, to effect very short WDs, two different monolithic 3D-printed glass optical systems, a doublet and triplet, were designed and fabricated.

2 Methods

2.1 Design Goals

For best performance in our microendoscopic application, the image must remain in focus for tissue very close to the endoscope (WD less than or equal to 2mm), the AFOV should be greater than or equal to 40°, and DOF should be greater than 1mm. Saline will be flushed to irrigate the lumen, clear the mucus, and/or displace cilia or plicae ahead of the endoscope, therefore the distal optical system's WD will be in 0.9% saline immersion. The overall system object space resolution requirement is driven by the ability to detect disease. For example, in the fallopian tubes, serous tubal intraepithelial carcinomas (STICs) may be as small as few hundred cells in size¹⁹. An object space resolution of 20μm may be sufficient to detect altered optical signatures of STICs and many other diseases. These goals are summarized in Table 1.

2.2 Distal Optic System Specifications

The specifications of the three distal optic systems utilized in this study are listed below in Table 2.

2.3 Optical Modeling

The three 0.5mm distal optic systems were modeled using OpticsStudio (Zemax); custom length GRIN singlet utilizing a glass catalog GRIN lens material (GTAGNEU, GRINTECH), and two different custom multi-element 3D printed lenses, a doublet and triplet. For the GRIN singlet, manufacturer's information about dispersion as a function of wavelength was imported into the material library of OpticsStudio and used to model the custom lenses. The GRIN lens rod length (pitch) was allowed to vary while OpticsStudio optimized for the smallest spot size with a WD of 1.5mm. A GRIN rod length of 1.28mm yielded the best results. For the 3D printed lens, the LSR was modeled to have a constant index of refraction of 1.44 at 550nm. The flexibility of the 3D printing process allowed for aspheric surfaces up to the 6th order term. As a result of the optimization process, two potential solutions were devised, a doublet with a nominal WD of 0.5mm, and a triplet with a nominal WD of 0.4mm. The simpler monolithic doublet contains a stop aperture mount and two bi-convex lenses with aspheric surfaces. The more complex monolithic triplet is designed to bring the WD even closer, and push the limits of the 3D printing process, as well as achieve excellent off-axis performance. It contains a stop aperture mount, and three lenses with aspheric surfaces, a bi-convex, meniscus, and plano-convex lens. The distal optic system layouts and modulation transfer function (MTF) plots of the three systems are shown in Figure 1. Our system utilizes a 10,000 element fiber bundle (PN: FIGH-10-350S, Fujikura) with a core-to-core spacing of 3.3μm, which serves as the limit of the intermediate image space resolution. This fiber bundle's limit is indicated as a vertical line in the MTF plots.

Based on the MTF, the theoretical best on-axis image space resolution of the GRIN singlet, 3D printed doublet and 3D printed triplet should be 3.015μm, 1.485μm, and 1.656μm respectively (with 10% contrast held as the threshold), much better than the fiber bundle's 3.3 μm resolution limit. To increase the DOF of all distal optic systems, the entrance pupil

diameter (EPD) was reduced to 250 μ m by use of an optical stop at the front of the first surface.

2.4 3D-printing process

The custom-length GRIN singlet was obtained from GRINTech. The 3D printed doublet and triplet were manufactured in the Precision Freeform Optics Design, Fabrication and Testing facility at the University of Arizona, via a two-photon polymerization (2PP) technique described previously.^{17,20} The technique uses a specially formulated, solvent-free, photosensitive, LSR that has been modified to reduce organic components that cause shrinkage during pyrolysis. Although the described process reduces shrinkage of the printed optic during pyrolysis, multiple trials were still required to determine the pre-pyrolysis dimensions that achieved the correct post-pyrolysis dimensions. It is critical to complete the pyrolysis process, as this heat treatment vaporizes the organic molecules that absorb shorter wavelengths, which cause a yellowish cast in the images. Figure 2 displays the transmittance of the pre-pyrolyzed and post-pyrolyzed material in the visible spectrum.

The LSR was polymerized by the 2PP system consisting of a 0.6 NA objective, 780nm fs laser, 1.4nJ pulse energy, and 75mm/s scanning speed. The 2PP setup is shown in Figure 3. The doublet and triplet systems were manufactured in 1.5 and 3 hours, respectively, the latter has greater structural complexity and number of elements. Figure 4 shows photographs of the two 3D printed distal optic systems mounted on the fiber bundle, as well as electron micrographs of the systems before experimentation.

2.5 Experimental Setup

To create the EPD of 250 μ m, the system stop, a blackened, stainless steel microwasher (Gateway Laser) with outer diameter 500 μ m and inner diameter 250 μ m, was glued to the front face of each lens, using UV-curing glue (PN: NOA68, Norland). The test bench setup consisted of resting the distal optic system in a stainless-steel V-groove holder and abutting the 10,000 element fiber bundle (PN: FIGH-10-350S, Fujikura) against the lens in the V-groove. The opposite end of the fiber bundle was positioned at the WD of a 20X microscope objective (PN: UAPON20XW340, Olympus), whose image was then magnified by a 75mm focal length achromat lens (PN: AC254-075-A, Thorlabs) onto a CMOS detector (PN: ON Semi PYTHON 1300BFS-U3-13Y3C-C, Teledyne FLIR), with auto-exposure activated. SpinView software (Spinnaker SDK, Teledyne FLIR) was used to display and save the images. For testing, a 3" x 1" positive variable line grating (R1L3S6P, Thorlabs), a 3" x 3" 1951 USAF Resolution Test Target (DA004, MaxLevy/II-VI), or a 100 μ m grid target (R1L3S3P, Thorlabs) was positioned at the WD of the distal optic system and translated by motorized linear actuator (T-NA08A25, Zaber). The test target was back-illuminated by a large, uniform LED source (CX Series, Advanced Illumination). Images were taken of different lp/mm resolution patterns at each object distance. These images were used to obtain the as-built WD, AFOV, DOF, distortion, and object space resolution of each test lens. Figure 5 illustrates the testing setup.

To obtain the as-built WD, the test target was moved to the position of highest contrast. To obtain the object size or field of view (FOV), images of the 26 lp/mm resolution pattern

were collected at the WD. The number of line pairs (lp) visible in each image was counted and divided by 26 to obtain the FOV in mm. With the knowledge of the as-built WD, the FOV was also translated into AFOV via geometric calculations.

To obtain DOF and object space resolution, the images were analyzed in ImageJ (National Institutes of Health, USA). The DOF was determined as the range of object distances over which the object, in this case the 26 lp/mm grating, remained resolvable to approximately 10% contrast. To calculate contrast, the average grayscale intensity of a sample of pixels in 3–4 fiber bundle cores imaging “white” (transparent) sections of the line grating was measured. Similarly, the average grayscale intensity of cores imaging “black” (opaque) locations was measured. These values were then used in the following contrast equation:

$$C = \frac{(I_{max} - I_{min})}{(I_{max} + I_{min})}$$

To determine the resolution of the distal optic system, the line grating was swapped out for the 1951 USAF Target. For this study, resolution was defined as the smallest group and element of the USAF Target that could be resolved with approximately 10% contrast, at the lens WD. According to MTF plots a, b, and c in Figure 1, the 10,000-element fiber bundle, rather than the distal optic system, is the limiting factor on overall system resolution, where each fiber core-to-core spacing in the bundle serves as the intermediate blur diameter criterion for the overall system. Therefore, the as-built theoretical best resolution due to the distinct cores of the fiber bundle was calculated by dividing the FOV(mm) by the number of cores across the diameter of the fiber bundle. This value describes the object size that corresponds to 1 fiber core. Any object below the as-built theoretical limit, will not be distinguishable through the fiber bundle system. Note that each “fiber pixel” at the proximal end of the fiber bundle was imaged on to the camera with sufficient camera pixels to ensure that detector resolution is not a limiting factor.

3 Results

The WD, FOV, AFOV, DOF, and resolution observed for each lens is collated in Table 3 below. The theoretical resolution limit due to the fiber bundle is also provided. Barrel distortion is evident in the images, of approximately 8.3%, 5.0%, and 3.5% at the edge of the field of view for the GRIN, 3D printed doublet, and 3D printed triplet respectively.

3.1 Working Distance and Full Field of View

The as-built WDs of the GRIN singlet, 3D printed doublet, and 3D printed triplet were 1.30mm, 0.81mm, and 0.55mm respectively. The FOV and AFOV of each distal optic system was 1.19mm, 1.00mm, and 0.731mm, or 49.3°, 63.4°, and 67.2°, respectively. Figure 6 shows images taken with each system, of the 26 lp/mm test pattern at the WD, to qualitatively illustrate the differences in FOV at WD, color rendition, and image contrast.

The FOV(mm) of the GRIN singlet is greater than the 3D-printed doublet or triplet systems, even though it has a smaller AFOV. This is unsurprising, as the object was held at the WD of each lens, of which the GRIN singlet has the largest. Figure 7 illustrates how a

lens with a smaller AFOV but larger WD can have a larger FOV than a lens with a larger AFOV. It is instructive to compare the object AFOV in degrees, as it is the more analogous measurement. The 3D doublet and triplet have similar AFOV (63.4° and 67.2° , respectively), with the GRIN singlet having the smallest AFOV at 49.3° .

3.2 Depth of Field

The GRIN singlet and 3D printed doublet featured almost identical DOF's of 0.7mm and 0.71mm respectively, with respective object distance ranges of 0.9mm-1.6mm and 0.5mm-1.21mm. The 3D printed triplet had a DOF of 0.55mm from 0.30mm-0.85mm.

3.3 Resolution

At the distal optic system's WD, it was possible to visualize, with 10% contrast or greater, group 5 element 5, group 5 element 6, and group 4 element 6, for the GRIN, 3D printed doublet, and 3D printed triplet lens, respectively. This corresponds to a spatial resolution of $9.84\mu\text{m}$, $8.77\mu\text{m}$, and $17.54\mu\text{m}$, respectively (Figure 8).

3.4 Performance in a microscope system

To qualitatively assess the performance of the distal optic systems without the resolution limitations of the fiber bundle, a 10X, 0.25 NA microscope was used to observe the intermediate image formed by the GRIN singlet and 3D printed doublet and triplet. The collected images of a histology section of fallopian tube (formalin fixed, paraffin embedded, $6\mu\text{m}$ thick section, stained with hematoxylin and eosin) are shown in Figure 9. The large differences in FOV are evident, although visualization of plicae is possible with all.

4 Discussion

This study characterized the properties of three 0.50mm diameter distal optic systems for close-focus, microendoscopic use. Table 3 compares the performance of the three distal optic systems against the goals set. All distal optic systems exceeded WD and AFOV expectations. However, none of the distal optic systems were able to achieve a DOF greater than 1mm. It is promising that the 3D printed doublet's DOF (0.71mm) was almost identical to the DOF performance of the GRIN singlet (0.7mm). Typically, the extremely close focus (0.81mm and 0.55mm) of the two 3D printed systems compared to the GRIN singlet (1.30mm) would cause a greater tradeoff in DOF, as seen in the 3D printed triplet. This would be expected, since the EPD of all lenses was the same (0.250mm), the object space numerical aperture is much higher for the 3D printed lenses, typically leading to a concurrent decrease in DOF. All lenses had a measured resolution poorer than the highest predicted by the MTF at 10% contrast. However, the GRIN singlet and 3D doublet had resolved test patterns close to that predicted based on the limit imposed by the fiber bundle. The slightly better measured, compared to as-built theoretical, resolution of the GRIN singlet and 3D printed doublet may be due to small measurement error or a fiber bundle with core-core spacing smaller than that specified by the manufacturer. The 3D printed triplet had poorer than expected resolution performance. It is possible that this very complex lens suffered from a final manufactured shape that deviated from design. Some core-to-core color variations are seen in all images. If desired, the transmission intensity and color variations

between fiber cores could be corrected by calibration using an image of a uniformly bright white target (a “flat” image)²¹.

CMOS detectors are now available with sub-micron size pixels. As overall sensor sizes shrink and number of pixels increases, miniature sensors should become more prevalent in microendoscopes. The resolution of these future optical systems may no longer be limited by a fiber element or pixel size; the limiting element will be the distal optic system. The wide availability and simplicity of the GRIN singlets makes it a current favorite for microendoscopic use^{22,23}. However, GRIN singlets lack the versatility of 3D printed monolithic optical systems, which can be designed for extremely short WDs and to mitigate aberrations. 3D-printed optical systems also have greater potential for multimodality use, with, e.g. multi focal capability¹⁵. The ability to integrate mounting surfaces in a monolithic design, including fiber bundle and aperture stop mounts, is a unique capability that can greatly ease assembly. Ongoing research in glass printing techniques is leading to more accurate/predictable shapes after processing. As a result, submillimeter endoscopes with a close focus will become feasible for imaging small and collapsed lumens such as the Eustachian tube, pancreatic ducts, or tear ducts. Overall, this study provides further evidence that 3D-printed lenses are a promising option for extending the capabilities of microendoscopes.

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Biography

Dominique Galvez is an Optical Sciences PhD student at the University of Arizona researching microendoscopes, 3D-printed optics, endoscopic OCT, multi-modal imaging, and endoscopic cell collection. She holds a BS in Optical Sciences and Engineering from the University of Arizona.

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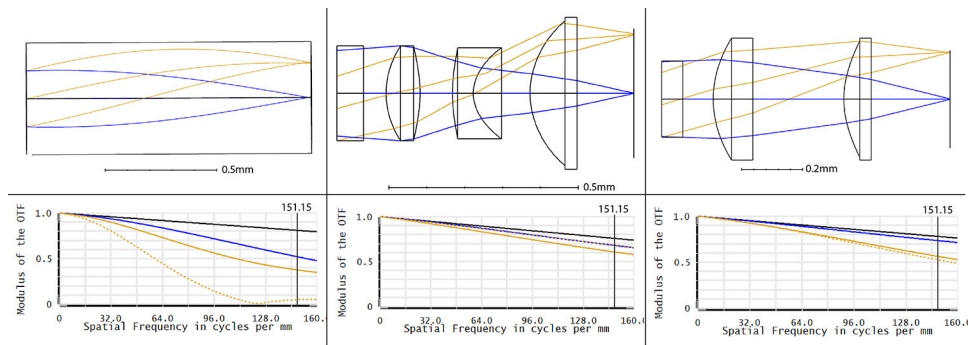


Fig 1. Distal optical layout and modulation transfer functions for (a) GRIN Singlet, (b) 3D printed Doublet, and (c) 3D printed Triplet. The blue lines represent the on-axis field, and the golden rays represent the off-axis field (sagittal rays are represented by the dotted lines, tangential rays by the solid lines). The vertical line on the MTF plots signifies the frequency cutoff imposed by the fiber bundle's core-to-core spacing ($3.3\mu\text{m}$) converted to line pairs(lp) per mm (151.2lp/mm).

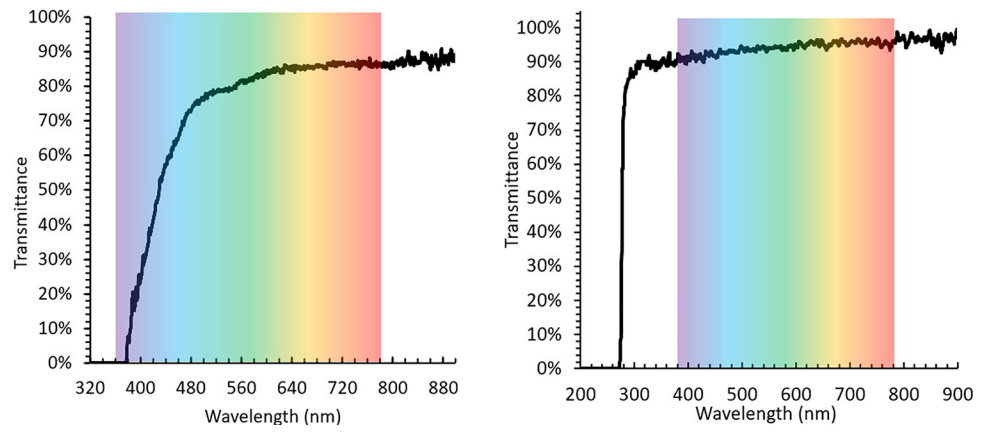


Fig 2. Transmittance of (left) the pre-pyrolysis LSR and (right) post-pyrolysis LSR in the visible spectrum

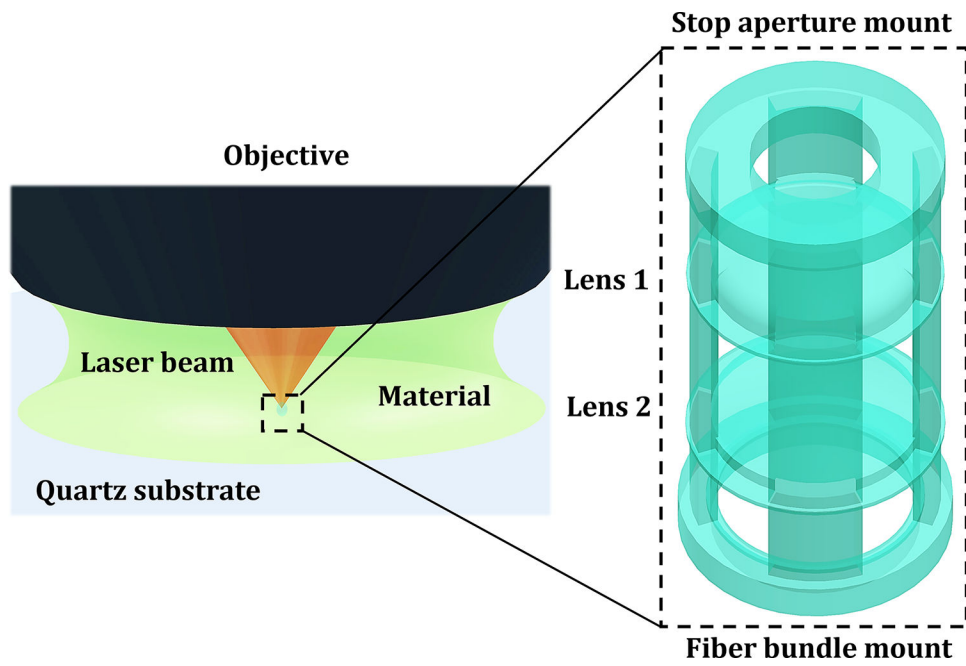


Fig 3.
Schematic of the 2PP-enabled 3D-printing setup and printing process

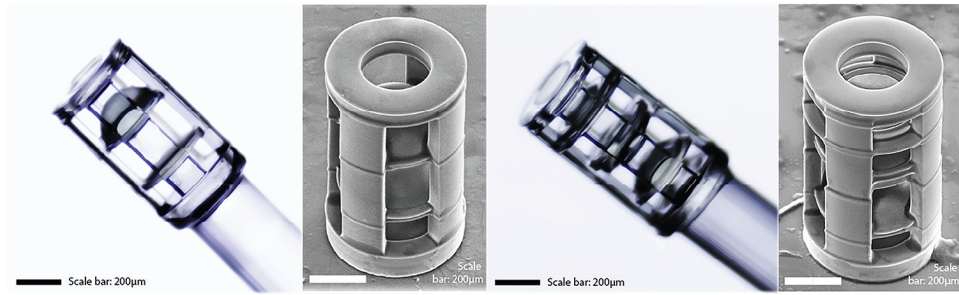


Fig 4. Photographs and electron micrographs of 3D printed doublet (a, b, respectively) and triplet (c, d, respectively). Scale bar is 200µm.

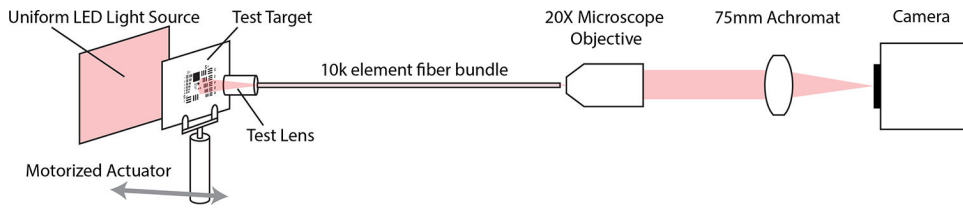


Fig 5.
Diagram of Test Setup

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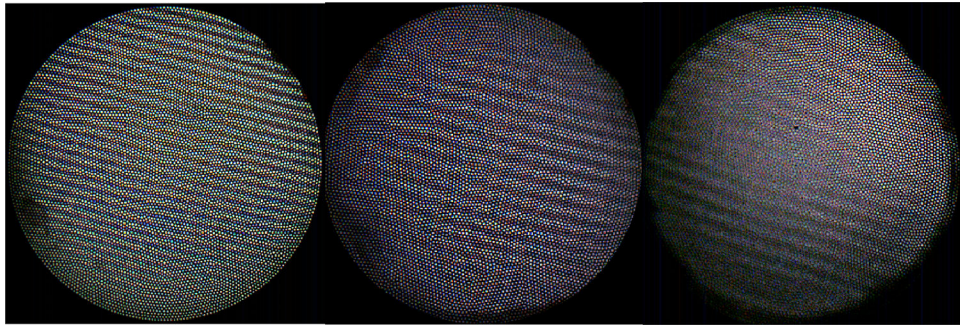


Fig 6. Images of the 26 lp/mm resolution pattern taken at the lens-specific WD for the a) GRIN singlet, b) 3D doublet, and c) 3D triplet.

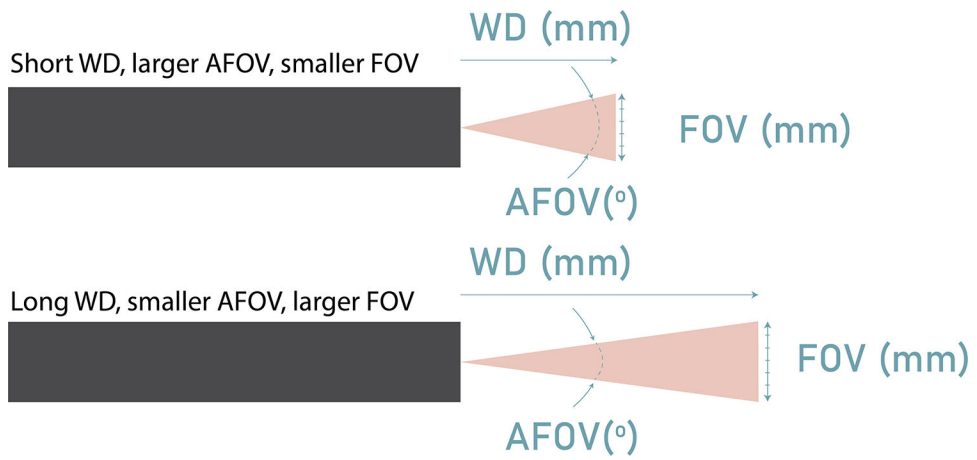


Fig 7. Diagram depicting how a lens with a smaller AFOV but larger WD can result in a larger FOV than a lens with a larger AFOV and shorter WD.

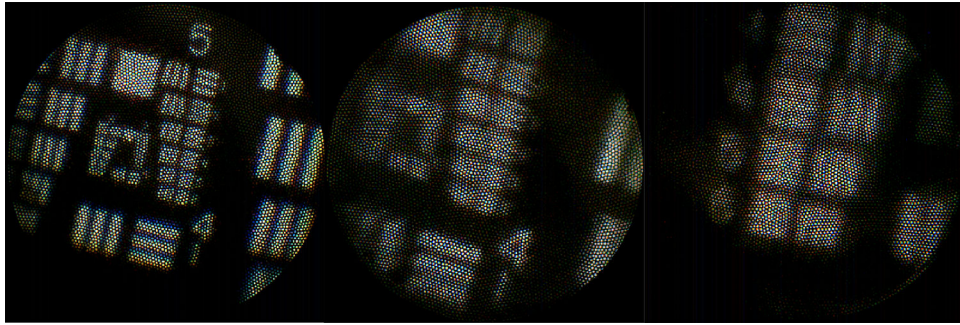


Fig 8. Images of the USAF Target taken at the corresponding WD's for the a) GRIN singlet showing Group 5, b) 3D printed doublet showing Group 5, and c) 3D-printed triplet showing Group 4 Elements 4, 5, and 6.

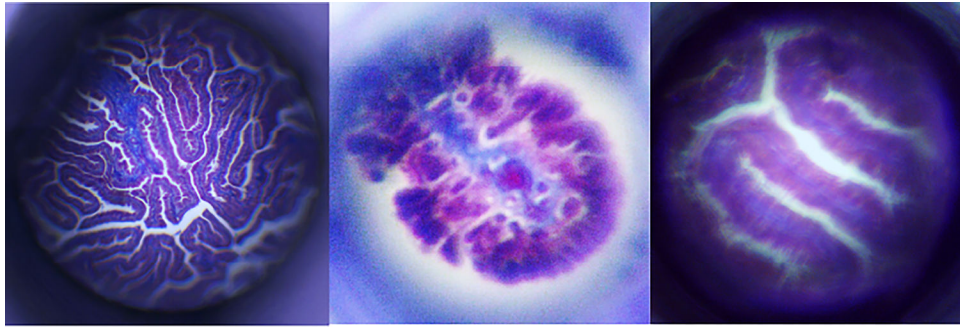


Fig 9.
Images of a histological slide of human fallopian tissue taken in a microscope setup with the
a) GRIN singlet b) 3D printed doublet and c) 3D printed triplet.

Table 1

Design goals for close-focus microendoscopic system

Optical Specification	Goal
WD	< 2mm in 0.9% saline
AFOV	> 40 degrees
DOF	> 1mm
Resolution	20 μ m

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Table 2

Distal Optic System Designed Specifications

Distal Optic System	Diameter (μm)	WD (mm)	Material
Custom GRIN Singlet	500	1.5	Non-toxic silver-based glass
3D Printed Doublet	500	0.5	<u>Pure Silica Glass</u>
3D Printed Triplet	500	0.4	<u>Pure Silica Glass</u>

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Table 3

As-Built Result Comparison

Distal Optic System	WD (mm)	FOV (mm)	AFOV (°)	DOF (mm)	Measured Resolution (μm)	Theoretical Resolution Limit (μm)
Goal	2		> 40	> 1.0	< 20	
Custom GRIN Singlet	1.3	1.19	49.3	0.7	9.84	12.11
3D Printed Doublet	0.81	1	63.4	0.71	8.77	10.15
3D Printed Triplet	0.55	0.731	67.2	0.55	17.54	7.42

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