A Portable, Optical Scanning System for Large Field of View, High Resolution Imaging of Biological Specimens †

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Abstract: Large field-of-view (FOV), high resolution imaging of biological specimens is a challenging task, requiring sophisticated and bulky optical systems. Such systems cannot be used for diagnosing or monitoring a disease at the point-of-care. To address this need, we developed a portable, optical system that can image—with a 2.88 μm resolution—large areas (6 mm × 40 mm) from various biological samples by performing scanning in one direction. This is achieved through the use of a microfabricated, mini-lens array. We demonstrated that our system can detect single cells from a smear blood test and thus validating our vision for its use at the point-of-care.

Keywords: point-of-care; microscopy; large field of view; optical imaging; lens array

1. Introduction

Examination of biological specimens is a crucial step for diagnosing and monitoring a broad range of diseases, extending from AIDS, tuberculosis and to sickle cell disease. Optical microscopy is widely used but it typically requires bulky, sophisticated and expensive imaging setups operated by trained personnel. In limited-resource settings in particular, a portable, easy to use system, that can automate the imaging process and obtain large field of view (FOV), high resolution images of the specimen, is needed [1]. Despite the numerous methodologies that have been proposed, implementing optical microscopy at the point-of-care still remains a challenge. In this work, we describe an imaging system that produces a 2 dimensional (2D) high resolution, large FOV image of a biological sample by performing scanning in only 1 direction.

2. Materials and Methods

We developed an imaging, scanning system (Figure 1) that comprises of: (a) an optical head that images the specimen of interest and (b) a motorized, translation stage that is responsible for moving the optical head along the specimen (1D scanning). The optical head is an assembly of a 5Mpixel CMOS sensor (Imaging Source, DMM72BUC02-ML) and of a custom-made mini-lens array. The optical head is attached to a translation stage that is connected to a computer-controlled, step motor through a lead screw. The specimen—typically sitting on a glass slide or coverslip—is placed at a short distance (~0–500 μm) from the optical head and is illuminated by an external white light LED array (Edmund Optics, #66-830) that provides homogeneous illumination. The CMOS/mini-lens array assembly moves in one direction and has a travelling distance of approximately 40 mm. This configuration creates a 40 mm × 6 mm image of the specimen. The distances between the mini-lens array, the CMOS sensor and the specimen are controlled by two step motors (20 μm step, 3 mm range).
that allows us to obtain a sharp, focused image of the specimen. The focusing procedure is performed manually prior to every scan.

**Figure 1.** (A) Schematic of the architecture of the integrated, portable scanning system and cross sectional view of the optical head; (B) photograph of the prototype and close-up view of the optical head.

Key element of the scanning system is the design of the 2D mini-lens array (Figure 2). It consists of 42, high numerical aperture (NA~0.7), 1 mm in diameter sapphire ball lenses (Edmund Optics, #43-638). They are glued on top of an array of 42 wells, DRIE-etched in a silicon chip [2] with a UV curable optical adhesive (Norland 60). The mini-lens array sits on top of the CMOS camera. All mini-lenses are equally spaced with an edge-to-edge distance of 50 μm. The lens array is placed with a tilt (17°) with respect to the scanning direction. The tilt design enables the FOV of a lens—estimated approx. 290 μm/lens—to overlap with the FOV of the neighboring one by ~20% during scanning. Having an overlapping FOV between neighboring lenses ensures that blind areas between them are also imaged.

**Figure 2.** (A) Top view schematic of the mini-lens array. The black, dotted outline indicates the effecting imaging area of the CMOS sensor, the green rectangular represents the silicon chip; (B) The tilt (17°) of the mini-lens array and the overlap between the FOV of neighboring lenses are crucial for covering the blind areas between the lenses; (C) Estimation of the FOV of a lens (~290 μm) based on a grid of lines imaged with the CMOS sensor in 1× magnification. The FOV is the diameter where the light intensity is 90% of its maximum value.
In order to quantify the optical performance of the scanner, we conducted two experiments: (a) we measured the uniformity in the optical resolution and contrast across the mini-lens array and (b) we measured the in-focus factor for each lens as they are not located in the same plane within the array. The in-focus factor represents the ability of each lens to focus the image to the CMOS sensor, when the mini-lens array is positioned at the plane that produces the best focused image. It has its maximum value at the best focused image plane. For quantifying the resolution and contrast, line pairs from the USAF 1951 resolution chart pattern (group 7) were imaged in 1× magnification. The contrast (C), obtained from greyscale images from each lens, was based on the Michelson formula [3]:

\[
C = \frac{\text{standard deviation of the light intensity gradient}}{\text{mean of light intensity gradient}},
\]

For quantifying the defocusing error, a series of images from a whole blood smear test were obtained by varying the mini-lens array to CMOS distance from 500 up to 1500 μm with a step of 20 μm. An image processing algorithm for the automatic detection of the best focused image was used [4,5].

3. Results

3.1. Optical System Characterization

The mean, optical resolution in the array, obtained by imaging the USAF 1951 resolution chart pattern, was 2.88 μm with a standard deviation of 0.5 μm (Figure 3A). 21 mini-lenses were selected for this experiment (out of 42), as those lenses captured images that fitted entirely into the CMOS effective detection area. The 0.5 μm standard deviation of the resolution, attributed to imperfections during the microfabrication process of the silicon chip and the assembly process, did not affect the image quality of the array as all lenses produce adequately focused images. The in-focus factor for each lens was measured using a smear blood test (Figure 3B). The best focused image produced by the array was selected as the one that the average in-focus factor (averaged from all lenses) was minimum. At the best focused image, defocusing error within the array was less than 8% while the contrast was within 30% of its maximum value.

![Figure 3.](image)

**Figure 3.** (A) Contrast and resolution distribution within the mini-lens array using a 1951 USAF resolution chart mask; (B) Normalized in-focus factor for each lens within the array, as measured by images obtained from a blood smear test. 5 lenses were excluded from this experiment as they did not produce high quality images.

3.2. Cell Detection

We performed single cell detection from a smear blood test (Figure 4) using an automated cell detection algorithm, specifically designed for bright field images [6]. Barrel distortion and vignetting effect corrections were performed prior to counting the cells. Dust particles—defined as particles with at least half the red blood cells size—were considered as background and were subtracted to provide enhanced contrast of the cell membranes with respect to the background. Hough transformation process was selected for image segmentation and for detecting single cells.
4. Discussion

We developed a portable, optical, scanning system that can image a large area (6 mm × 40 mm) with approximately 2.88 μm optical resolution in 1× magnification. The system operates in white light transmission mode and completes a full scan in a few minutes. We validated our system by performing single cell count from a smear blood test in an automated fashion. We envision our imaging system to be used for imaging biological specimens for the diagnosis and monitoring of a wide variety of diseases and conditions at the point-of-care.

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References


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