

Single Objective Light Sheet Microscope (OPM)

Oblique Plane Microscope (OPM) Acknowledging the Arnold & Mabel Beckman Foundation for their support in financing the equipment for this instrument.

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Specifications and primary uses of the OPM

Single Objective Light Sheet Microscope

The **Single Objective Light Sheet Microscope** is also referred to as **Oblique Plane Microscope** or **OPM**. It is a custom design Light Sheet microscope that enables high resolution and can be used with standard coverslipped samples. There is virtually no sacrifice in numerical aperture (NA) or loss in resolution - unlike most light sheet microscopes.

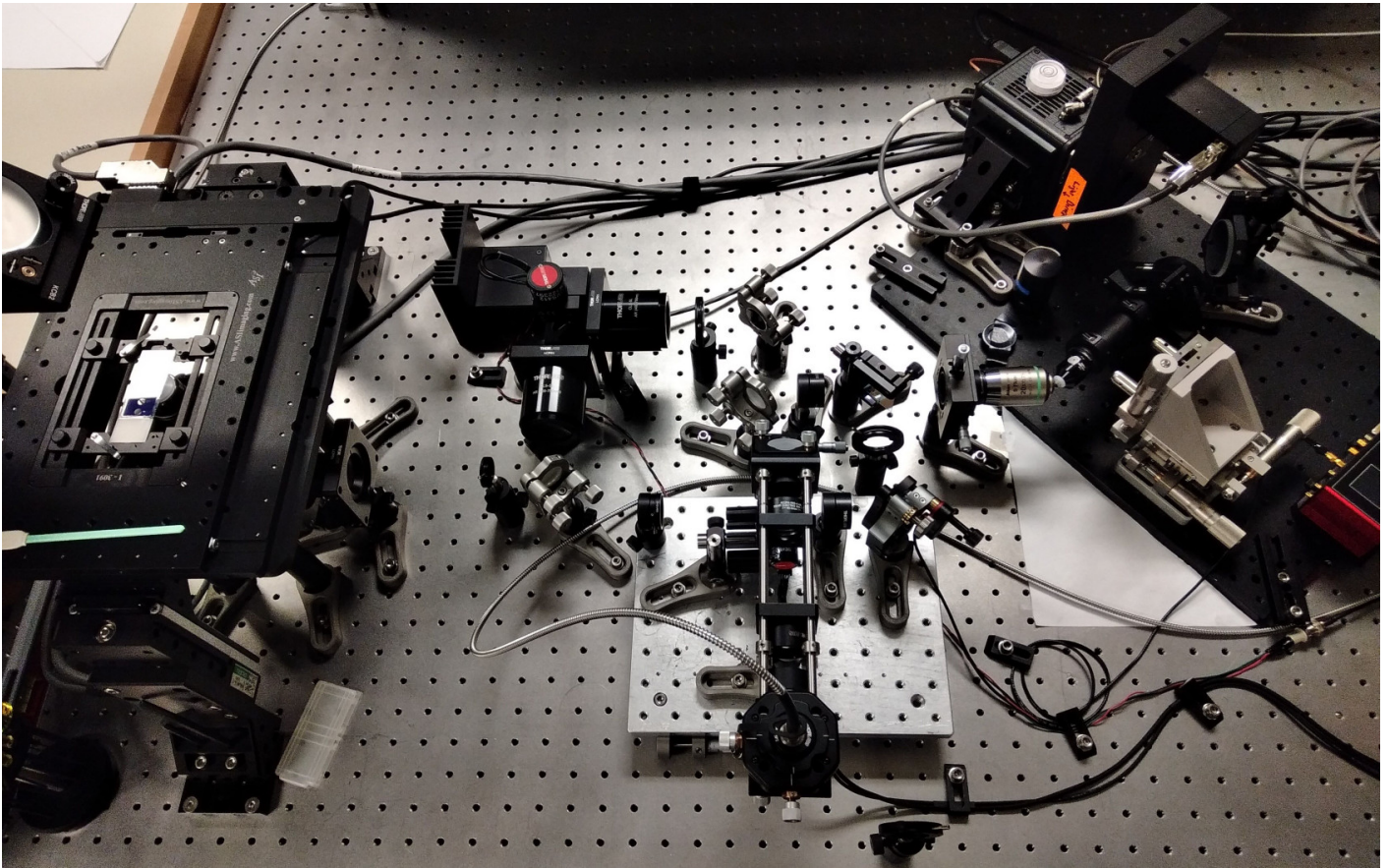
The OPM therefore presents a strict improvement over standard confocal microscopy. It has the sectioning capability and reduced photo-toxicity of light sheet microscopes while preserving image quality.

The animation below illustrates how it works.

[High_NA_single-objective_light-sheet_scan.mp4](#)

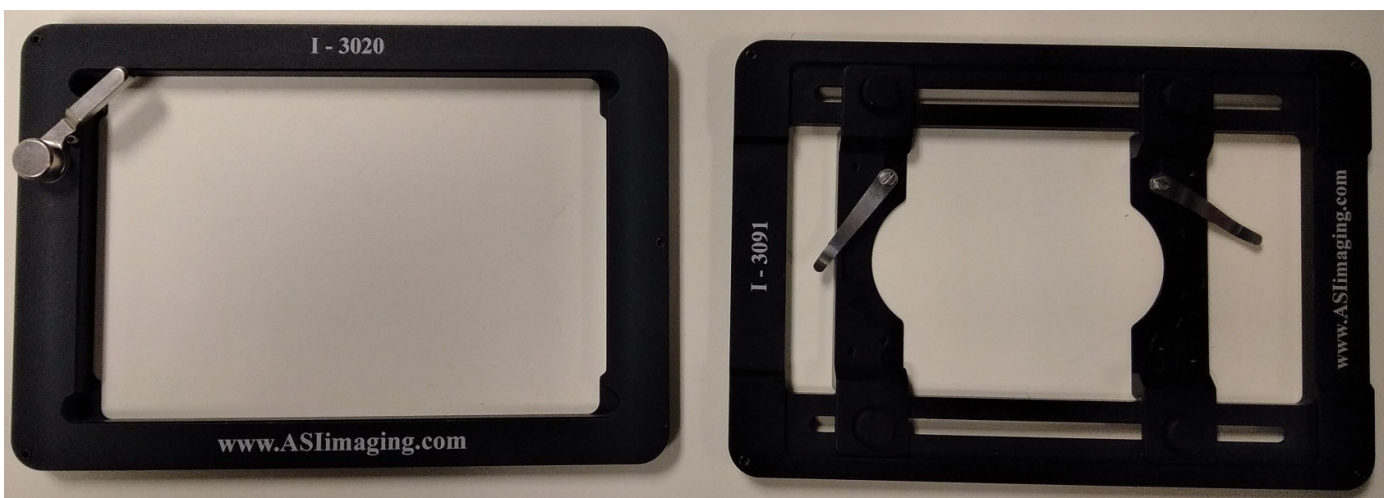
For more technical information on the OPM design, check [this open source paper](#).

The Beckman Center Single Objective Light Sheet Microscope



- Primary objective: Olympus 20X 1.0NA water dipping
- Secondary objective: Nikon 20X 0.75NA
- Tertiary objective: AMS-AGY V1.0 "Snouty"
- Hamamatsu Fusion sCMOS camera
- ASI FTP2000 XYZ stage
- 4 Channels: 405, 488, 561, 640

The sample stage has several sample holders for slides, round dishes and multi-well plates.



Primary uses

The size range of the samples is roughly from **single cells** to the bottom layer of **small drosophila** or **zebra fish embryos** and **small organoids**. Typically the OPM is used for **1 to 3 cells deep into the sample**.

While this limits the variety of sample that can be imaged on the OPM, the flexibility of sample preparation is a considerable advantage. The fact that it uses the same dishes as other standard commercial microscope allows for straightforward comparison of images between a confocal microscope and the single objective light sheet microscope.

Examples

Some imaging examples using the OPM technology by The Dean Lab at UT South-Western, Dallas Texas:

- **Biological imaging of clathrin-mediated endocytosis, vimentin and membrane dynamics**
- **Natural killer cell mediated cytotoxicity**
- **Imaging in biological microchannels - microtubules and nuclear shielding**
- **High-speed imaging of calcium transduction and cytoplasmic flows**
- **Simultaneous volumetric imaging and optogenetic stimulation**
- **Large field of view imaging of cortical neurons and ventral furrow formation**
- **Tissue-scale imaging**

For further details and reference, the original article "**A versatile oblique plane microscope for large-scale and high-resolution imaging of subcellular dynamics**" can be found [here](#).

Below is an example of what the OPM can do for tissue scale imaging:

"In addition to the rapid laser scan/descan illumination geometry, OPM is also compatible with a sample scanning acquisition format that is essentially field of view unlimited. Indeed, by combining scan optimized equipment with fully automated fluidic handling, it is possible to image $\sim 1 \text{ cm}^2$ of a thin tissue in less than 45 min per color and perform biochemistry, such as sequential multiplexed labeling. To demonstrate this, we imaged an entire 30-micron thick slice of coronal mouse brain tissue ([Figure 9A](#), [Video 15](#)) labeled with the nuclear marker DAPI. Within these data, even small features like nucleoli are clearly resolved from both lateral and axial viewing perspectives throughout the entire $\sim 6 \times 8 \text{ mm}$ tissue slice ([Figure 9B and C](#)). Likewise, we also imaged a $\sim 4 \times 14 \text{ mm}$ slice of 12-micron thick human lung tissue labeled for nuclei, angiotension-converting enzyme 2 (*ACE2*) mRNA, and surfactant protein C (*SFTPC*) protein ([Figure 9D](#)). Here, characteristic histological features, including bronchiole, alveoli and vasculature, are readily visible, albeit with molecular contrast and sub-cellular resolution ([Figure 9E,F and G](#), and [Video 16](#)). Quantification of molecular expression within this tissue section provides spatial information on $\sim 20,000$ cells, and

verifies our previous limited quantification of *ACE2* expression in alveolar epithelial type II cells using confocal microscopy ([Muus and Luecken, 2020](#)). Indeed, because we were not sterically restricted by the orthogonal illumination and detection geometry ([Figure 9—figure supplement 1](#)), the lateral dimensions of this human lung specimen were 8- and 1.5-fold larger than those of the biggest sample imaged with lattice light-sheet microscopy ([Gao et al., 2019](#)). However, in the third dimension, lattice light-sheet microscopy has in principle a 6.7x larger reach (2 mm working distance of the typically employed NA 1.1/25X detection objective compared to 300 microns working distance of our primary objective). In practice, optical aberrations limit high-resolution light-sheet microscopy to depths of a few hundreds of microns, even for highly transparent samples. Furthermore, our approach is fully compatible with automated fluid exchange, which is increasingly important for projects like the Human Cell Atlas that necessitate iterative imaging approaches for spatial *-omics* of RNAs and proteins at the single-cell level throughout entire tissues ([Chen et al., 2015](#))."

Using the OPM

The Single Objective Light Sheet Microscope is a custom-designed microscope and is a **VERY** sensitive instrument. It is very easy to bump components out of alignment.

For this reason, you will not be allowed to use it in autonomy. There are no trainings provided at the moment, all your experiments will require the presence and supervision of the Beckman Center staff.

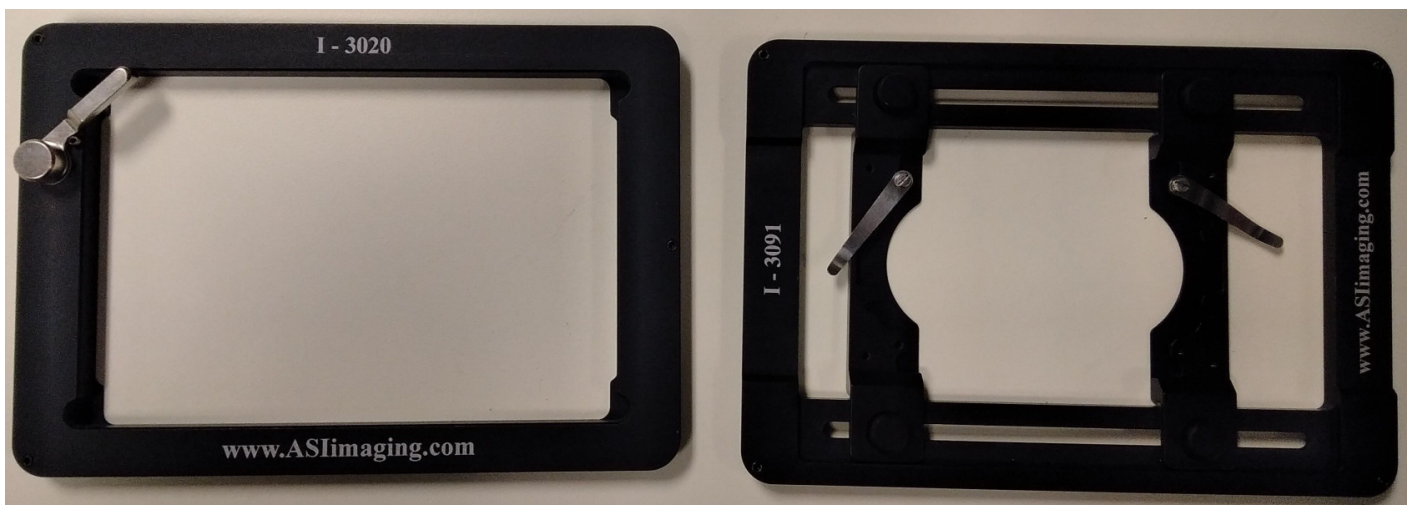
Your path to imaging your samples with the OPM

- Email the ticketing system at biof-imaging@colorado.edu and request a consultation with the Light Sheet Specialist.
- Book an appointment using the Microsoft booking link you will receive in the email from the ticketing system.
- Come prepared to the consultation with a clear idea of what your imaging needs are.
- You will then schedule a time on the instrument with the Beckman Center staff for a general overview of the instrument, the software and the Data generated. You will do some preliminary imaging then if you have your sample ready. This session will be similar to a classical training except you will only watch.
- Immediately after, or at another time, you will learn to process and visualize the data you acquired.
- Later on, always inquire with the Beckman Center staff about availability for scheduling your next imaging session.

Sample preparation for the OPM

Good news: Traditional sample preparation!

The OPM sample holders are similar to that of other commercial microscopes. Slides, round dishes and multi-well plates can be used. There is nothing different for your sample preparation so far.



Setting up an experiment on the OPM

This page is a guide to set-up an experiment with the OPM.

All experiments on the OPM will be done under the supervision of the Beckman staff and you will get help with processing the data to make it ready to analyze.

Turning the system on

Inserting the sample on the microscope

Turning Navigate on

Finding the sample and an area of interest

Setting up a basic experiment

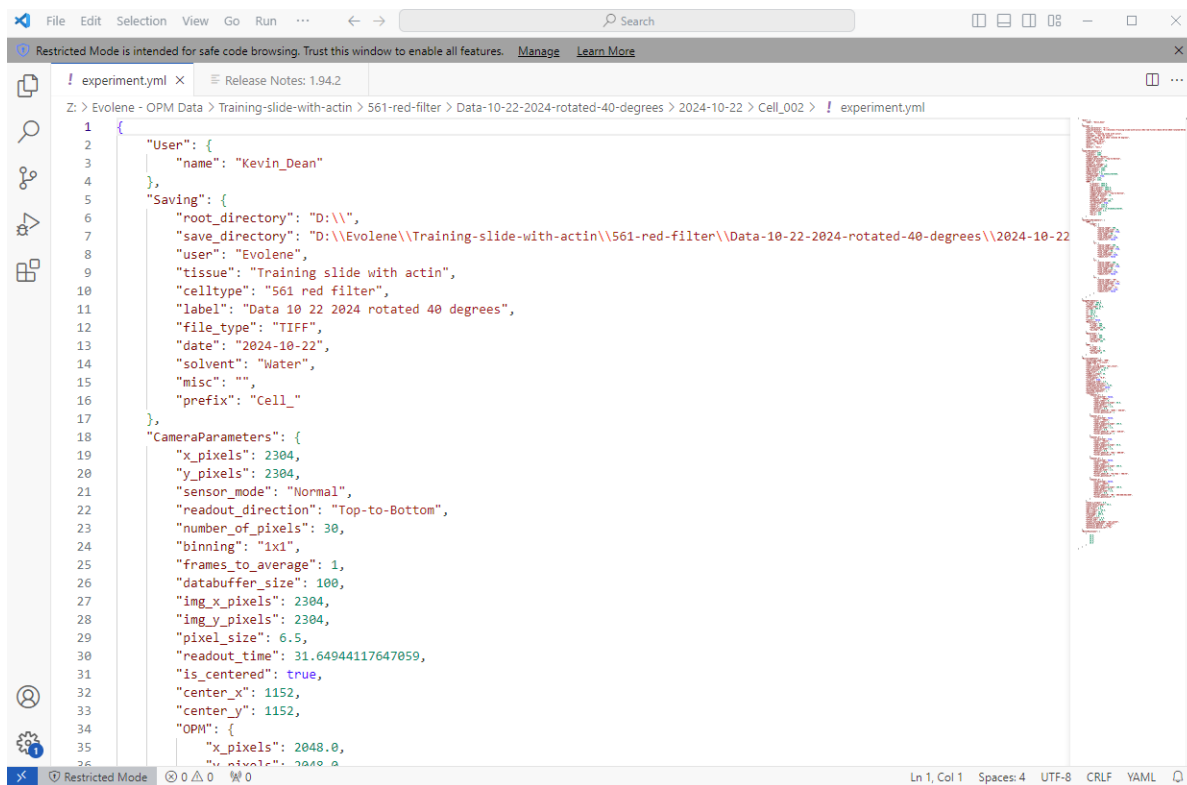
How is the Data saved

OPM Data processing in Fiji

OPM Data

The data generated by the OPM is in TIFF format. Whether you acquire a single image or a z-stack, you will get a single TIFF file containing all the images.

The software also saves an "experiment" file in Yaml format that contains all the experiment settings as well as a "waveform_constants" file in Yaml format that contains the waveform settings. These files can be opened by different applications such as notepad, etc. If you have Visual Studio on your computer, it is a great application to open the file in a structured way. See example below.

A screenshot of the Visual Studio Code editor interface. The main window displays a file named 'experiment.yaml' with the following content:

```
1 {
2   "User": {
3     "name": "Kevin_Dean"
4   },
5   "Saving": {
6     "root_directory": "D:\\",
7     "save_directory": "D:\\Evolene\\Training-slide-with-actin\\561-red-filter\\Data-10-22-2024-rotated-40-degrees\\2024-10-22",
8     "user": "Evolene",
9     "tissue": "Training slide with actin",
10    "celltype": "561 red filter",
11    "label": "Data 10 22 2024 rotated 40 degrees",
12    "file_type": "TIFF",
13    "date": "2024-10-22",
14    "solvent": "Water",
15    "misc": "",
16    "prefix": "Cell_"
17  },
18  "CameraParameters": {
19    "x_pixels": 2304,
20    "y_pixels": 2304,
21    "sensor_mode": "Normal",
22    "readout_direction": "Top-to-Bottom",
23    "number_of_pixels": 30,
24    "binning": "1x1",
25    "frames_to_average": 1,
26    "databuffer_size": 100,
27    "img_x_pixels": 2304,
28    "img_y_pixels": 2304,
29    "pixel_size": 6.5,
30    "readout_time": 31.64944117647059,
31    "is_centered": true,
32    "center_x": 1152,
33    "center_y": 1152,
34    "OPM": {
35      "x_pixels": 2048.0,
36      "y_pixels": 2048.0
37    }
38  }
39 }
```

The editor's status bar at the bottom indicates 'Ln 1, Col 1', 'Spaces: 4', 'UTF-8', 'CRLF', and 'YAML'. The left sidebar shows the Explorer view with a file tree containing 'experiment.yaml' and 'Release Notes: 1.94.2'.

Data processing with Fiji/ImageJ

Of course, you are welcome to use any software/method you like to process, analyze and render your data. The TIFF file format should make it easy to open by most softwares.

However, until further development of the instrument, the acquired data is skewed due to the angle of the beam scanning to capture the z-stack. Before visualizing it, you will need to perform a shearing operation to "de-skew" the data.

THE BECKMAN CENTER STAFF WILL DO THIS FOR YOU!!

But if you are curious about how to do it yourself, here are the steps:

There is a way to do this using Fiji/ImageJ. It requires a **GPU** and **CLIJ** installed on ImageJ.

- **How to install CLIJ on ImageJ:**

- Follow the steps indicated on this [github page](#). A Wiki page about CLIJ will be available soon as well.

- **Shearing the data:**

- Drag and drop the Fiji macros code in Fiji (provided to you by Beckman Center staff).
- Go in your experiment folder and copy the TIFF file of your z-stack into an empty separate folder (call it "Raw Data" to avoid confusion)
- On the code, update the angle values if different than 40 degrees, and the z step size. The xy pixel size should not change for this microscope.
- Click Run and when prompted, select the TIFF file of your z-stack in the Raw Data folder.

- **Rotate and visualize the data:**

- Two more scripts are used to visualize and rotate the data to top view and side view. They can be provided to you by the Beckman center staff.

In practice, once your experiment is finished, the Beckman Center staff will give you the processed data. However, if you are an enthusiastic Fiji user and want to further process and analyze your data in Fiji you might find it useful to be able to perform the de-skewing operations yourself to tweak them to your needs.