I3D:bio OMERO user training slides

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https://www.i3dbio.de

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Research Data Management for Bioimage Data at the ADD INSTITUTE HERE

How to connect Fiji and OMERO?



ADD AUTHOR / RESPONSIBLE PERSON FROM YOUR INSTITUE



Connect Fiji and OMERO (1/2)

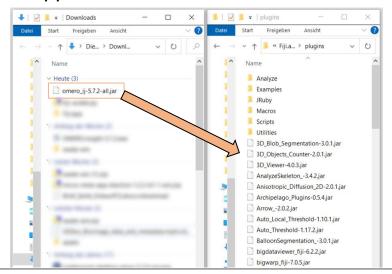
Prerequisite:

You have downloaded Fiji (https://fiji.sc) and have access to the OMERO instance (direct or VPN)

 Download the OMERO plugin for Fiji from the OME downloads website: https://www.openmicroscopy.org/omero/downloads



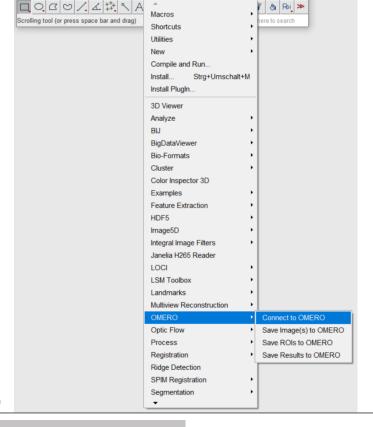
2. Move the *omero-ij-x.x.x-all.jar* file to the *Plugins* folder of your Fiji application



Connect Fiji and OMERO (2/2)

- Open Fiji and go to Plugins → OMERO → Connect to OMERO
- Log in to OMERO with your user credentials.





Plugins Window Help

III (Fiji Is Just) ImageJ

File Edit Image Process Analyze

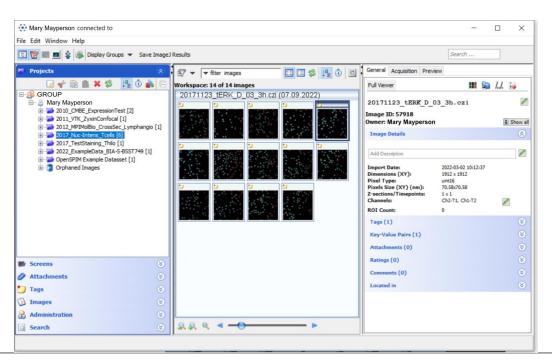
The Fiji-OMERO plugin looks almost precisely like OMERO.insight, but is, in fact, part of the open Fiji application



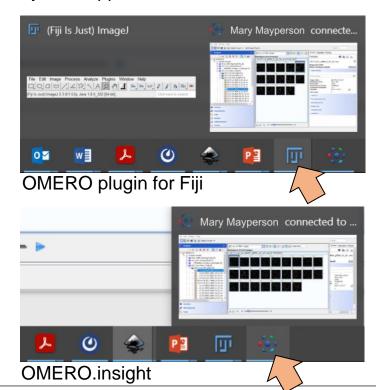
p 5

OMERO plugin for Fiji versus OMERO.insight

NOTE: The window looks similar to OMERO.insight, but it is a different application. For example, OMERO.insight has no View in ImageJ function nor allows Save ImageJ Results.



You can distinguish the applications by their appearance in the task bar.

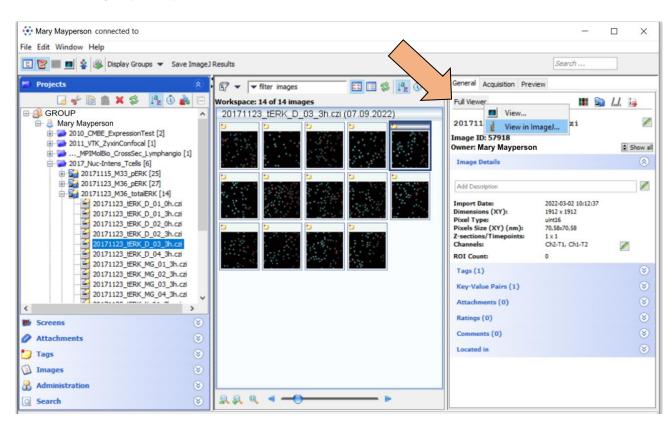




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Select image(s) to open in Fiji (1/2)

- Select image(s) from the file tree
- Open in Fiji by clicking Full Viewer and then View in ImageJ...



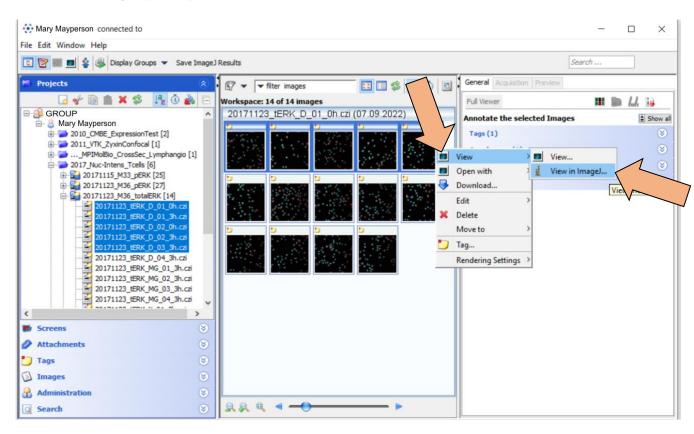


Select image(s) to open in Fiji (2/2)

OR

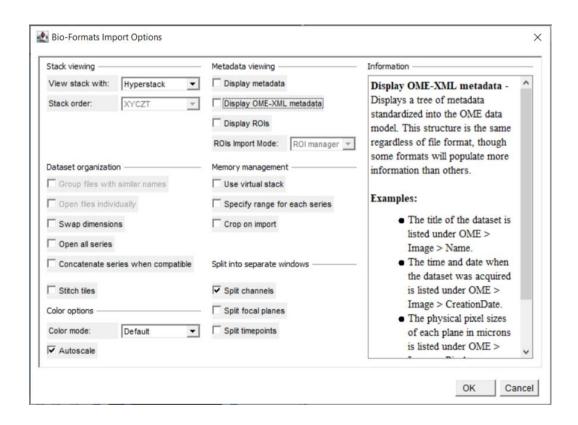
Open image(s) by right-click
View and then
View in ImageJ...

(or double-click)

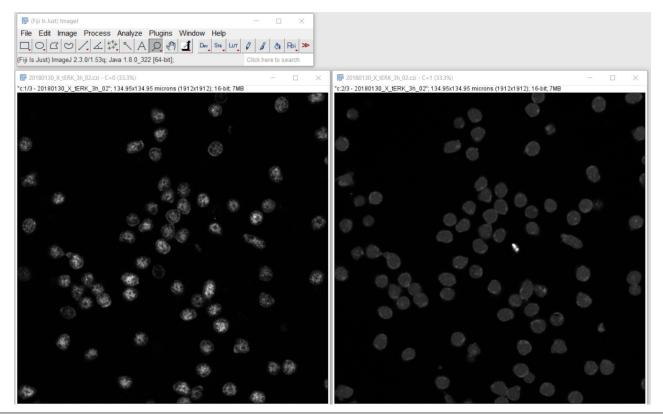


Choose settings for loading the image(s) in Fiji

Use your prefered settings to open the image(s) as required for your work



View your images in Fiji and work with the image for processing and analysis



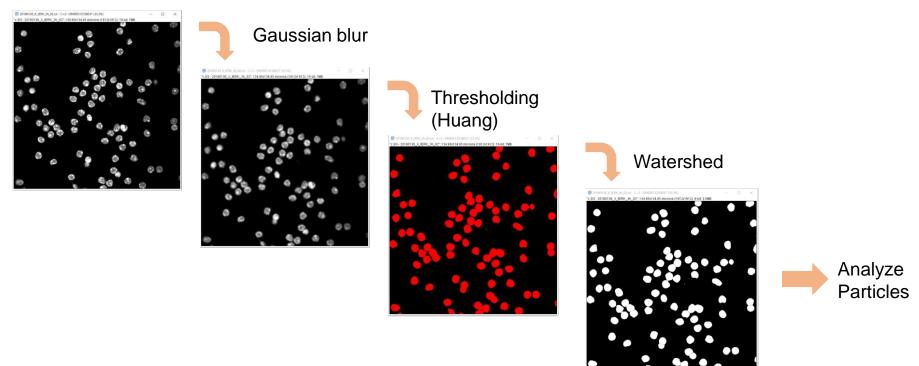


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Example – processing & analysis workflow to segment and count nuclei

Perform your workflow in Fiji

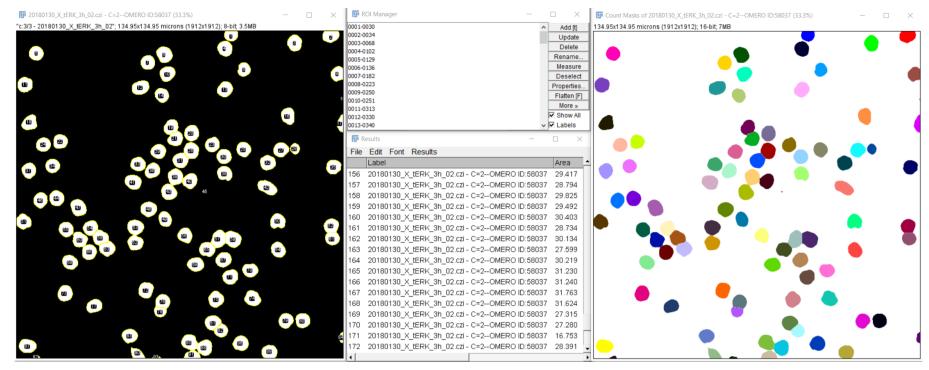
(here: segmentation and cell counting based on nuclear staining with DAPI)





Example – processing & analysis workflow to segment and count nuclei

Perform your workflow in Fiji (here: segmentation and cell counting based on nuclear staining with DAPI)



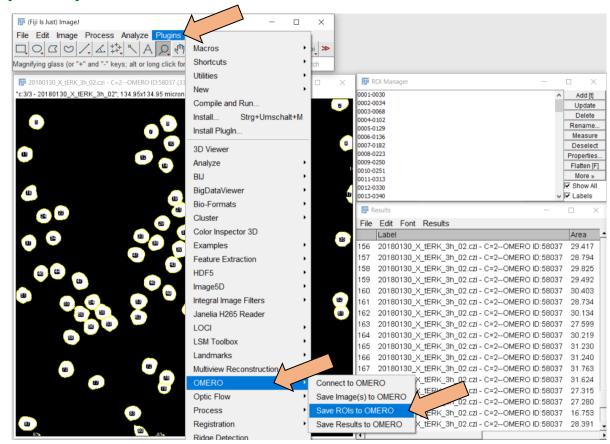


Save Regions of Interest (ROIs) and Measurement Results to OMERO

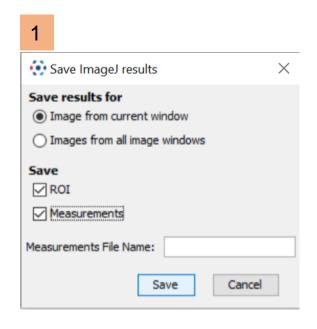
Save to OMERO using the plugin

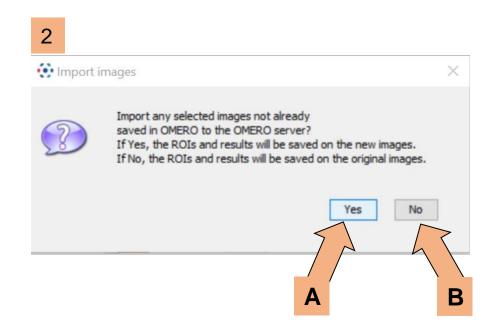
Plugins

- → OMERO
- → Save ROIs to OMERO



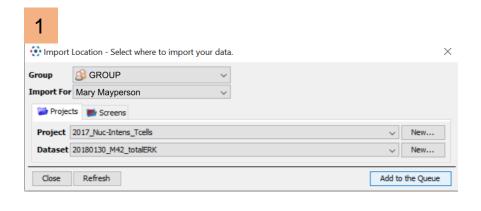
Choose settings for saving in OMERO

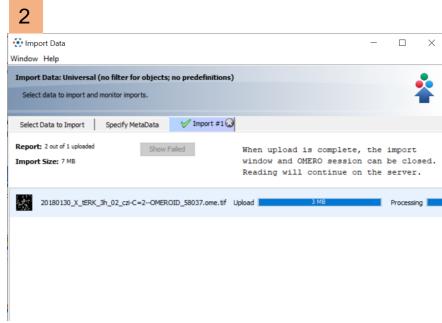




A – Upload the mask image to OMERO as a new image

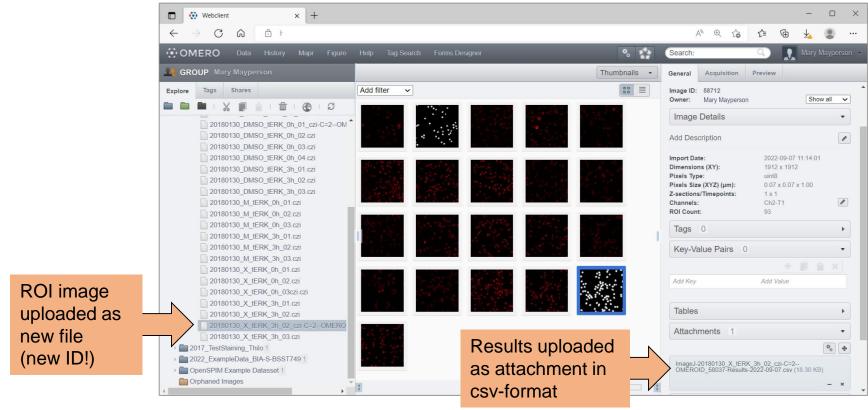
Choose upload destination (Group, User, Project, Dataset) and upload





A – View the imported mask image and the analysis results

(e.g., in OMERO.web)

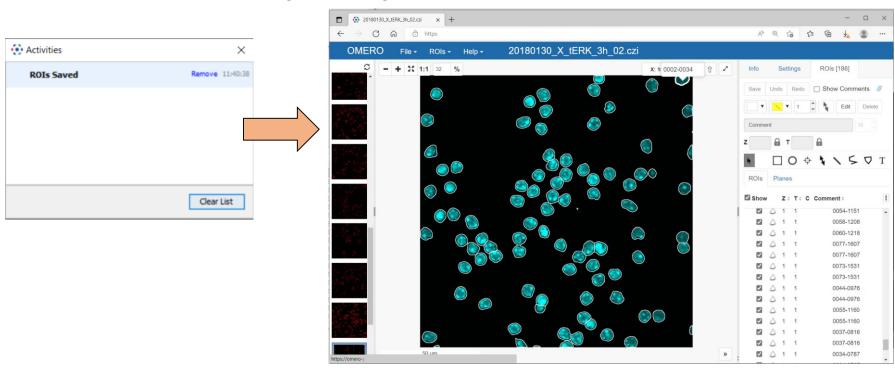




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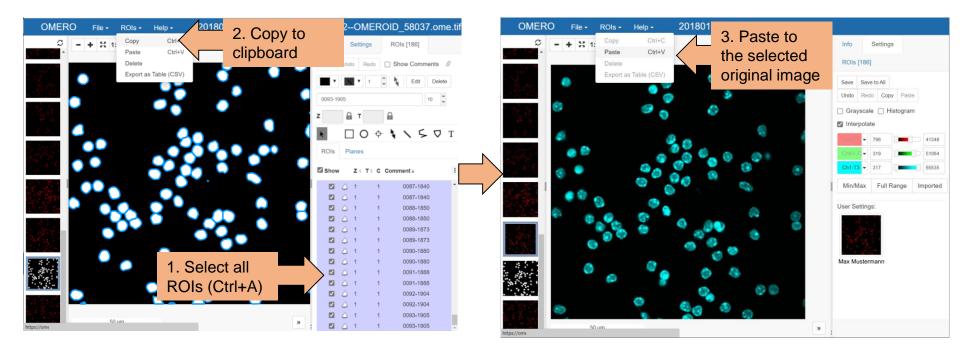
B – Add the ROI to the orginal image in OMERO

Review the ROIs on the original image with OMERO.iviewer



A and B combined

In the OMERO.iviewer you can copy the ROI from the segementation image to the original image manually, too.



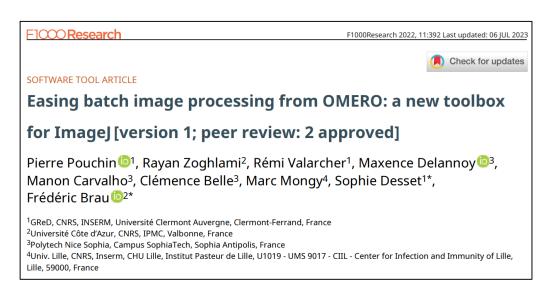


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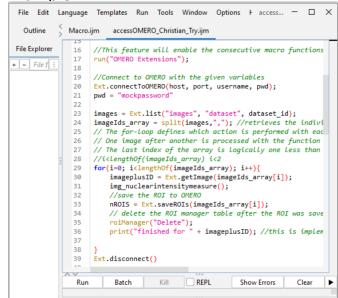
Batch processing, macros and scripts for Fiji and OMERO

Running image analysis pipelines with Fiji over several images from OMERO in batch is possible, too.

Recommended: OMERO Macro Extension & OMERO batch plugin



Example:



https://omero-guides.readthedocs.io/en/latest/fiji/docs/index.html



JiPipe visual macro programming with a connection to OMERO

Batch processing in Fiji/ImageJ and the connection to OMERO can now be established with a graphical user interface (GUI) in the software JiPipe:

Correspondence

https://doi.org/10.1038/s41592-022-01744-4

JIPipe: visual batch processing for ImageJ



he growth in microscopy adoption has led to a concomitant upsurge in the development of software tools for the automated analysis of image data. Pillars among these tools are ImageJ¹ and its Fiji² distribution, which have been serving the imaging community for decades and continue to gain public support to keep up with the quantification

needs of the newest and most-demanding microscopy techniques. The hallmark of ImageJ is its intuitive graphical user interface, which provides access to its many tools. On the other hand, the creation of reproducible batch-processing workflows is only possible using a macro language. As programming skills are uncommon among experimentalists³, the need for scripting contributes to an

already-existing communication gap between life and computer scientists. Visual programming languages that replace the writing of text commands with the design of a flowchart offer a solution. Existing tools contribute to this effort by providing a visual way to build pipelines or by simplifying the scripting procedure (Supplementary Information, section 1). Our newly developed visual programming











Extended resources on using Fiji and OMERO

Official OMERO guide:

https://omero-guides.readthedocs.io/en/latest/fiji/docs/index.html

A workshop on image analysis with Fiji and OMERO:

https://learning.rc.virginia.edu/notes/fiji-omero/

Workshop recordings by the Open Microscopy Environment Consortium on YouTube, including scripting in Fiji:

https://www.youtube.com/watch?v=W5EDx3yKA_o (https://www.youtube.com/watch?v=dOtnEO-nmlg)

Image Analysis Lecture by Robert Haase (TU Dresden): https://www.youtube.com/playlist?list=PL5ESQNfM5lc7SAMstEu082ivW4BDMvd0U

Help for Image Analysis or OMERO-related issues - Image.sc forum: https://image.sc

