

MRI-CIA has been created in 2023, to answer the bioimage analysis needs of the users of the core imageing facility Monpellier Ressources Imagerie and the wider community. We offer comprehensive support for bioimage analysis, from the start of a research project to publication. The support we provide is tailored to each individual case and can include oneon-one software training or the creation of custom analysis tools. We handle various aspects of image analysis, such as for example:

- \bigcirc image data handling
- \bigcirc image restoration and registration
- \bigcirc quantitative image analysis and visualization

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- \bigcirc instance segmentation and tracking
- \bigcirc morphological and intensity quantification
- O colocalization analysis and spatial statistics

OPEN DESK SESSIONS

We do an open desk session every fortnight. Do you have questions about image analysis? Do you need help with a script or macro? Are you planning to start an image analysis project with us? Then come join us at the next open-desk session. The access is free and no inscription is needed, but we ask you to kindly give us some information, that will help us planning the session.



tinyurl.com/v9suznr6





PROJECTS

The service is open to the public and private sector without any institutional or geographic conditions. MRI-CIA accepts image analysis projects within the field of life-science. The images to be analyzed, might be acquired on the MRI imaging core facility, but this is not a precondition for accessing the image analysis service.

The services provided by MRI-CIA include training and consulting in bio-image analysis and finding and implementing solutions for bio-image analysis problems. In the second case the preferred way is to develop tools and train the biologists to use them to perform their own analyses. In some cases, especially when new methods and technologies are involved, it can be preferable that MRI-CIA executes the analysis for the user.

The services of MRI-CIA, beyond the open desk sessions, are charged. We provide a free quote on request. Please find the current tariffs on our website.

Topics we often work on (non exhaustive) are:

Colocalization Intensity measurements Spot detection Cell morphology Filament Tracing A Tracking Cell motility Cell lineage Object counting Clustering





MRI CENTER FOR IMAGE ANALYSIS

Clément Benedetti, Volker Baecker

mri-cia@mri.cnrs.fr

TOOLS DEVELOPED BY MRI-CIA





tinyurl.com/ms824s4p

MD YEASTS







The tool measures the intensity of a fluorescent channel in the membranes of the mother and daughter cells of budding yeast.

Moving n-gons are used to find the membrane and to split the cells. An ngon starts as an ellipse or box around the budding yeast. It moves towards the center and stops according to some criteria. The curvature is used to find the splitting points.

SPOTS IN YEASTS



NEURITE ANALYZER

In s d l f bn h5 i bs bd bf b ≫



The workflow allows to analyze **RNA-FISH** spots on neurites. It counts the spots per cell and measures their distance to the soma along the neurite.

The images are acquired with the Phenix Opera and consist of 10x10 fields of 1000x1000 pixels each. They are exported and reconstructed with the help of the MRI-**Opera-Export-Tools**.

The neurites are segmented with a pretrained Ilastik classifier. The distances are calculated using the Geodesic Distance Map from the FIJI-plugin MorphoLibJ. Each point on a neurite is labelled with the identity of the closest connected soma

The tools estimate the width-profile of an object given as a binary mask. Four methods are provided: as local thickness, as voronoi distance between two parts of the contour-line, perpendicular to the axis of inertia and at regular distances using rays perpendicular to a centerline segment.

WIDTH PROFILE TOOLS





The napari-plugin analyzes lipid droplets in budding yeast cells. The cells are segmented with cellpose in the brightfield channel. In the result the mother-daugter pairs are separated. They are reconnected using the Hopcraft-Karp algorithm. The cells are considered as a bipartite graph, in which the cells with nucleus form one partition and those without the other. The tool reports the number of droplets per cell, their areas and intensities and their positions within the cell



Current features are:

- \bigcirc get the active image from FIJI
- \bigcirc send a screenshot to FIJI
- get a set of points from the FIJI results table
- \bigcirc filter the points in napari
- \bigcirc send the filtered points back to FIJI



(1) gt5 (1) gt5	(1) noise (1) n2v	19.20405144 38.72119838	26.77
(1) gt5	(1) gaussian-filter	34.12542003	41.69
(1) gt5	(1) gaussian-filter	34.12542003	

ImageJ Macro Programming

setBatchMode(true); while(next()) { run("Analyze Image"); setBatchMode("exit and display"); https://tinyurl.com/2p8tsnbc

3D Bio-Image Analysis





NAPARI-J

A plugin to exchange data with FIJI and to use FIJI image analysis from napari.



DL4MIC

WORKSHOPS

Machine Learning (ML/DL) for bio-image analysis



https://tinyurl.com/daxxb3mn

Check out the upcoming Biocampus workshops: www.biocampus.cnrs.fr/index.php/en/workshops-registration