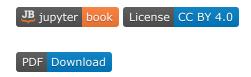
Fiji Exercises Collection

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Fiji Exercises

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By the **Image Analysis Collaboratory** @ Harvard Medical School

This is a collection of exercises for the open-source Fiji software.

The exercises are designed to be used in a classroom setting, but can also be used for self-study. They are divided into sections, each of which covers a different topic and/or a different aspect of Fiji.

Here is an example illustrating the structure of the exercises:

1.1 - Image inspection

- 1. open the single_channel.tif image (drag & drop, or File > Open)
- 2. Image > Duplicate (IJ:28.9) (shift + d)
- 3. Analyze > Histogram (IJ:30.10) (**h**)
- 4. click Live

- 5. Image > Adjust > Brightness/Contrast (shift + c)
 - play with Maximum, Minimum, Brightness and Contrast sliders and with Auto, Set
 and Reset
 - observe the histogram window, what is changing, what is not?
 - click Apply
 - what changed?

You can perform each exercise using the **provided image(s)** (click the image name to download it - note that downloads may not work properly in Safari) or with **your own images**.

For certain Fiji commands, references to the <u>ImageJ User Guide</u> are included (e.g. **IJ:28.9**) as a helpful resource for learning more about Fiji's features and capabilities.

If a command has a keyboard shortcut, it is indicated in parentheses in **bold** (e.g. **shift + d**).

Have fun!

1 - Working with Single-Channel Images

1.1 - Image inspection

```
1. open the \underline{\underline{*}} single_channel.tif image (drag & drop, or File > 0pen)
```

```
2. Image > Duplicate (IJ:28.9) (shift + d)
```

- 3. Analyze > Histogram (IJ:30.10) (h)
- 4. click Live
- 5. Image > Adjust > Brightness/Contrast (shift + c)
 - play with Maximum, Minimum, Brightness and Contrast sliders and with Auto, Set
 and Reset
 - observe the histogram window, what is changing, what is not?
 - click (Apply)
 - what changed?

1.2 - Adjust brightness/contrast of all open images

```
    open the three single_channel_...tif images in folder sfor_exercise_1.2 (drag & drop, or File > Open)
    for all images: Analyze > Histogram (IJ:30.10) (h)

            click Live

    for one image: Image > Adjust > Brightness/Contrast (shift + c)
    adjust contrast with Maximum, Minimum, Brightness or Contrast sliders or Auto
    click Set
    check Propagate to all other open images checkbox
    click OK
    what happened to the histograms?
    what happened to the images?
    did pixel values change?
    pick another image and repeat steps 4-7
    what happened, what is different, is it better/worse?
```

1.3 - File handling and non-invasive editing

- o saving the image as tiff keeps the scale bar as an overlay, so pixel values below it are kept
- NOTE: you can see the ovarlay scale bar in the saved image only if you open it in Fiji
- continue with <u>1.4</u>

1.4 - File handling and invasive editing

```
    open the blobs1.tif saved in 1.3 (drag & drop, or File > Open)
    Image > Info (IJ:28.3) (i)
    Image > Overlay > Remove Overlay (IJ:28.14.7)

            the scale bar should disappear since it was saved as an overlay

    Analyze > Set Scale (IJ:30.8)

            set scale such as image is 100 um long in each dimension

    Analyze > Tools > Scale Bar (IJ:30.14.6)

            set Color, Background, Location
            uncheck Overlay

    File > Save As > Tiff -> blobs2.tif (IJ:26.10.1)
```

2 - Working with Multi-Channel Images

pixel values)

2.1 - composite images - splitting and merging

7. load again to check the difference (e.g. check info, hover over the scale bar and look at the

```
    File > Open Samples > Fluorescent Cells (IJ:26.4)
    Image > Color > Arrange Channels

            click on New 1, and select Magenta

    split channels

            Image > Color > Split Channels (IJ:28.5.1)

    merge channels to composite

            Image > Color > Merge Channels (IJ:28.5.2)
            check create composite checkbox (IJ:28.5.2)

    LUT (look-up table) (IJ:19.17 & IJ: 28.15)

            change LUT of the channels (LUT in the startup tools or Image > Lookup Tables)

    color blindness
```

```
    convert to RGB Color: Image > Type > RGB Color (IJ:7)
    Image > Color > Simulate Color Blindness
    Image > Color > Dichromacy
```

3 - Image Processing: Filters

3.1 - Edge filters - vertical stripes

```
1. open the <a href="mailto:vertical_stripes.tif">vertical_stripes.tif</a> image (drag & drop, or File > 0pen)
```

```
2. Image > Duplicate (IJ:28.9) (shift + d)
```

3. apply horizontal Prewitt filter:

```
• [Process > Filters > Convolve]
```

• as kernel input (Prewitt):

- click on OK.
- do you understand the output of this process?
- 4. select the original vertical_stripes.tif image and duplicate it again.

```
• Image > Duplicate (IJ:28.9) (shift + d)
```

5. apply vertical Prewitt filter:

```
• Process > Filters > Convolve
```

• as kernel input (Prewitt):

- click on OK.
- o do you understand the output of this process?

3.2 - Edge filters - horizontal stripes

The steps in this exercise are identical to 3.1, only the input image differs.

```
1. open the <a href="https://www.horizontal_stripes.tif">horizontal_stripes.tif</a> image (drag & drop, or [File > Open])
```

```
2. Image > Duplicate (IJ:28.9) (shift + d)
```

3. apply horizontal Prewitt filter:

```
• Process > Filters > Convolve
```

• as kernel input (Prewitt):

```
-1 -1 -1
0 0 0
1 1 1
```

- click on OK.
- do you understand the output of this process?
- 4. select the original vertical_stripes.tif image and duplicate it again.
 - Image > Duplicate (IJ:28.9) (shift + d)
- 5. apply vertical Prewitt filter:
 - [Process > Filters > Convolve]
 - as kernel input (Prewitt):

- click on OK.
- o do you understand the output of this process?

3.3 - Edge filters - checkerboard

The steps in this exercise are identical to 3.1 and 3.2 only the input image differs.

1. open the <u>★ checkerboard.tif</u> image (drag & drop, or File > Open)

- 2. Image > Duplicate (IJ:28.9) (shift + d)
- 3. apply horizontal Prewitt filter:
 - Process > Filters > Convolve
 - as kernel input (Prewitt):

```
-1 -1 -1
0 0 0
1 1 1
```

- click on OK.
- o do you understand the output of this process?
- 4. select the original checkerboard.tif image and duplicate it again.
 - Image > Duplicate (IJ:28.9) (shift + d)
- 5. apply vertical Prewitt filter:
 - Process > Filters > Convolve
 - as kernel input (Prewitt):

- o click on OK.
- o do you understand the output of this process?

3.4 - Morphological filters - Binary

- 1. open the $\underline{\underline{}}$ exercise_morphology.tif image (drag & drop, or [File > 0pen])
- 2. duplicate the image 4 times and name each copy as follow:
 - Eroded
 - Dilated
 - Closed
 - Opened
- 3. apply binary morphological filters:
 - Process > Binary (Erode, Dilate, Close, Open)

- Note: Make sure you apply the operation that corresponds to the name of each image.
- 4. report findings

3.5 - Morphological filters - Gray

- 1. open the <u>kexercise_morphology.tif</u> image (drag & drop, or [File > Open])
- 2. apply Gray Morphological filters:

```
• Process > Morphology > Gray Morphology
```

- 3. apply image **opening** with **circular** structuring element with **radius 3** removes noise
- 4. Report: did it get rid of the noise?
- 5. apply image opening with circular structuring element with radius 5
- 6. report: did it get rid of the tentacles from the top-left object?
- 7. with the line tool (IJ:19.2.1), measure the diameter of the holes in the bottom-right object.
- 8. apply image closing with circular structuring element with radius slightly larger than the measured radius (diameter/2).
- 9. report: did it get rid of the tentacles from the top-left circle?

4 - Segmentation

4.1 - DAPI segmentation with thresholding

```
1. open the LDAPI.tif image (drag & drop, or File > Open)
```

- 2. change LUT to Grays
- 3. Image > Duplicate (IJ:28.9) (shift + d)
- 4. Image > Adjust > Threshold (IJ:28.2.4)
- 5. understand the function of the Dark Background checkbox (inspect pixel values)
- 6. try setting sliders manually. Can you find a good threshold range?
- 7. try different algorithms by selecting them in the left dropdown menu. Can you find one that gives a good result?
 - NOTE: Image > Adjust > Auto Threshold, if you want to see all at the same time

- 8. try different display options (Red, B&W, Over/Under) by selecting them in the right dropdown menu, do you understand what they show?
- 9. when happy with result, click Apply
- 10. save the resulting binary image: File > Save As > Tiff
- 11. apply watershed to divide touching objects
 - select the binary image
 - Process > Binary > Watershed
- 12. proceed with Analyze > Analyze Particles (IJ:30.2)
 - select Exclude on Edges and Add to Manager
 - click on 0K
- 13. bonus: repeat step 12 but use the Size and Circularity options to try to exclude some particles and the Show dropdown menu to visualize different outputs.
- 14. set the parameters you want to measure:
 - Analyze > Set Measurement (IJ:30.2)
 - select Area, Mean gray value, Min & max gray value, Display label
 - click on [0K]
- 15. select the original image (open it again as in step 1 if you do not have it)
- 16. in the ROI Manager, click on Deselect and then on Measure
- 17. save the Results table as .csv: select the table and click on File > Save As

4.2 - DAPI segmentation with filters and thresholding

- 1. open the LDAPI_noise.tif image (drag & drop, or File > Open)
- 2. change LUT to Grays
- 3. Image > Duplicate (IJ:28.9) (shift + d)
- 4. Image > Adjust > Threshold (IJ:28.2.4)
- 5. understand the function of the Dark Background checkbox (inspect pixel values)
- 6. try setting sliders manually. Can you find a good threshold range?
- 7. try different algorithms by selecting them in the left dropdown menu. Can you find one that gives a good result?
 - NOTE: Image > Adjust > Auto Threshold, if you want to see all at the same time
- 8. apply a filter of your choice (Mean), Gaussian Blur, Median, ...)

```
• Process > Filters
```

- check the Preview checkbox
- change the Radius / Sigma. What happens to the image?
- when you are happy, click on 0K
- 9. repeat steps 3, 7 and 8 until happy with result, then click Apply
- 10. save the resulting binary image: File > Save As > Tiff
- 11. apply watershed to divide touching objects
 - select the binary image
 - o Process > Binary > Watershed
- 12. proceed with [Analyze > Analyze Particles] (IJ:30.2)
 - select Exclude on Edges and Add to Manager
 - click on OK
- 13. bonus: repeat step 12 but use the Size and Circularity options to try to exclude some particles and the Show dropdown menu to visualize different outputs.
- 14. set the parameters you want to measure:
 - [Analyze > Set Measurement] (IJ:30.2)
 - select Area, Mean gray value, Min & max gray value, Display label
 - click on OK
- 15. select the original image (open it again as in step 1 if you do not have it)
- 16. in the ROI Manager, click on Deselect and then on Measure
- 17. save the Results table as .csv: select the table and click on File > Save As

4.3 - DAPI segmentation with Weka

- 1. open the <u>khela.tif</u> image (drag & drop, or File > 0pen)
- 2. change LUT to Grays
- 3. Plugins > Segmentation > Trainable Weka Segmentation
- 4. draw a line outside a nucleus
- 5. click Add to class 1
- 6. draw a line inside a nucleus
- 7. click Add to class 2

- 8. click Train classifier
- 9. repeat the last four steps until happy with result
- 10. click Get probability
- 11. inspect the probability maps, do you understand the meaning of the values of the pixels in the different channels?
- 12. click Create result
- 13. inspect results, do you understand the meaning of the values of the pixels in the different channels?
 - you now have a binary image, but not the kind Fiji likes
 - to measure, proceed by thresholding ("set" both threshold values to 1), then Analyze Particles, etc
 - o alternatively, multiply all values in the Weka output image by 255, then apply Binarize, etc
- 14. save the resulting image with name "myWekaHeLa1.tif": File > Save As > Tiff
- 15. bonus round: play with Settings

5 - Spot Detection

5.1 - Manual spot detection with the Multi-point Tool

- 1. read all the next steps before beginning
- 2. open the beads_001.tif image in folder <u>spot_detection</u> (drag & drop, or File > 0pen)
- 3. right-click (ctrl-click) on the Point Tool in the tool bar and select MultiTool (IJ:19.5)
- 4. double-click on the Point Tool in the toolbar and customize to your liking (test on image, clear points by shift + a)
- 5. count all beads by clicking on them one by one
- 6. how long did it take you to click on all the beads (measure with stopwatch the time it took to complete the previous step)?
- 7. add the points to the ROI Manager (e.g. press t on your keyboard)
 - NOTE: only one ROI will be added, it contains all the points
- 8. save the ROI to file (in the ROI Manager window, More > Save)
- 9. click measure in the ROI Manager to get number of spots

5.2 - Algorithmic spot detection with Find Maxima

useful Find Maxima info on the image.sc forum.

- 1. open the beads_001.tif image in folder <u>spot_detection</u> (drag & drop, or File > Open)
- 2. Process > Find Maxima
- 3. check Preview point selection
- 4. try different values for Prominence and the three checkboxes
- 5. try each of the possibilities in the pull-down (remember to click OK to apply your selection)
 - what is the difference and what could the different outputs be used for?
 - tip: shift + u allows you to pin sub-menus of the control panel for quick selection (e.g. the Process menu)
- 6. add you detected maxima to the ROI Manager and save to file
 - using your preferred settings, set the Output type to Point Selection and press OK
 - add the points to the ROI Manager (e.g. press t on your keyboard)
 - NOTE: only one ROI will be added, it contains all the points.
 - save the ROI to file (in the ROI Manager window, More > Save).
- 7. load saved ROIs from the manual detection exercise (5.1) (e.g. drag & drop on Fiji the ROIs .zip file)
- 8. compare results: how many beads do you get and how does the number compare to your manual count?

5.3 - Automatic spot segmentation with thresholding

- 1. open the beads_001.tif image in folder <u>spot_detection</u> (drag & drop, or File > 0pen)
- 2. Image > Adjust > Threshold (IJ:28.2.4)
 - select Otsu, then Apply
- 3. Analyze > Set Measurements (IJ:30.7)
 - select (Area), (Mean gray value), (Display label)
 - click on OK
- 4. Analyze > Analyze Particles (IJ:30.2)

- select the Overlay Masks option in the Show dropdown menu
- select Display results, Clear Results, Summarize, Add to Manager
- 5. save the ROIs to file (in the ROI Manager window, More > Save)
- 6. Compare results to the previous two approaches ($\underline{5.1}$ and $\underline{5.2}$)
 - Do they differ significantly from each other?
 - Which performed better?

5.4 - Spot detection with noise

- 1. Repeat 5.2 with the with_noise_8000.tif image in folder & spot_detection
 - consider smoothing the image first. You can try any of the filter in the Process >
 Filters menu:
 - e.g. Process > Filters > Gaussian Blur, Process > Filters > Mean, ...
 - NOTE: the Smooth operation under the Process > Smooth is a mean filter wuth radius 1 (3x3 kernel)
- 2. Repeat 5.3 with the with_noise_8000.tif image in folder <u>spot_detection</u>
 - consider smoothing the image first. You can try any of the filter in the Process >
 Filters menu:
 - e.g. Process > Filters > Gaussian Blur, Process > Filters > Mean, ...
 - NOTE: the Smooth operation under the Process > Smooth is a mean filter wuth radius 1 (3x3 kernel)
- 3. Compare and comment on the performance of the two methods
 - Which one is better?
 - Why/how is it better?

5.5 - Spot detection with variable background

- 1. Repeat <u>5.2</u> with the beads_001_ramp.tif image in folder <u>spot_detection</u>
- 2. Repeat 5.3 with the beads_001_ramp.tif image in folder & spot_detection
- 3. Compare and comment on the performance of the two methods
 - Which one is better?
 - Why/how is it better?

4. What could be done to the image to make thresholding work better?	