Fiji Exercises Collection

Image Analysis Collaboratory @ Harvard Medical School

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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) (\mathbf{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 ImageJ User Guide 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!

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1 - WORKING WITH SINGLE-CHANNEL IMAGES

1.1 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) (\mathbf{h})
- 4. click Live
- 5. Image > Adjust > Brightness/Contrast $(\mathbf{shift} + \mathbf{c})$
 - 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 1.2 - Adjust brightness/contrast of all open images

- 1. open the three single_channel_...tif images in folder for_exercise_1.2 (drag & drop, or File > Open)
- 2. for all images: Analyze > Histogram (IJ:30.10) (h)
 - click Live
- 3. for one image: Image > Adjust > Brightness/Contrast (shift + c)
- 4. adjust contrast with Maximum, Minimum, Brightness or Contrast sliders or Auto
- 5. click Set
- $6.\ check\ {\hbox{\sc Propagate}}$ to all other open images checkbox
- 7. click OK
- 8. what happened to the histograms?
- 9. what happened to the images?
- 10. did pixel values change?
- 11. pick another image and repeat steps 4–7
- 12. what happened, what is different, is it better/worse?

1.3 1.3 - File handling and non-invasive editing

- 1. File > Open Samples > Blobs (IJ:26.4) ($\mathbf{shift} + \mathbf{b}$)
- 2. Analyze > Tools > Scale Bar (IJ:30.14.6)
 - set Color, Background, Location
 - check: Overlay, Bold Text
- 3. File > Save As > Tiff->blobs1.tif(IJ:26.10.1)
 - 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 1.4

1.4 1.4 - File handling and invasive editing

- 1. open the blobs1.tif saved in 1.3 (drag & drop, or File > Open)
- 2. Image > Info (IJ:28.3) (i)
- 3. Image > Overlay > Remove Overlay (IJ:28.14.7)
 - the scale bar should disappear since it was saved as an overlay
- 4. Analyze > Set Scale (IJ:30.8)
 - set scale such as image is 100 um long in each dimension
- 5. Analyze > Tools > Scale Bar (IJ:30.14.6)
 - set Color, Background, Location
 - uncheck Overlay
- 6. File > Save As > Tiff -> blobs2.tif (IJ:26.10.1)
- 7. load again to check the difference (e.g. check info, hover over the scale bar and look at the pixel values)

2 - WORKING WITH MULTI-CHANNEL IMAGES

2.1 2.1 - composite images - splitting and merging

```
1. File > Open Samples > Fluorescent Cells (IJ:26.4)
```

- 2. Image > Color > Arrange Channels
 - click on New 1, and select Magenta
- 3. split channels
 - Image > Color > Split Channels (IJ:28.5.1)
- 4. merge channels to composite
 - Image > Color > Merge Channels (IJ:28.5.2)
 - check create composite checkbox (IJ:28.5.2)
- 5. LUT (look-up table) (IJ:19.17 & IJ: 28.15)
 - change LUT of the channels (LUT in the startup tools or Image > Lookup Tables)
- 6. 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 3.1 - Edge filters - vertical stripes

- 1. open the vertical_stripes.tif image (drag & drop, or File > Open)
- 2. Image > Duplicate (IJ:28.9) ($\mathbf{shift} + \mathbf{d}$)
- 3. apply horizontal Prewitt filter:
 - Process > Filters > Convolve
 - as kernel input (Prewitt):

```
\begin{pmatrix}
-1 & -1 & -1 \\
0 & 0 & 0 \\
1 & 1 & 1
\end{pmatrix}
```

- 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.
- do you understand the output of this process?

3.2 3.2 - Edge filters - horizontal stripes

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

- 1. open the horizontal_stripes.tif 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):

- 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):

```
\begin{pmatrix}
-1 & 0 & 1 \\
-1 & 0 & 1 \\
-1 & 0 & 1
\end{pmatrix}
```

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

3.3 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 checkerboard.tif image (drag & drop, or File > Open)
- 2. Image > Duplicate (IJ:28.9) ($\mathbf{shift} + \mathbf{d}$)
- 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 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):

```
\begin{pmatrix}
-1 & 0 & 1 \\
-1 & 0 & 1 \\
-1 & 0 & 1
\end{pmatrix}
```

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

3.4 3.4 - Morphological filters - Binary

- 1. open the exercise_morphology.tif image (drag & drop, or File > Open)
- 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 3.5 - Morphological filters - Gray

- 1. open the exercise_morphology.tif 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?

FOUR

4 - SEGMENTATION

4.1 4.1 - DAPI segmentation with thresholding

- 1. open the DAPI.tif image (drag & drop, or File > Open)
- 2. change LUT to Grays
- 3. Image > Duplicate (IJ:28.9) ($\mathbf{shift} + \mathbf{d}$)
- 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 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

4.2 4.2 - DAPI segmentation with filters and thresholding

- 1. open the DAPI_noise.tif image (drag & drop, or File > Open)
- 2. change LUT to Grays
- 3. Image > Duplicate (IJ:28.9) ($\mathbf{shift} + \mathbf{d}$)
- 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 OK
- 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
 - 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 4.3 - DAPI segmentation with Weka

- 1. open the hela.tif image (drag & drop, or File > Open)
- 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
 - 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 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 spot_detection (drag & drop, or File > Open)
- 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 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 spot_detection (drag & drop, or File > Open)
- 2. Process > Find Maxima
- $3.\ check\ { t Preview\ point\ selection}$
- 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 5.3 - Automatic spot segmentation with thresholding

- 1. open the beads_001.tif image in folder spot_detection (drag & drop, or File > Open)
- 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 (5.1 and 5.2)
 - Do they differ significantly from each other?
 - Which performed better?

5.4 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 with radius 1 (3x3 kernel)
- 2. Repeat 5.3 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)
- 3. Compare and comment on the performance of the two methods
 - Which one is better?
 - Why/how is it better?

5.5 5.5 - Spot detection with variable background

- 1. Repeat 5.2 with the <code>beads_001_ramp.tif</code> image in folder spot_detection
- 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?