

MTQuant DOCUMENTATION

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1 MTQuant Overview

This document details the usage of the MATLAB function `MTQuant`. `MTQuant` rapidly quantifies neuronal microtubule organization by extracting three parameters from fluorescence images of neuronal microtubules. Collectively referred to as the "organization parameters," the three parameters are:

- **Spacing** (\bar{S}): average number of microns between microtubule ends
- **Coverage** (\bar{C}): average number of microtubules in bundle cross-section
- **Length** (\bar{L}): average length of microtubules in a single neuron

The `MTQuant` code allows users to perform four tasks:

- **Task 1:** Generate Maximum Projections
- **Task 2:** Trace and Generate Neurite Masks
- **Task 3:** Generate Intensity Line Scans
- **Task 4:** Calculate Organization Parameters

Throughout this document, we assume a basic working knowledge of MATLAB: strings, cell arrays of strings, comments, using the command line, etc. Typically, a user will begin with either image stacks or maximum projection images, and then complete Tasks 2-4. Task 2, Trace Neurite Masks, is the only step that *requires* user input. The remainder of this document will describe the four tasks in greater detail and provide some usage examples. The specifics of the input arguments, output files, and naming conventions are included as well.

1.1 Citation

If you use this code in your research, please cite it as:

R. Cooper, S. Yogev, K. Shen, and M. Horowitz, *MTQuant: "Seeing" Beyond the Diffraction Limit in Fluorescence Images to Quantify Neuronal Microtubule Organization*, (in preparation, 2016).

1.2 Setup & System Requirements

`MTQuant` and its supporting code can be downloaded at:

<http://roscoope.github.io/MTQuant>

The code can be downloaded as a ZIP file or checked out as a Github repository. Download all of the files and keep them together. The either run the code in the top-level `MTQuant`

directory, or elsewhere by typing the following commands into MATLAB to add the appropriate directories to the path:

```
>> addpath('C:/path/to/MTQuant')
>> addpath('C:/path/to/MTQuant/em')
>> addpath('C:/path/to/MTQuant/misc')
>> addpath('C:/path/to/MTQuant/neuronTracing')
>> addpath('C:/path/to/MTQuant/redScan')
>> addpath('C:/path/to/MTQuant/singleMT')
```

If you copy and paste commands from this document into MATLAB directly, some characters, such as curly braces (`{}`) might not copy correctly; be sure to review copied commands before running them.

Thus far, `MTQuant` has been tested on Windows 7 and 8, using MATLAB versions R2014b and R2013b.

2 MTQuant Functionality Details

This section will describe the inputs and outputs of each of the tasks, and also provide detailed instructions on the user input required in Task 2. There are a few interspersed examples of command line execution of `MTQuant` in MATLAB, which are collected and augmented in Section 3. By default, `MTQuant` completes all four of the tasks mentioned above.

`MTQuant` always requires two inputs: a list of directories in which to find the images to process, and the name of a CSV file to store the data. The program searches the directories in the list, as well as all subdirectories, for the files to analyze. ***The list of directories must be a cell array of strings.*** Furthermore, the output filename must ***not*** have an extension. For example:

```
>> dirList = {'C:/path/to/images/'; './'};
>> fileOut = 'testOutput';
>> MTQuant(dirList,fileOut);
```

To change the default values of various variables, `MTQuant` uses MATLAB's name-value pair format. Examples, once again, are in Section 3, and a full list of the input and output arguments is in Section 5.1.

2.1 Task 1: Generate Maximum Projections

When generating maximum projections, by default `MTQuant` searches for green and red image stacks that end in `'*_w1488-single.TIF'` and `'*_w2561-single.TIF'`, respectively. Both of these strings can be changed with an input argument name-value pair. The program will simply ignore processing of the red channel if there are no red stacks present. Thus

far, **MTQuant** has been tested with 16-bit .TIF images. The output projections are written with the same absolute path as the stack images, but with '`*_w1488-single.TIF`' and '`*_w2561-single.TIF`' replaced by '`*_g_proj.tif`' and '`*_r_proj.tif`', respectively. Naming conventions are described in more detail in Section 5.3. **MTQuant** also writes a CSV file which contains the stack slice from which each pixel was selected in a file saved with the suffix '`*_stackInds.csv`'.

2.2 Task 2: Trace and Generate Neurite Masks

The program requires some user input in order to determine which portions of the neuron are of interest to the researcher. Figure 1 shows example images of the necessary human input. For each image ending in '`*_g_proj.tif`', the image is displayed to the user (Figure 1(a)), and the user is asked whether or not they would like to trace a neurite. When the user selects yes, they are prompted to click along the neuron. The user needs to click points along the section of the neuron that they are interested in, beginning and ending with the endpoints of that section. For example, to trace part of the axon in, the user needs to click on the points labeled in Figure 1(b).

Then the user is prompted to click on the end of the trace that corresponds to the minus-ends of the MTs, denoted by the green star in Figure 1(c).

Finally, the user needs to click on a box inside the animal to capture the background intensity so that it can be subtracted later from the line scans. See Figure 1(d).

At each step along the way, the program asks the user to confirm whether they are satisfied with their selection. If the user wants to change the trace, they can simply type 'n' at those prompts and re-select the trace or the background box. An example command line interaction is shown in Section 3.4.

This function writes out two files: one is an image file (.TIF) of the mask, and another is a .CSV file which stores the mask as well as the background and endpoint information. These files end in '`*_Mask.tif`' and '`*_Mask.csv`', respectively. To allow for separate labeling of various sections, the user can use the '`indivFileOut`' input argument to prepend a label to the mask files, such as '`axon`', yielding, for example, '`_axonMask.csv`'.

2.3 Task 3: Generate Intensity Line Scans

MTQuant generates line scans and saves them in files with suffix '`*_LineScans.csv`'. The first column of this file is the green line scan, with the first value corresponding to the location of the green star in Figure 1(c), or the minus-ends of the MTs. The second column is the red line scan, oriented the same way.

The program also saves the pixel locations that were summed to generate the line scans in the file '`*_LineScanInds.csv`'. As with the saved mask files, the '`indivFileOut`' input argument allows the user to prepend a label to these suffixes. The format of this file is as follows: If

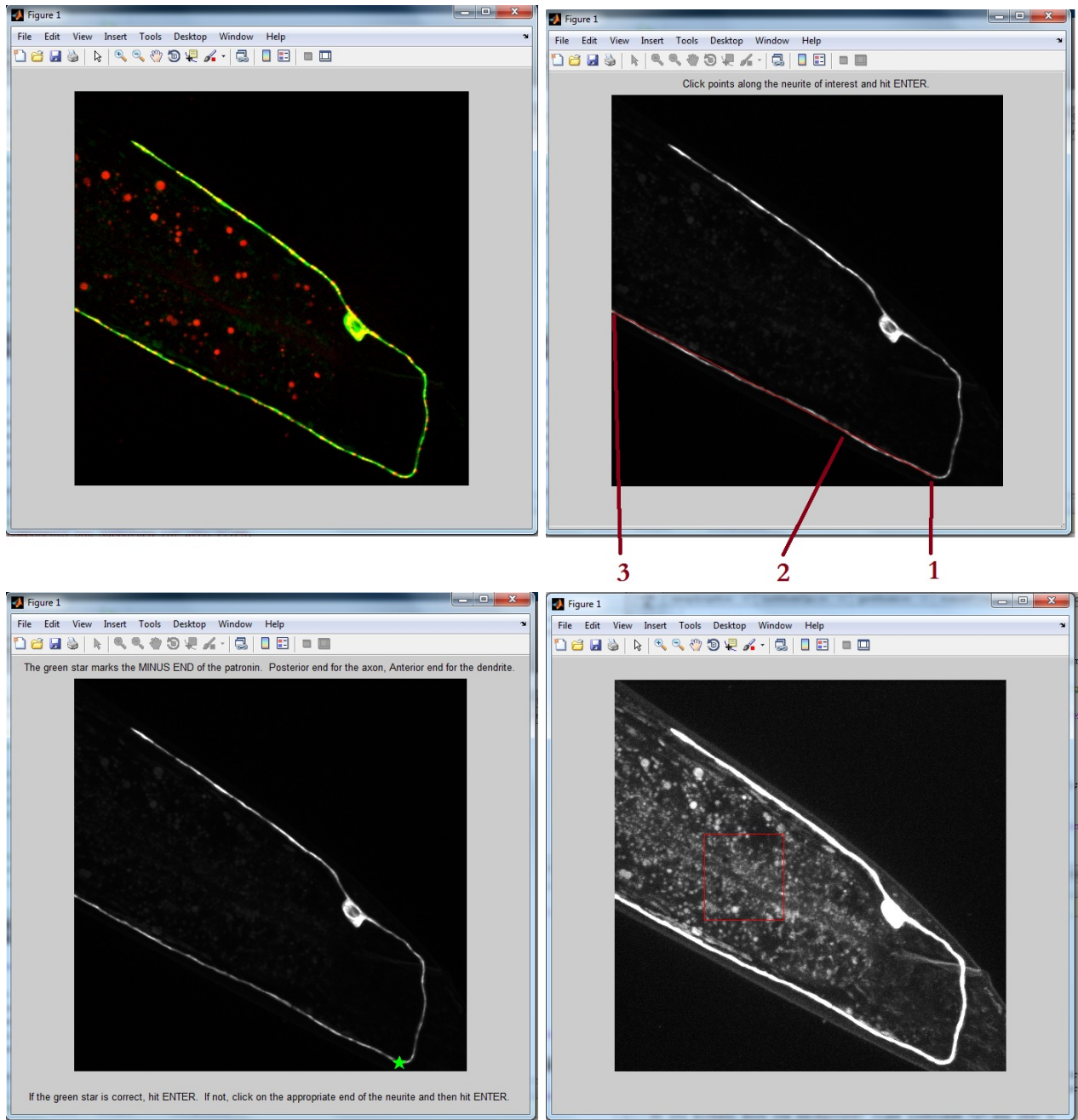


Figure 1: Clock-wise from top left: (a) Multi-color Image. (b) Tracing the Axon Mask: The numbers 1, 2, and 3 correspond to the locations clicked by the user to generate the red line tracing the axon. (c) Select the orientation of the MT minus-ends. (d) Select the background box.

the line scan has length d , Its dimensions are $d \times 402$. The first 201 columns refer to the y -coordinate of each point, and the second set of 201 columns refers to the x -coordinate. Hence, the starting (x, y) location of the spline is in $(1, 302)$ and $(1, 101)$ in this CSV file.

2.4 Task 4: Calculate Organization Parameters

The final task of **MTQuant** is to calculate the organization parameters of the animals. If the flag `toRateImgs` is set to true (it is false by default), the program stops after each image to ask the user for a rating: 2 for a good image, 1 for a barely acceptable image, and 0 for a bad image. Then, it generates two CSV files for each of the following groupings: all good images (group 2), all acceptable images (groups 1 and 2), and all images (groups 0, 1, 2).

The first CSV file, with the name of the input argument `fileOut`, contains an entry for every image analyzed. The list of entries in the table is summarized in Section 5.2.

The second file is has the suffix `'*_grouped.csv'`. The program assumes that, if the input directory list is only one element long, that each sub-folder of that directory is a separate group of worms. **MTQuant** assigns each group a number, which is included in `fileOut`, and used for both the distribution refinement. The `'*_grouped.csv'` file calculates the mean and standard deviation for each parameter in Section 5.2 for each separate group of worms.

3 Example Usage

3.1 Setup

```
>> addpath('C:/path/to/MTQuant')
>> addpath('C:/path/to/MTQuant/em')
>> addpath('C:/path/to/MTQuant/misc')
>> addpath('C:/path/to/MTQuant/neuronTracing')
>> addpath('C:/path/to/MTQuant/redScan')
>> addpath('C:/path/to/MTQuant/singleMT')
```

3.2 Run MTQuant with Default Settings

```
>> dirList = {'C:/path/to/images/', './'};
>> fileOut = 'testOutput';
>> MTQuant(dirList,fileOut);
```

3.3 Run MTQuant with Various Name-Value Input Argument Pairs

```
>> dirList = {'C:/path/to/images/', './'};
>> fileOut = 'testOutput';
>> taskList = 14; % we have already made the projections
>> toRateImgs = 1; % we suspect many images are bad, let's rate them
>> MTQuant(dirList,fileOut,'taskList',taskList,'toRateImgs',toRateImgs);
```

3.4 Example Output During Task 2 (Neurite Tracing)

```
>> Do you want to trace an axon? (lowercase "n" for no) y
>> Are you happy with this trace? (lowercase "n" for no) y
>> The green star marks the MINUS END of the patronin. Posterior end for the axon, Anterior
    end for the dendrite.
>> If the green star is correct, hit ENTER. If not, click on the appropriate end of the neurite
    and then hit ENTER.
>> Select the green background.
>> Do you already know the background? (type lowercase "y" for yes) n
>> Please click on two opposite corners of the box you want to average for the background
>> Are you happy with the location of this rectangle? (type lowercase "y" for yes) y
>> background = 2146.8372
>> Select the red background.
>> background = 1984.9885
```

4 Directory Organization Naming Conventions

Every analyzed line scan is given a group number (described in `DirNum` in Table 2). `DirNum` is assigned as described below. First, consider the following directories:

- `C:/path/to/data`
- `C:/path/to/data/wild_type`
- `C:/path/to/data/mutant1`
- `C:/path/to/data/mutant2`

Assume that `C:/path/to/data` contains the three subdirectories `wild_type`, `mutant1`, and `mutant2`, and no other images. The three subdirectories can contain images, subdirectories with more images, or a combination of both. Depending on the input `dirList` to `MTQuant`, the assignment of the output `DirNum` varies.

If `dirList` is just a single directory (e.g., `dirList = {'C:/path/to/data'};`), then all subdirectories contained in `dirList` are considered a separate group. Hence, images in the `wild_type` subdirectory (and its subdirectories) would be in group 1, `mutant1` images in group 2, and `mutant2` images in group 3. The output `DirNum` is 1 for the images in group 1 (`wild_type`), 2 for images in group 2 (`mutant1`), and 3 for images in group 3 (`mutant2`).

If `dirList` is a list of directories (e.g., `dirList = {'C:/path/to/data/mutant2'; 'C:/path/to/data/mutant1'};`), then each directory in the list is considered a separate group, and numbered based on the order it appears in `dirList`. Therefore, images in the `mutant2` subdirectory (and its subdirectories) would be in group 1 and `mutant1` images in group 2. Again, `DirNum` is 1 for the images in group 1 (`mutant2`) and 2 for images in group 2 (`mutant1`).

5 Appendix

5.1 Full List of Input Arguments

Input Argument	Description	Default Value
REQUIRED ARGUMENTS		
<code>dirList</code>	List of directories in which to find the images to process The program searches the directories in <code>dirList</code> , as well as all subdirectories, for the files to analyze. <i>Must be a cell array of strings.</i>	
<code>fileOut</code>	Name of the file to store the data. Must <i>not</i> have an extension.	
GENERAL ARGUMENTS		
<code>taskList</code>	Tasks MTQuant will complete. Add the values of the various tasks to complete in order to perform more than one task at a time. 1. 1 to make stacks into projections 2. 2 to make masks of projections 3. 4 to generate line scans 4. 8 to analyze line scans	15
<code>verbose</code>	How often to display which file is being processed 2 = every new file 1 = every 50 files 0 = never	1
<code>indivFileOut</code>	String to prepend to the outputs of MTQuant, such as 'axon'	'
ARGUMENTS FOR TASK 1: GENERATE MAXIMUM PROJECTIONS		
<code>gFileExt</code>	Ending of green stack files to search for	'*_w1488-single.TIF'
<code>rFileExt</code>	Ending of red stack files to search for	*'_w2561-single.TIF'
ARGUMENTS FOR TASK 2: TRACE NEURITE MASKS		
<code>diPix</code>	Number of pixels by which to dilate the user-selected line when creating the neurite mask	30
ARGUMENTS FOR TASK 3: GENERATE LINE SCANS		

Input Argument	Description	Default Value
toAvg	0 = sum together intensities along perpendiculars to the neuron's spline 1 = average together intensities instead (NOT recommended)	0
ARGUMENTS FOR TASK 4: CALCULATE ORG PARAMS (MT MINUS-END DETECTION)		
toUseManualRPeaks	0 = detect MT minus-ends automatically 1 = User is asked to enter number of MT minus-ends for each image	0
toUseRandRPeaks	0 = No correction to count single red puncta as multiple MT minus-ends 1 = Randomly count some single red puncta as multiple MT minus-ends	1
rPeakTol	Tolerance of peak selection. This value is multiplied by the maximum value of the red line scan. For a peak to be considered real (not noise), the difference between the peak and the nearest valley must be at least $\text{rPeakTol} \times \text{max}(\text{rLineScan})$. A higher value for rPeakTol means fewer peaks are identified.	0.05
rPeakCorr	Weight on Bernoulli random variable that decided if a single red dot is 1 or 2 MT minus-ends. As coverage increases, this value should increase as well (up to 20 if necessary)	0.1
ARGUMENTS FOR TASK 4: CALCULATE ORG PARAMS (SINGLE MT BRIGHTNESS EXTRACTION)		
toPreproc	0 = Use the green line scan directly 1 = Smooth and sharpen the green line scan before extracting single MT brightness	0
toUseBlurCorr	0 = Don't correct for overlapping MT starts and ends (not recommended) 1 = Correct for overlapping MT starts and ends	1
roundLevel	Rounding level for quantized signal. A fractional number of MTs below roundLevel is rounded down when forming the quantized signal, while a fractional number above roundLevel is rounded up.	0.5
distThresh	When calculating single MT brightness, this value is the number of allowable pixels between a step up in the quantized signal and a MT minus-end before the MT minus-end is classified as a mislocalization	3
alpha1	Weight on Mean Squared Error Cost	$\frac{0.26}{\text{mean}(\text{gLineScan})}$
alpha2	Weight on Cost of mislocalized steps up	0.0013
alpha3	Weight on Error in number of steps up	1

Input Argument	Description	Default Value
ARGUMENTS FOR TASK 4: CALCULATE ORG PARAMS (MISCELLANEOUS)		
toRateImgs	<p>0 = do not rate images 1 = rate images After each image's line scan has been extracted, the user is asked to rate the image: 2 = good image 1 = ok image 0 = bad image Then the program outputs a set of CSV files for each of the following combinations of images: good images only (2), good and ok images (1 & 2), and all images (0, 1, 2) If toRateImgs is false, then all images are given the rating of 2.</p>	0
toUseHalfMTs	<p>How to handle the number of MTs at the beginning and end of the signal when calculating length. Makes very inconsequential changes; can be ignored. 0 = no correction (Use the number of minus-ends detected, called numMTs) 1 = Use numMTs - (value of quantized signal at end of line scan) + (average value of quantized signal at beginning and end of line scan) 2 = Use numMTs + (value of quantized signal at end of line scan)</p>	0
toEM	<p>0 = do not implement distribution refinement 1 = implement distribution refinement</p>	1

Table 1: Input Arguments to MTQuant

5.2 Output Table Column Names

#	Output Argument	Description	Units/ Possible Values
IMAGE INFORMATION			
1	Directory	Directory containing the processed image	String
2	DataFile	Name of the line scan file that was processed to generate this row of data	String
3	DirNum	The group number of the line scan file. If the <code>dirList</code> sent to <code>MTQuant</code> had one entry, then each sub-folder inside of <code>dirList</code> is considered a separate group, and receives a distinct <code>DirNum</code> . Otherwise, each directory in <code>dirList</code> is considered a distinct group.	Integer
4	Rating	If the argument <code>toRateImgs</code> is false (as it is by default), this column is simply 2 for every image. However, if <code>toRateImgs</code> is true, then this column contains the user's rating: 2 = good image 1 = ok image 0 = bad image	0, 1, or 2
5	Max_G	Maximum intensity value of the green line scan	Intensity (AU)
6	Avg_G	Average intensity value of the green line scan	Intensity (AU)
7	Scan_Length	Length of the green line scan	Pixels
8	Num_MTs	Number of MT minus-ends detected in the red line scan	Integer
ORGANIZATION PARAMETERS			
9	Avg_Spacing	Average Spacing of MT minus-ends in the line scan	Pixels
10	Std_Dev_Spacing	Standard Deviation of Spacing of MT minus-ends in the line scan	Pixels
11	Single_MT_Brightness	Extracted Single MT Brightness	Intensity (AU)
12	Avg_Coverage	Average Coverage (number of MTs in neuron cross-section)	Number of MTs
13	Std_Dev_Coverage	Standard Deviation of Coverage	Number of MTs
14	Avg_Length	Average MT Length	Pixels

Table 2: Outputs of MTQuant

In the '`*_grouped.csv`' files, the data in each column of the output data has been consolidated by the group numbers (`DirNum`). For each group identified as described in Section 4, the mean and standard deviation are calculated for each of the parameters in Table 2. The names of the columns are the same in the '`*_grouped.csv`' files as for the non-grouped data, but a '`M_`' or '`S_`' is prepended to the name in the '`*_grouped.csv`' files, representing the mean and standard deviation, respectively.

5.3 Input and Output Filenames for Each Task

File Suffixes	Description	Output of Task...	Input to Task...
'*_w1488-single.TIF' gFileExt	Green stack file	-	1
'*_w1561-single.TIF' rFileExt	Red stack file	-	1
'*_g_proj.tif'	Green maximum projection file	1	2,3
'*_r_proj.tif'	Red maximum projection file	1	2,3
'*_stackInds.csv'	Stack slice containing green maximum projection values	1	-
'*_Mask.csv' [indivFileOut, '*_Mask.csv']	Mask info CSV file	2	3
'*_Mask.tif' [indivFileOut, '*_Mask.tif']	Mask image file	2	3
'*_LineScans.csv' [indivFileOut, '*_LineScans.csv']	Line scans	3	4
'*_LineScanInds.csv' [indivFileOut, '*_LineScanInds.csv']	Pixel locations that were summed to generate the line scans	3	4
[fileOut, '_2.csv']	All data for images classified as "good" (all images by default)	4	-
[fileOut, '_2_grouped.csv']	Grouped data for images classified as "good" (all images by default)	4	-
[fileOut, '_12.csv']	All data for images classified as "good" or "ok" (Only if toRateImgs is 1)	4	-
[fileOut, '_12_grouped.csv']	Grouped data for images classified as "good" or "ok" (Only if toRateImgs is 1)	4	-
[fileOut, '_012.csv']	All data for all images (Only if toRateImgs is 1)	4	-
[fileOut, '_012_grouped.csv']	Grouped data for all images (Only if toRateImgs is 1)	4	-

Table 3: File Naming Convention. *=wild card