

ΒΙΟΛ-491 Ειδικά Κεφάλαια Βιοτεχνολογίας και Απεικονιστικής Φυτών



<https://www.imbb.forth.gr/en/research-en/plant-molecular-biology/item/4119-panagiotis-n-moschou>

<https://pmoschoulab.blog/home/>



FORTH

INSTITUTE OF MOLECULAR BIOLOGY & BIOTECHNOLOGY

<https://twitter.com/Panosmoschou>



Χρήση FIJI

Νέες δεξιότητες-Διάλεξη 3

➤ Χρήση Fiji

Τι είναι το FIJI

- Το ImageJ και το FIJI είναι ελεύθερα λογισμικά που χρησιμοποιούνται για την ανάλυση εικόνας στις επιστήμες ζωής *used for image analysis in life sciences*

Collaboration

The Fiji project is driven by a strong desire to improve the tools available for life sciences to process and analyze data. To this end, Fiji collaborates closely with the following projects:



ImageJ2



Bio-Formats



OME



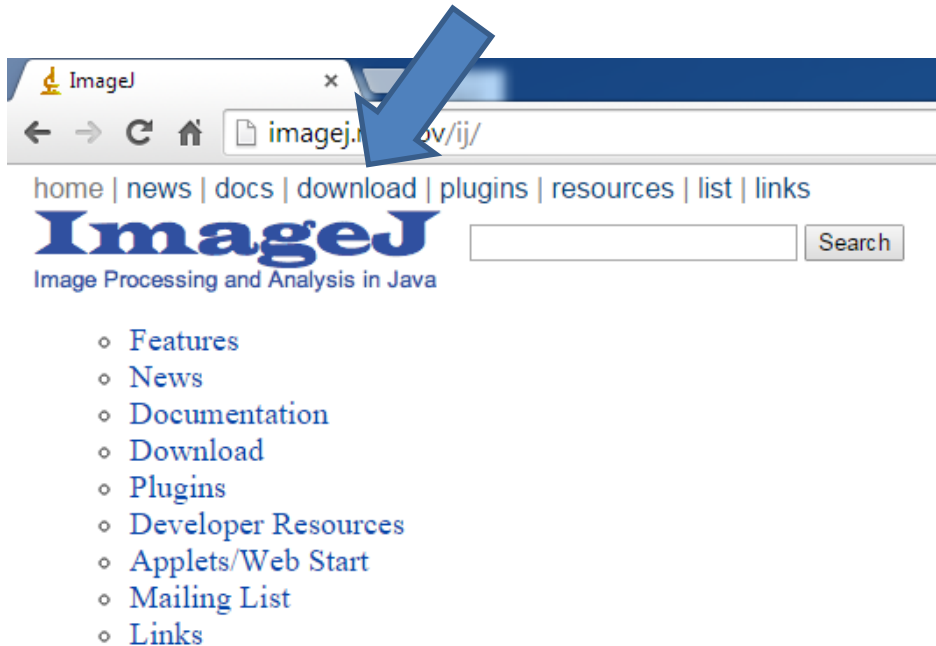
µManager



KNIME

Πώς ξεκινάμε

Step 1



This page has been visited **11,177,869** times. Send comments to wsr@nih.gov. [Disclaimer](#)

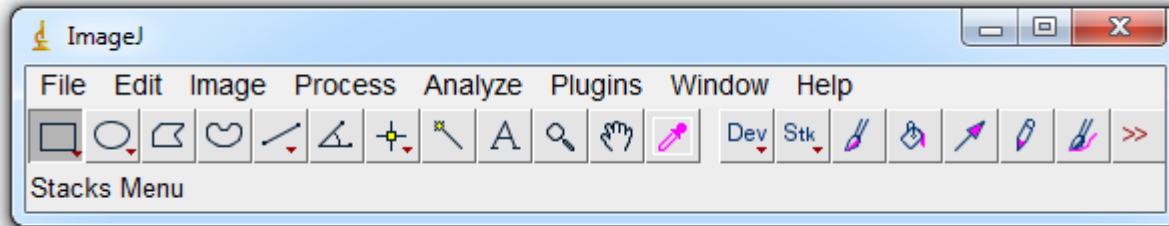
Step 2



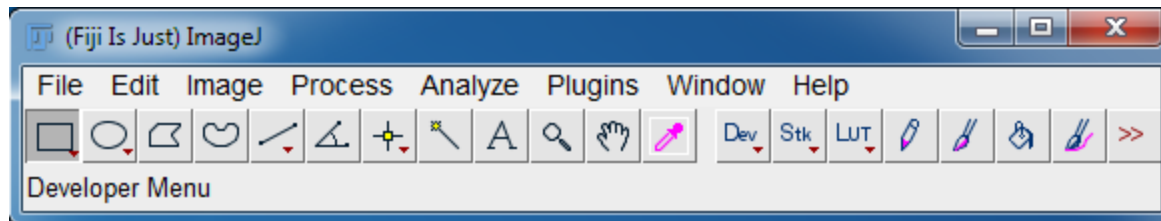
You can also browse the ImageJ download directory at imagej.nih.gov/ij/download/. Refer to the [release notes](#) for a list of new features and bug fixes.

[top](#) | [home](#) | [news](#) | [docs](#) | [download](#) | [plugins](#) | [resources](#) | [list](#) | [links](#)

Πώς μοιάζει το περιβάλλον του



OR?



Έστω ότι έχουμε μια εικόνα χρονικής αλληλουχίας, τι κάνουμε μετά;

- Έλεγχος εικόνας (ανάλυση 512x512 dpi κατά τη λήψη; Δες και French et al., 2008)
- Χρειάζεται να κάνουμε ανάλυση;

Πώς μπορώ να βρω το μέγεθος της εικόνας μου;

The image shows the ZEN 2012 software interface. On the left, the 'Method' dropdown menu is open, showing various processing options. The 'Info' tab is selected in the left sidebar, and a blue arrow points to it. The main panel displays the 'Info' tab for the file 'TSN2.TIP8 30 min 5.avi'. The 'Dimensions' field is highlighted with a red box, showing 'x: 463, y: 512, time: 18, channels: 2, 8-bit'. A calculator window is open in the center, displaying the calculation $0.227 \times 463 = 105.101$. The status bar at the bottom shows CPU usage at 0%, Free HD at 0.33 TB, and Free Ram at 2.4 GB.

Method

- Maximum intensity projection
- Color-coded projection
- Image calculator
- Average
- Filter
- Linear unmixing
- Ion Concentration
- Correlation
- Modify Series
- HDR- imaging
- Channel Alignment
- Copy

Method Parameters

Input image:

Info

Name: TSN2.TIP8 30 min 5

Description:

Notes:

User: eogu0001

Scaling X: 0.227 μm

Scaling Y: 0.227 μm

Scaling Z:

Dimensions: **x: 463, y: 512, time: 18, channels: 2, 8-bit**

Image size: x: 104.99 μm , y: 116.12 μm

Scan Mode: plane, time series

Zoom: 3.7

Objective: Plan-Apochromat 20x/0.8 M27

Pixel dwell: 1.75 μs

Average: line 4

Master gain: Ch1 : 928
Ch2 : 535

Digital gain: 1.00

Digital offset: 0.00

Pinhole: 250 μm

Filters: Ch1 : 431 - 504
Ch2 : 529 - 595

Beam splitters: MBS : MBS 458/514
MBS_InVis : MBS -405

Lasers: 458 nm : 5.0 %
514 nm : 0.3000 %

Calculator

View Edit Help

0.227 *
463

MC MR MS M+ M-
← CE C ± √
7 8 9 / %
4 5 6 * 1/x
1 2 3 - =
0 . +

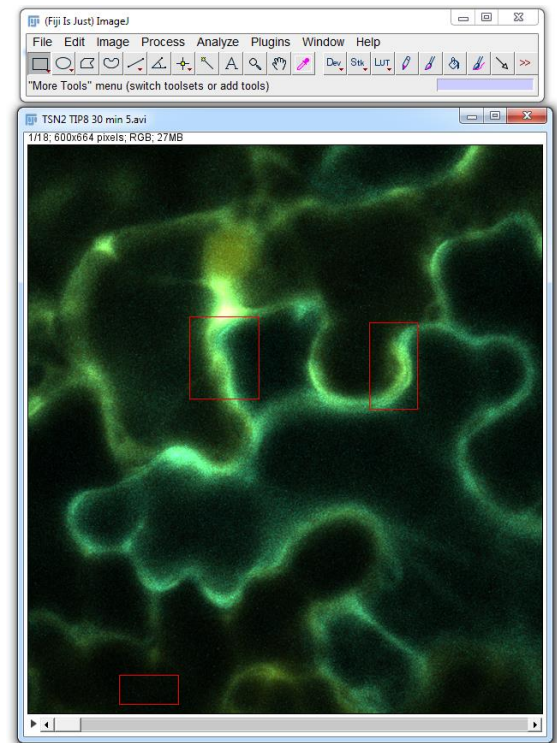
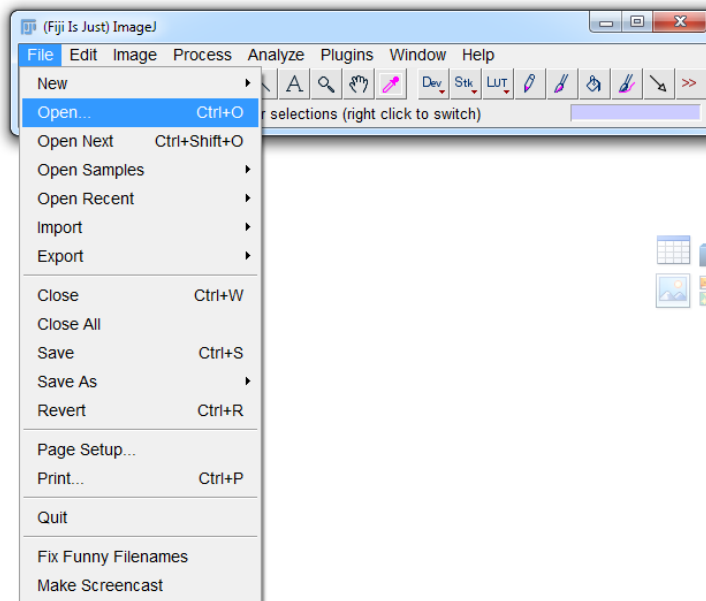
Images and Documents

- 3 b.lsm *****
0.25 MB
- 3 b.lsm *****
0.25 MB
- TSN2 ...n 5.avi *****
8.1 MB

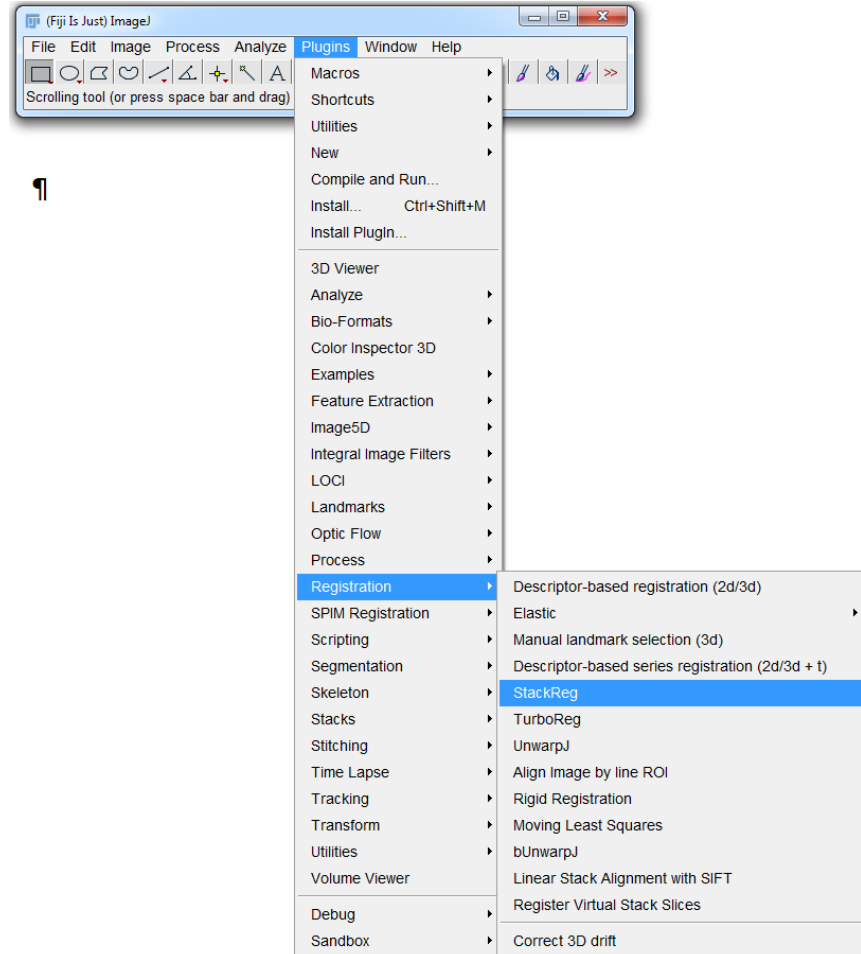
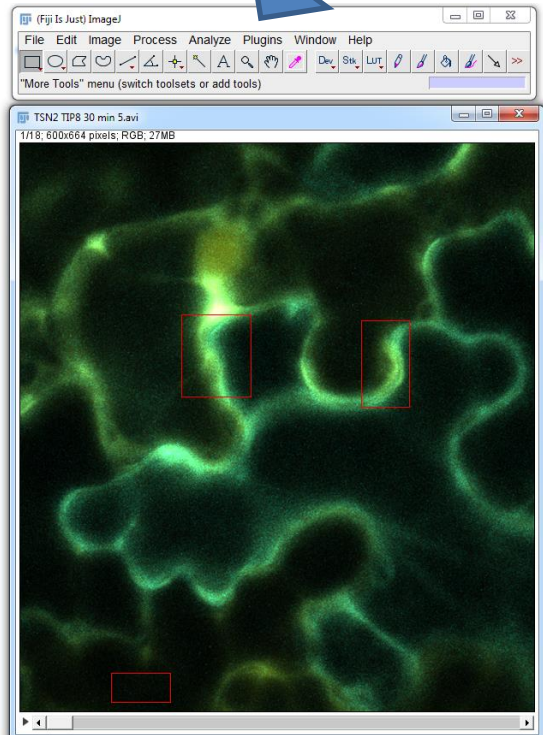
CPU 0% Free HD 0.33 TB Free Ram 2.4 GB

Άνοιξε το αρχείο με το FIJI

- Το FIJI ανοίγει πολλούς τύπους αρχείων!!!!



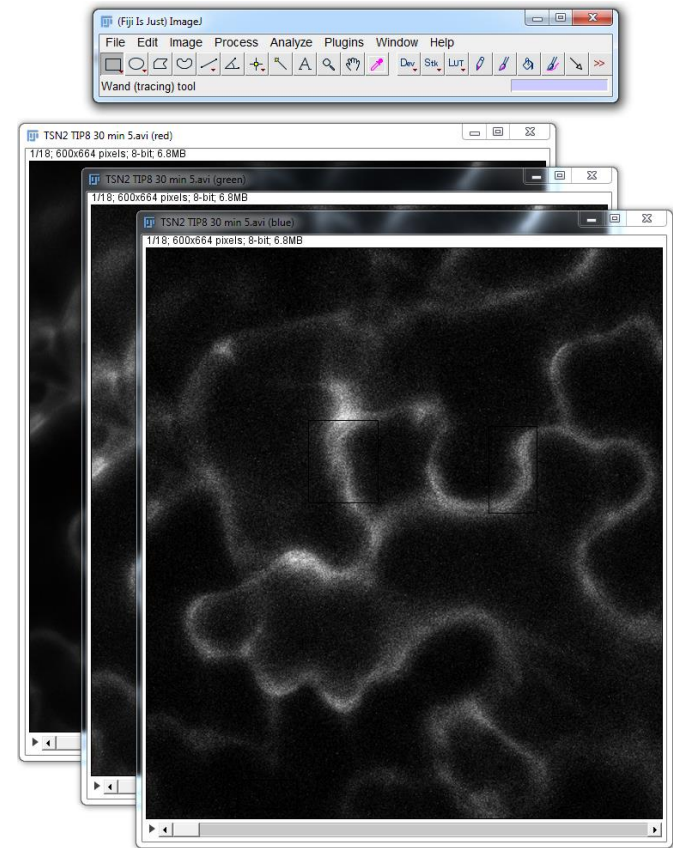
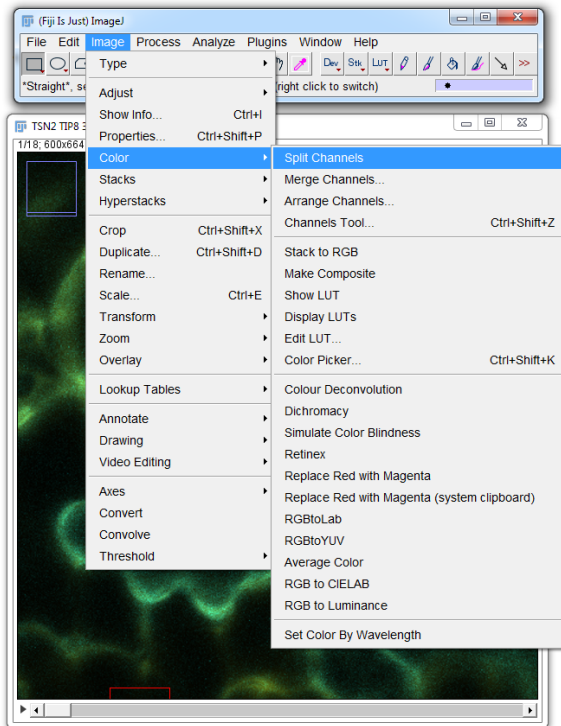
Η εικόνα μου κουνιέται, τι μπορώ να κάνω;



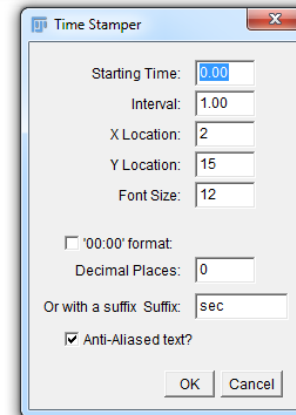
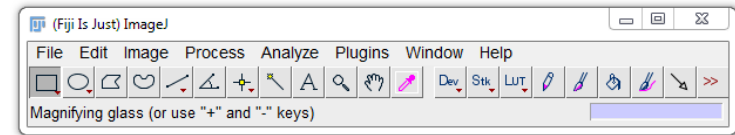
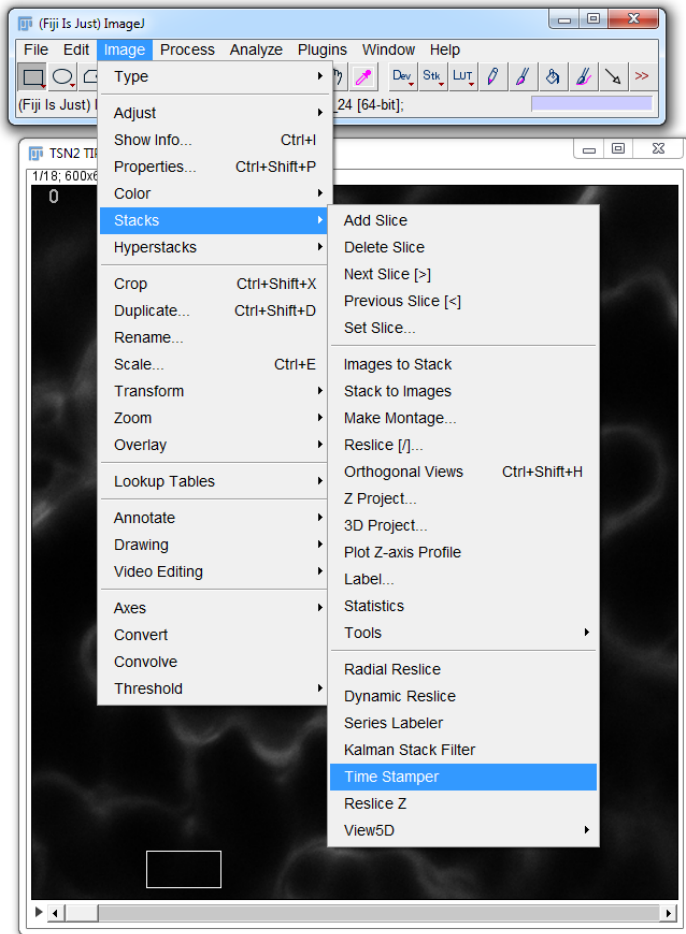
How does it work?

- [We are using Rigid body transformations](#)
- <http://bigwww.epfl.ch/thevenaz/stackreg/>

Αν έχεις διάφορα κανάλια μπορείς να τα διαχωρίσεις



Μπορούμε να βάλουμε χρονικές ενδείξεις

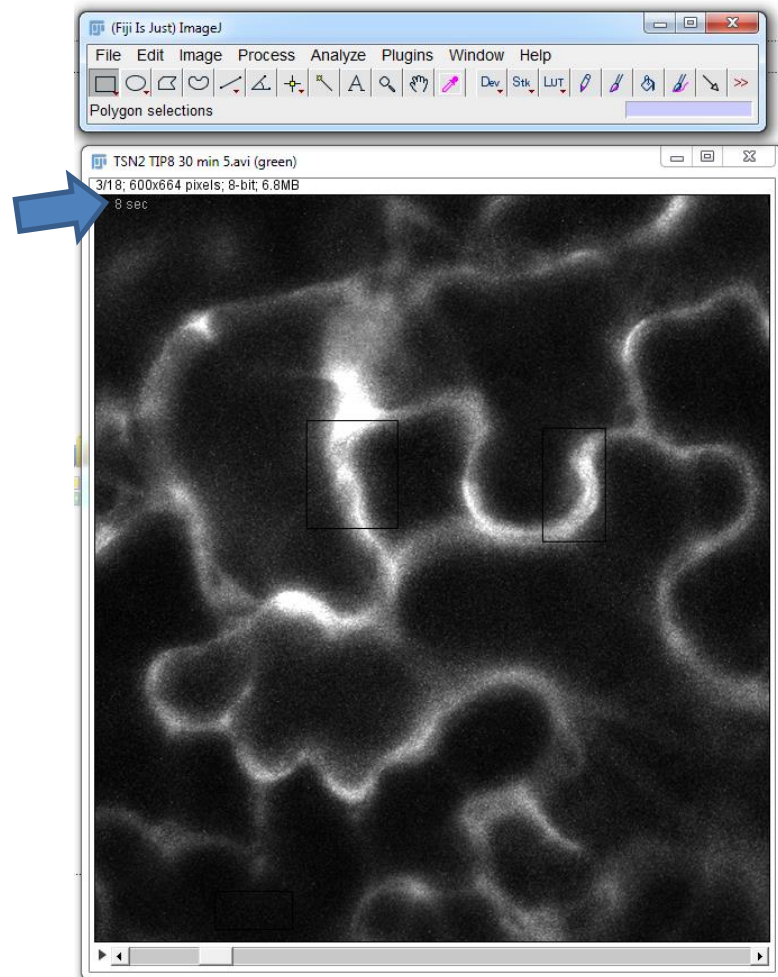
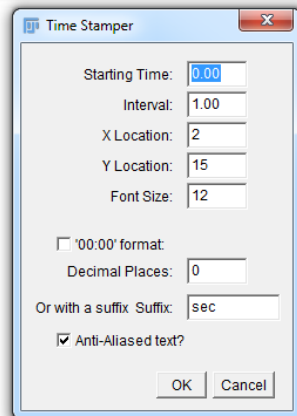
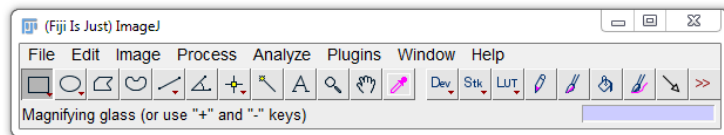


Υπολογισμός ενδιάμεσου χρόνου

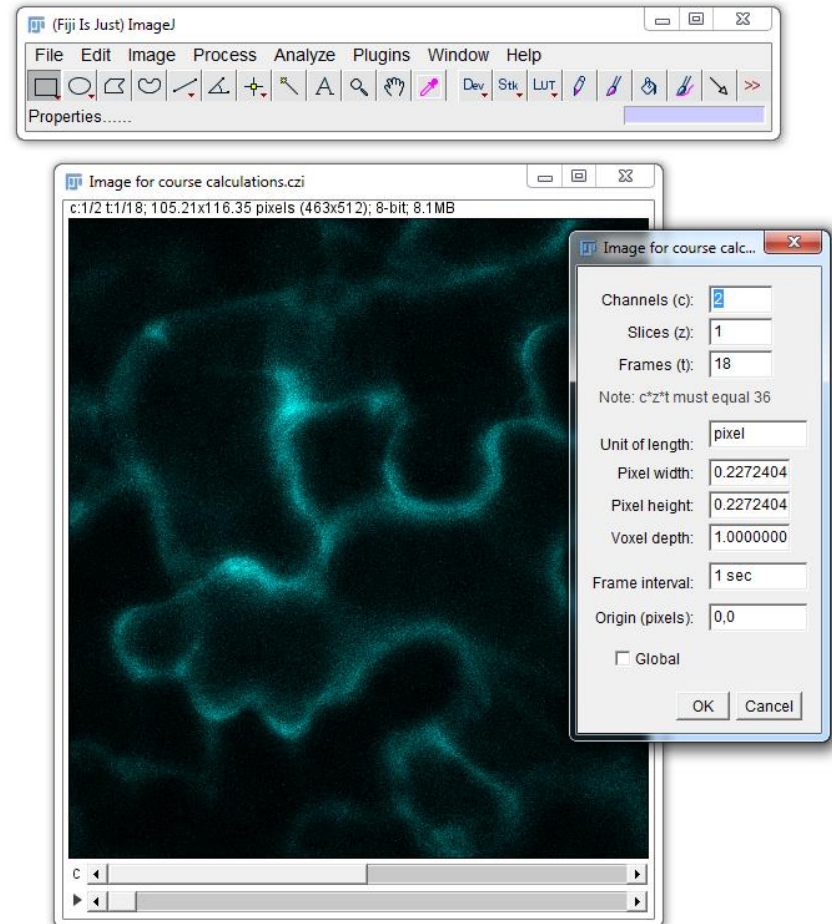
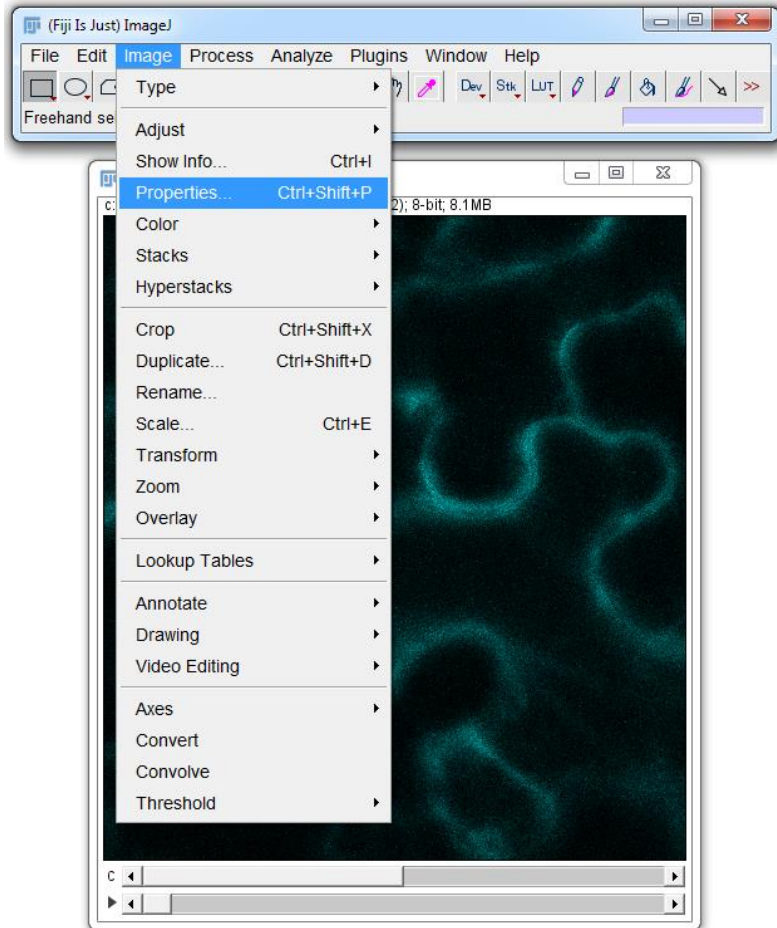
The screenshot displays the Zen 2012 software interface. The main window shows a fluorescence image with two red rectangular boxes highlighting specific regions. A Windows Calculator window is open over the image, showing the calculation $30.6 - 28.6$. The interface includes several panels:

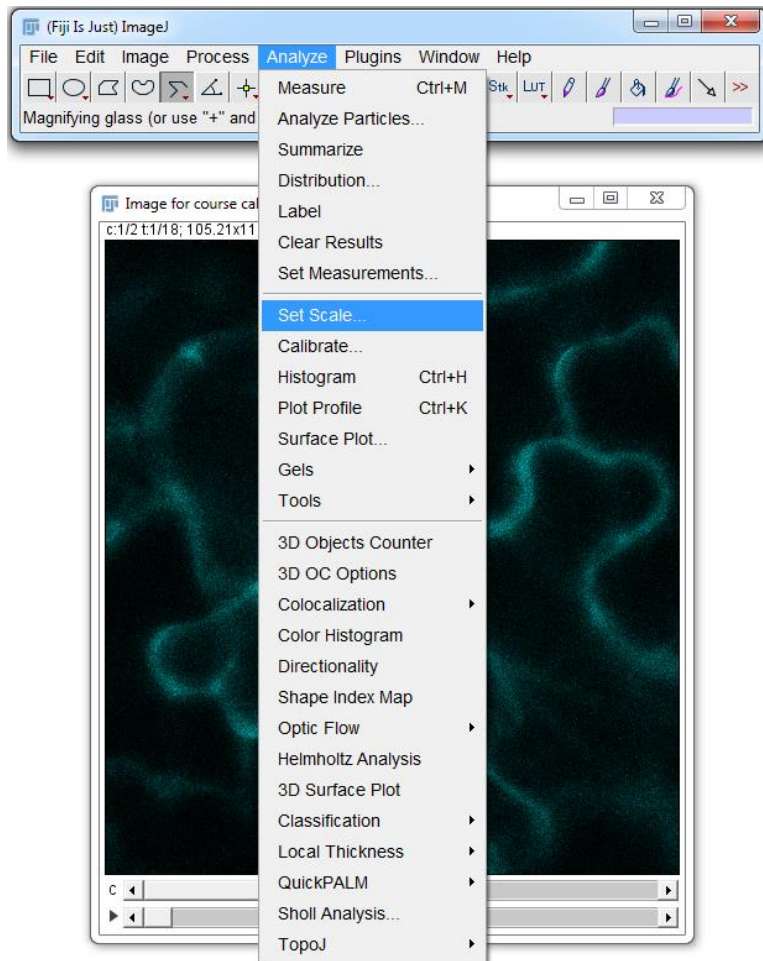
- Left Panel:** Contains the 'Method' dropdown menu with options like 'Maximum intensity projection', 'Color-coded projection', 'Image calculator', 'Average', 'Filter', 'Linear unmixing', 'Ion Concentration', 'Correlation', 'Modify Series', 'HDR- Imaging', 'Channel Alignment', and 'Copy'. Below it is the 'Method Parameters' section with an 'Input Image' field and a 'Select' button.
- Top Panel:** Shows the 'Workspace Zoom' and 'Workspace Configuration' controls.
- Right Panel:** Displays the 'Images and Documents' list, showing files like '3 b.lsm' (0.25 MB), '3 b.lsm' (0.25 MB), and 'Imag...ons.czi' (8.1 MB).
- Bottom Panel:** Contains the 'Dimensions' and 'Graphics' sections. The 'Dimensions' section shows 'Time' (1 to 18) and 'Zoom' (100% to 130). The 'Graphics' section shows 'Ch1' and 'Ch2' channels, 'Single Channel' and 'Range Indicator' checkboxes, and a 'Display' section with a graph showing a curve and a line.

The status bar at the bottom indicates system information: CPU 0%, Free Ram 1.6 GB, Free HD 0.33 TB, and Resolution 1920x1080.

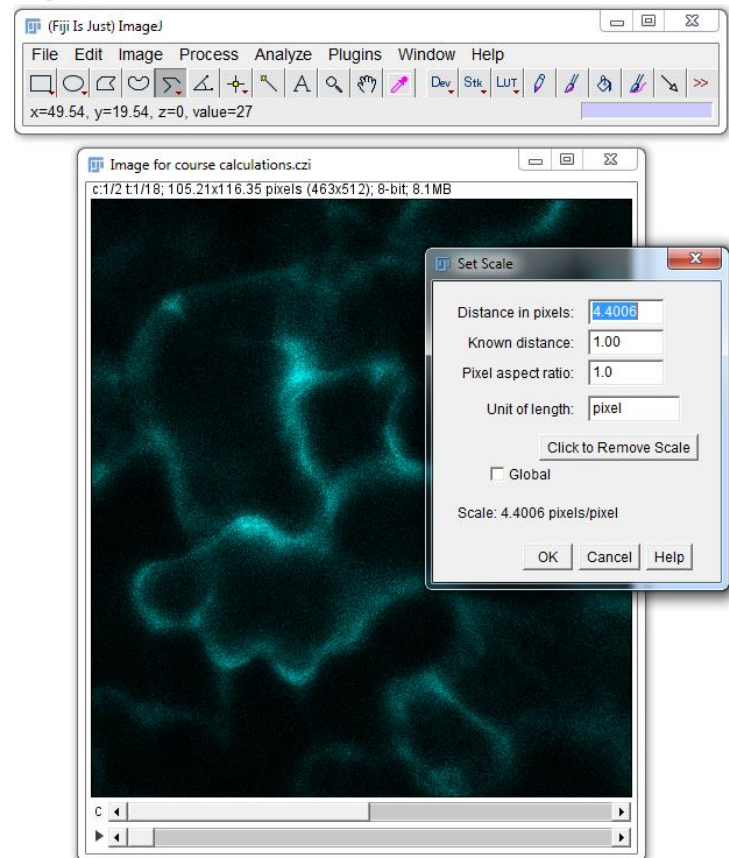


Πώς βάζω κλίμακα;

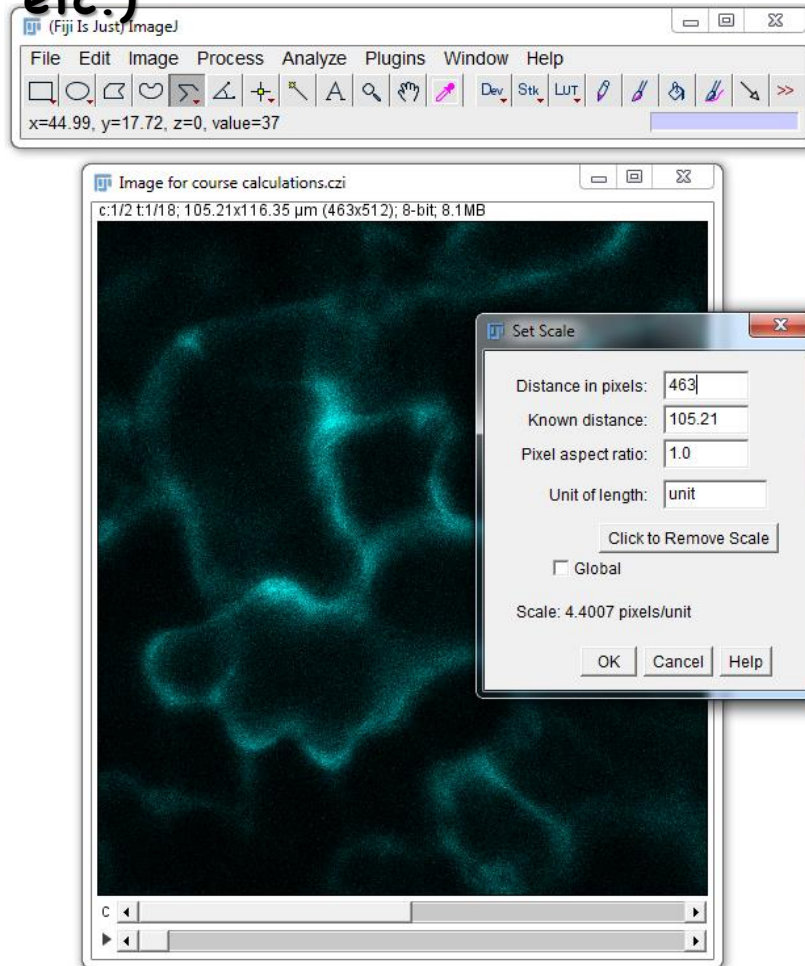




1. Εάν έχεις το αυθεντικό αρχείο



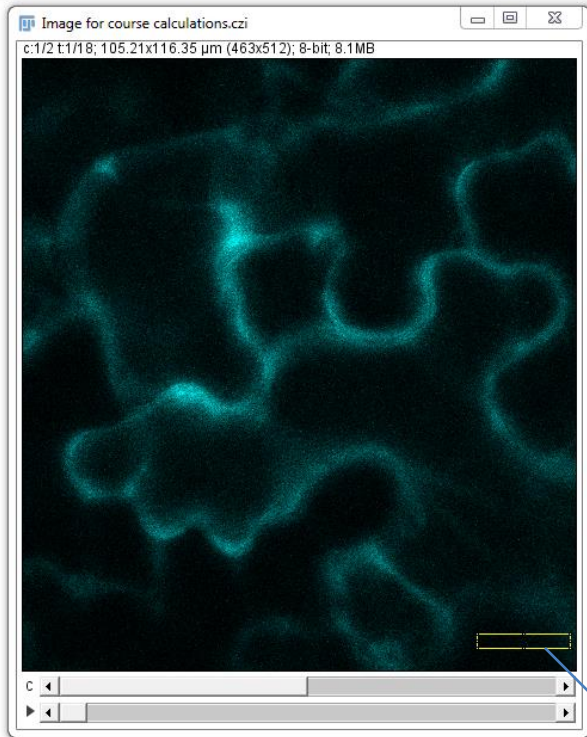
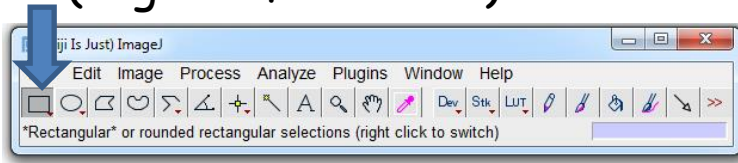
2. Εάν δεν έχεις το αρχείο στην αρχική του μορφή (π.χ. 'avi' files etc.)



Πώς βάζω τη μπάρα

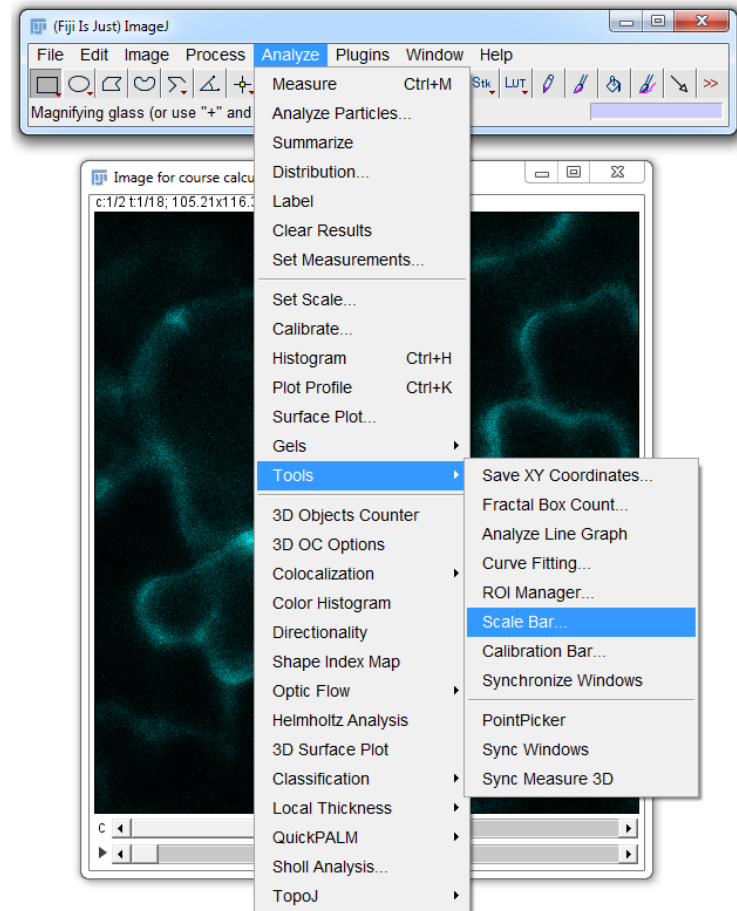
Step 1: πού θα μπει

ROI (region of interest)

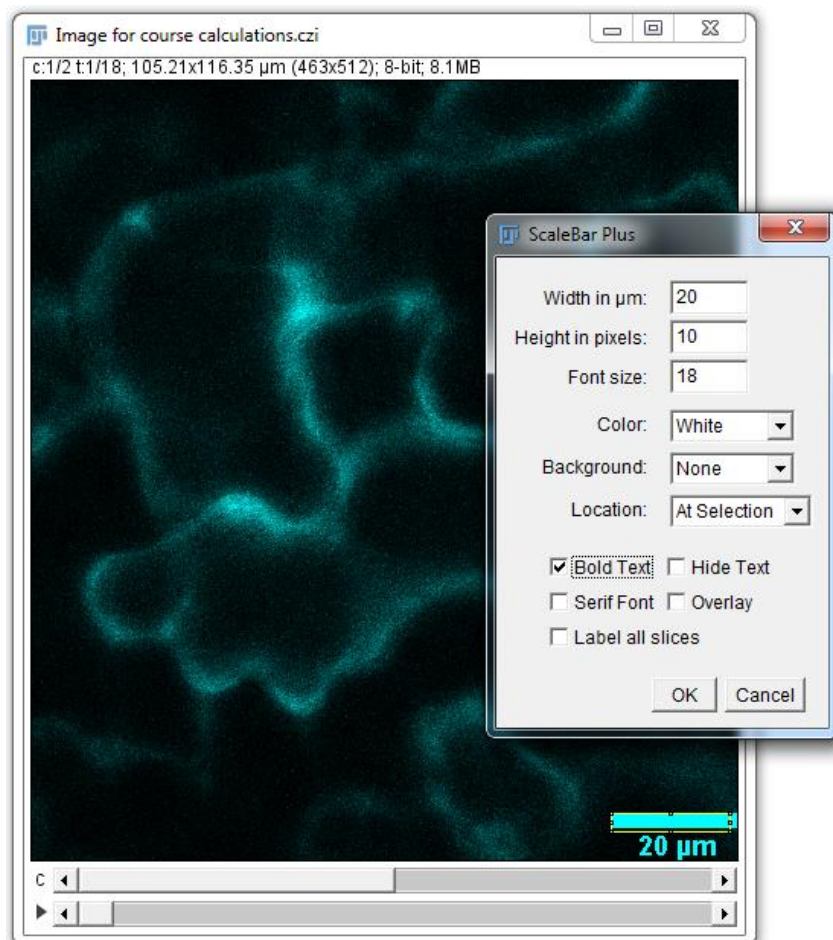


Εδώ

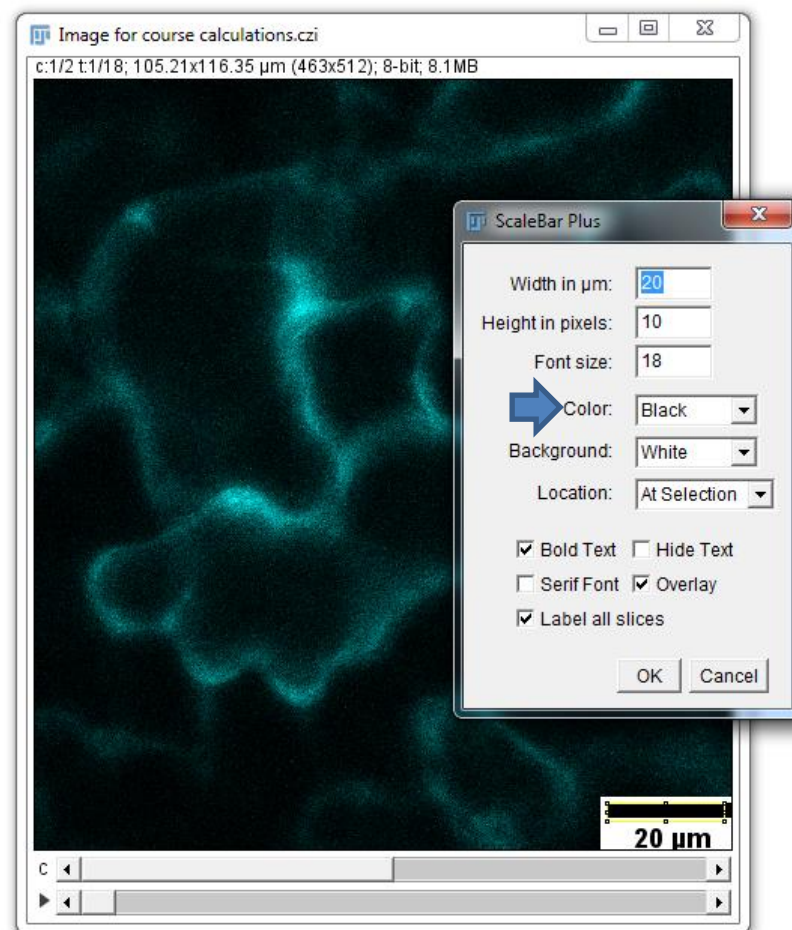
Step 2: κάνε τα εξής



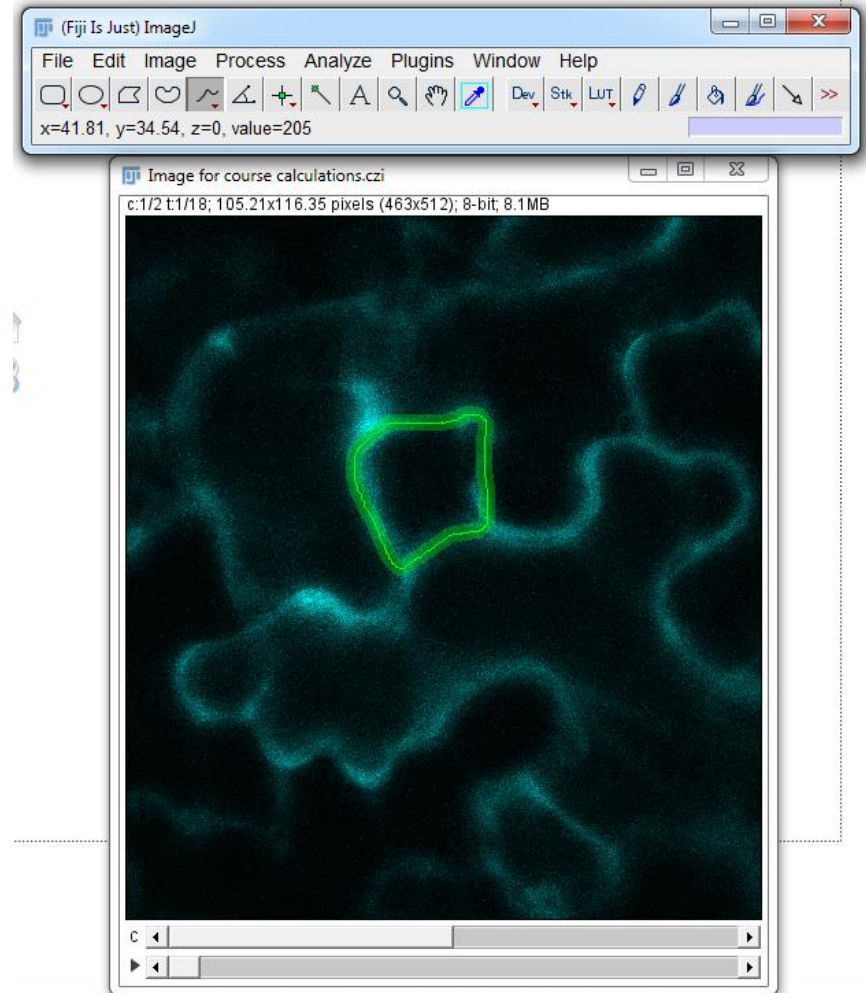
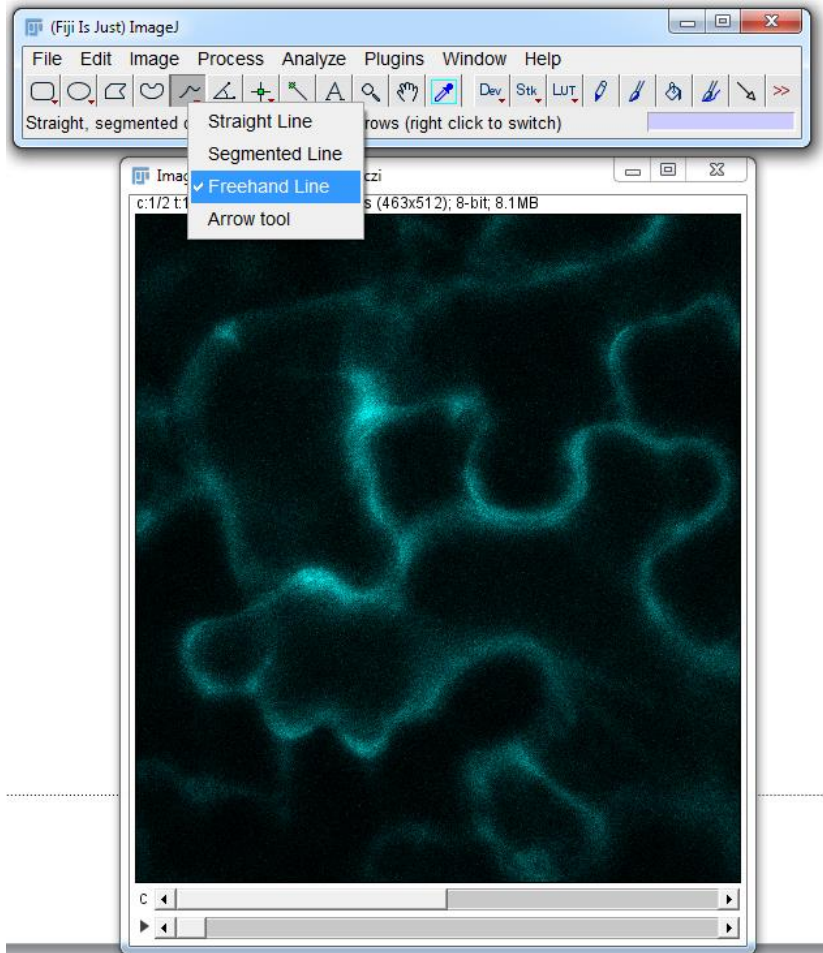
Step 3:

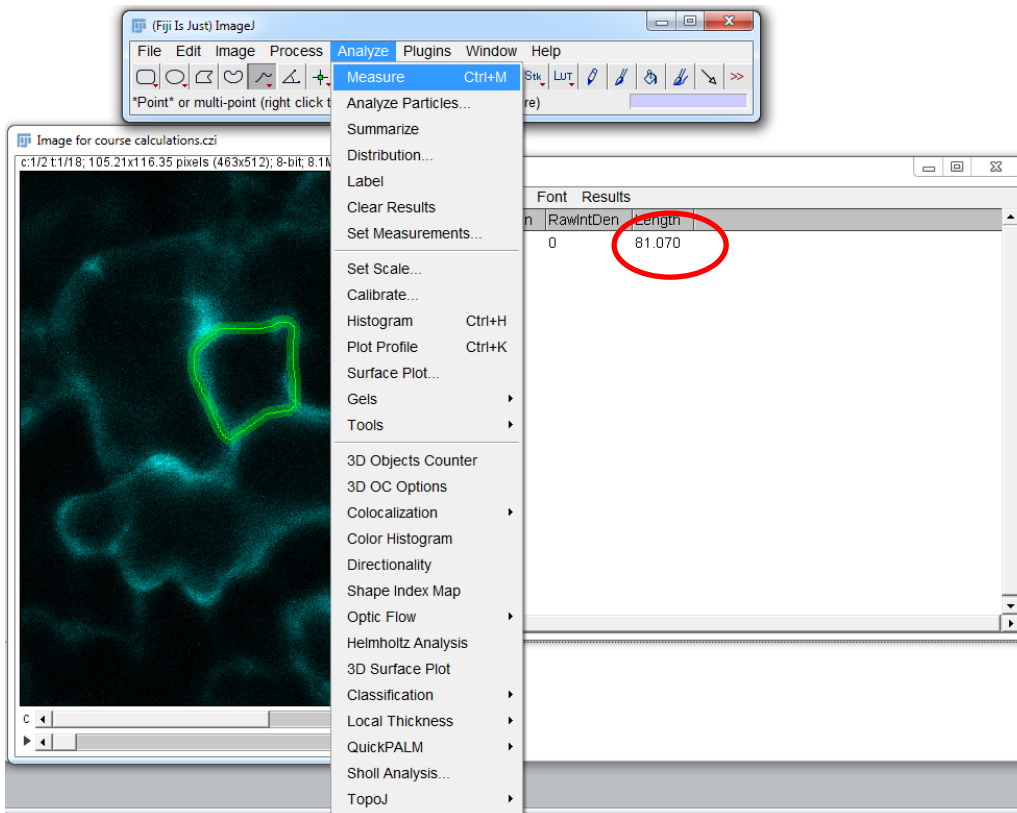


Step 4: Χρώμα κτλ.

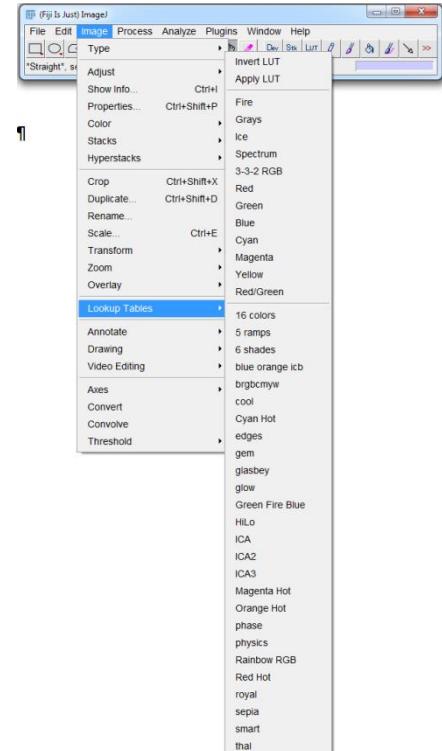
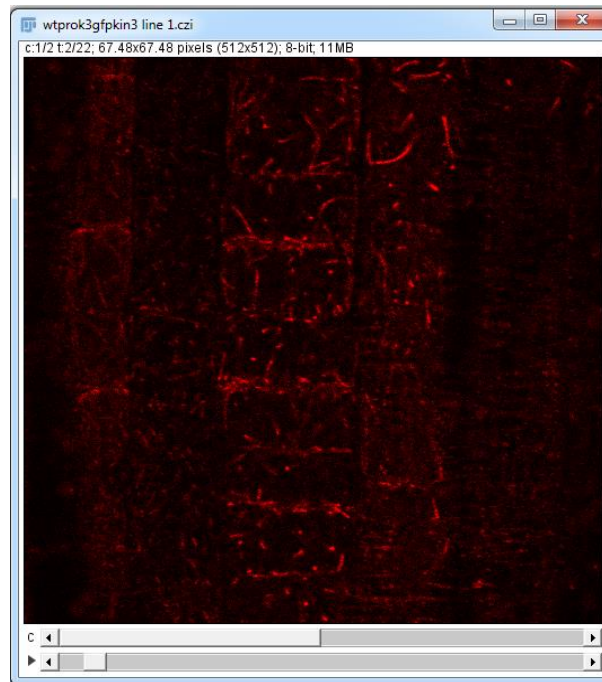
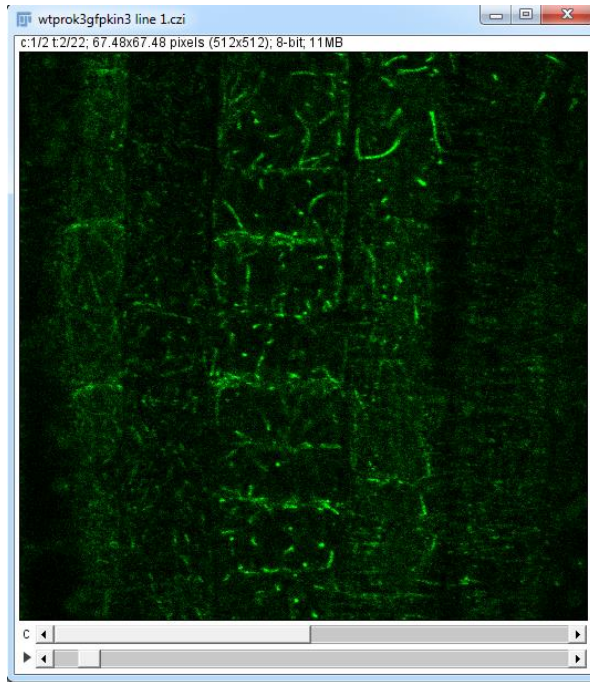


Τώρα μπορώ να μετρήσω αποστάσεις

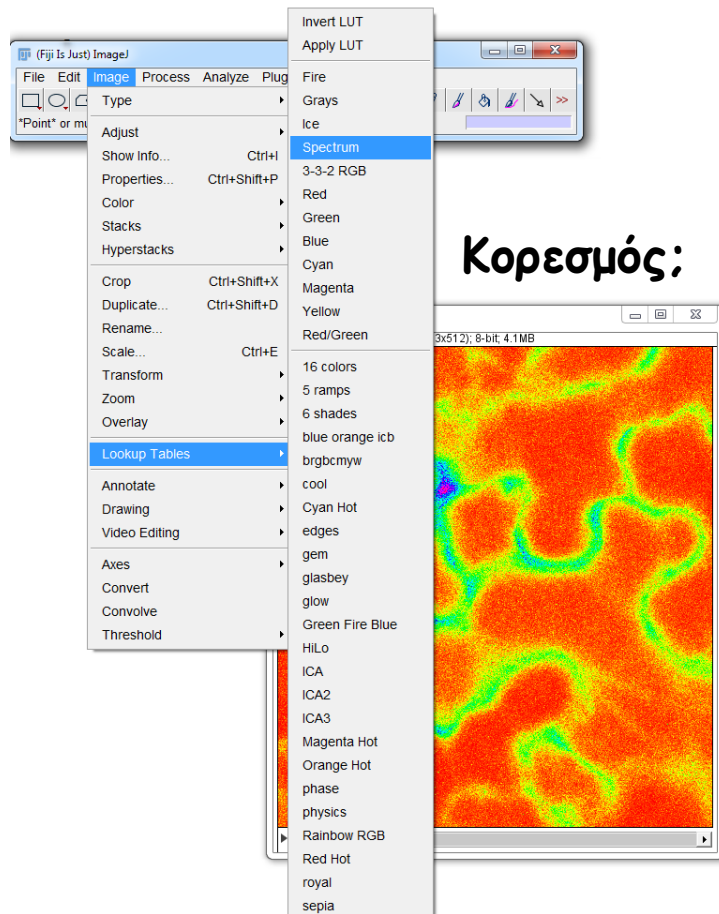




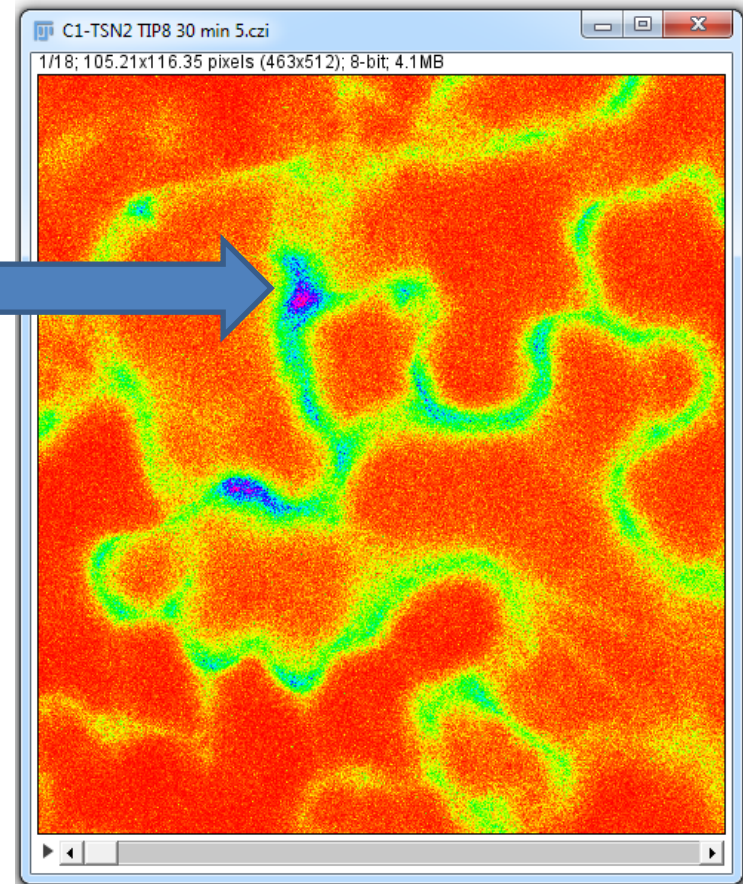
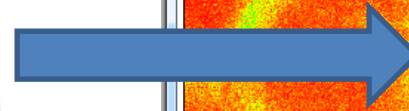
Πώς μπορώ να προσαρμόσω τα χρώματα;



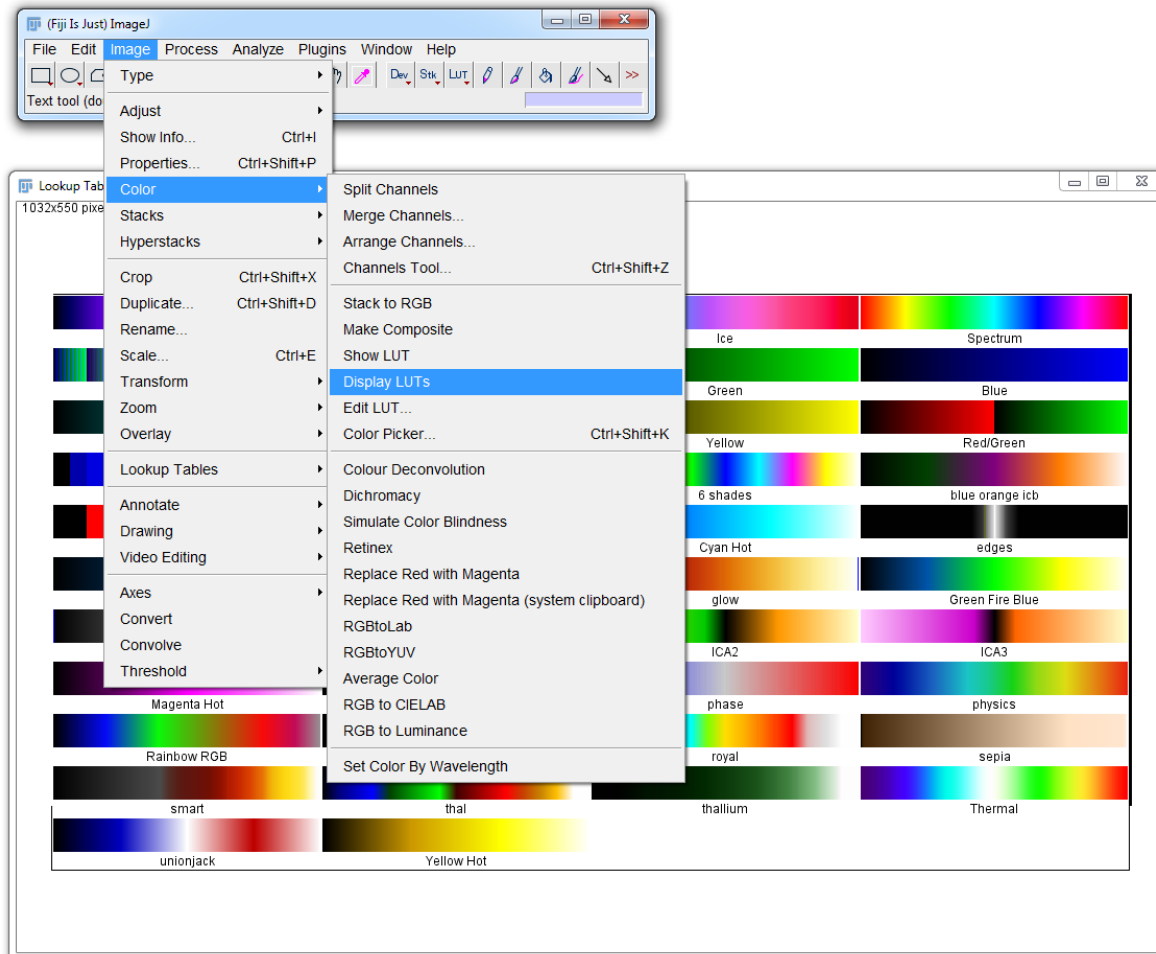
Λύση: αποχρώσεις του γκρι



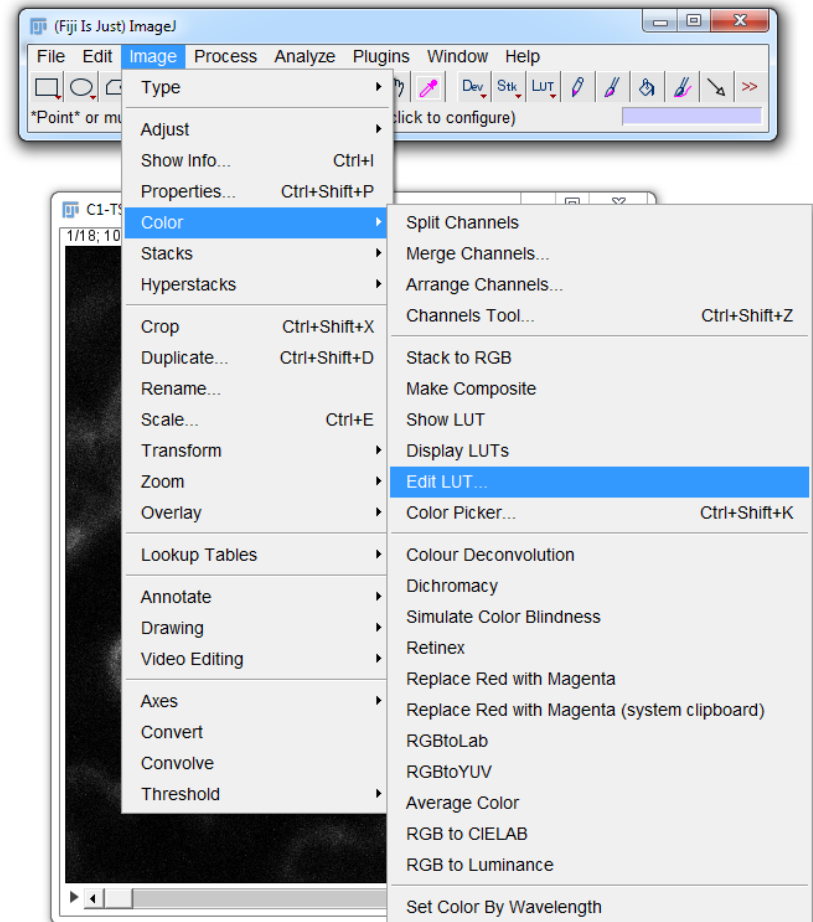
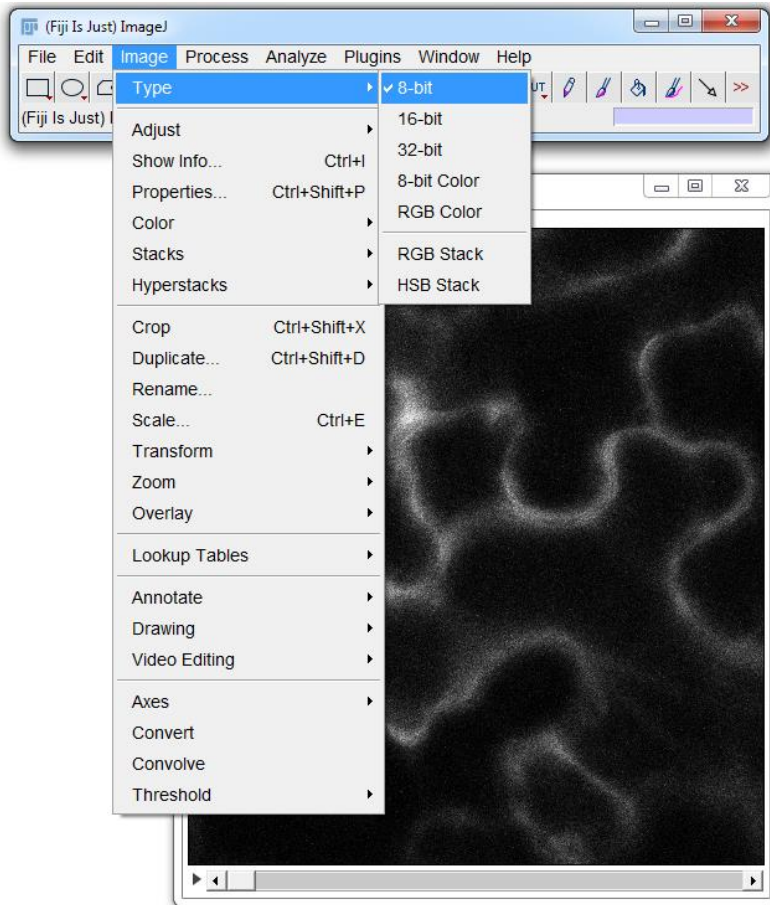
Κορεσμός:



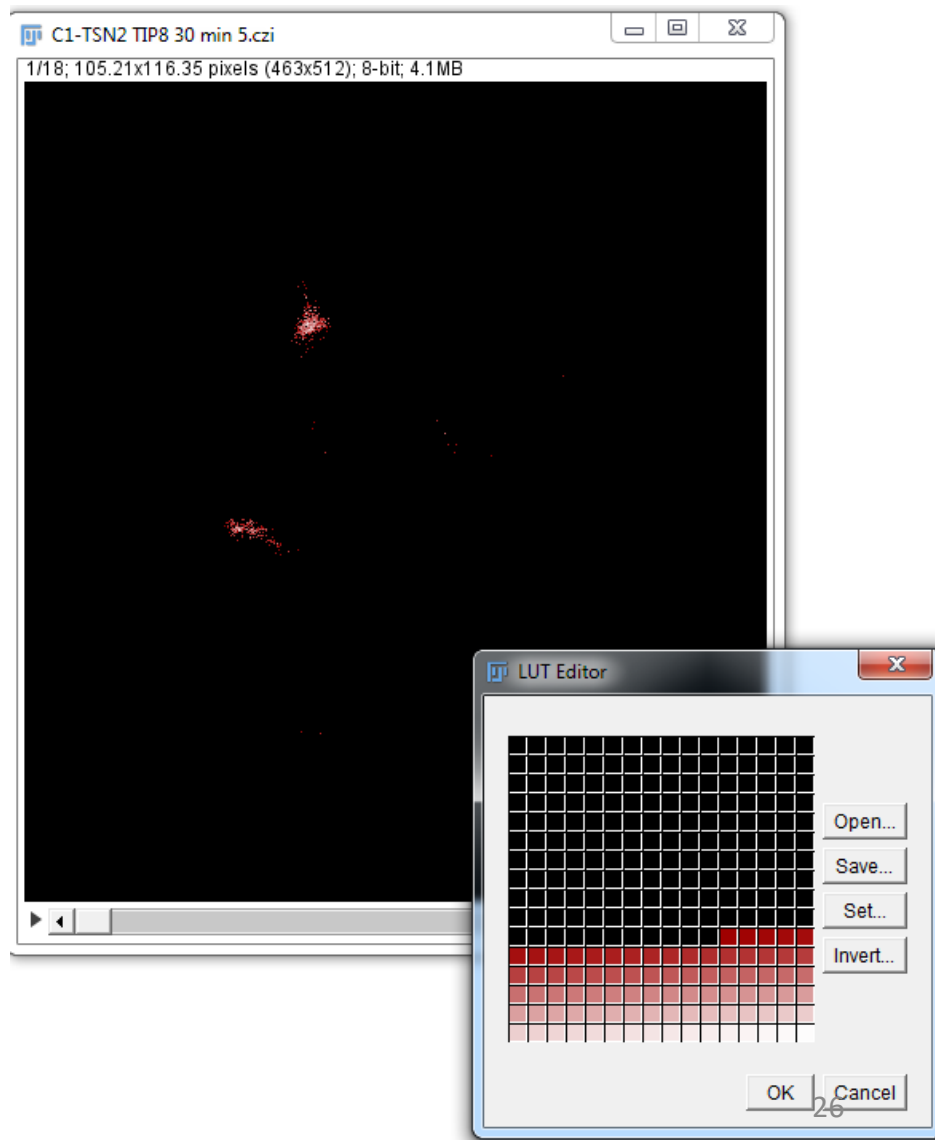
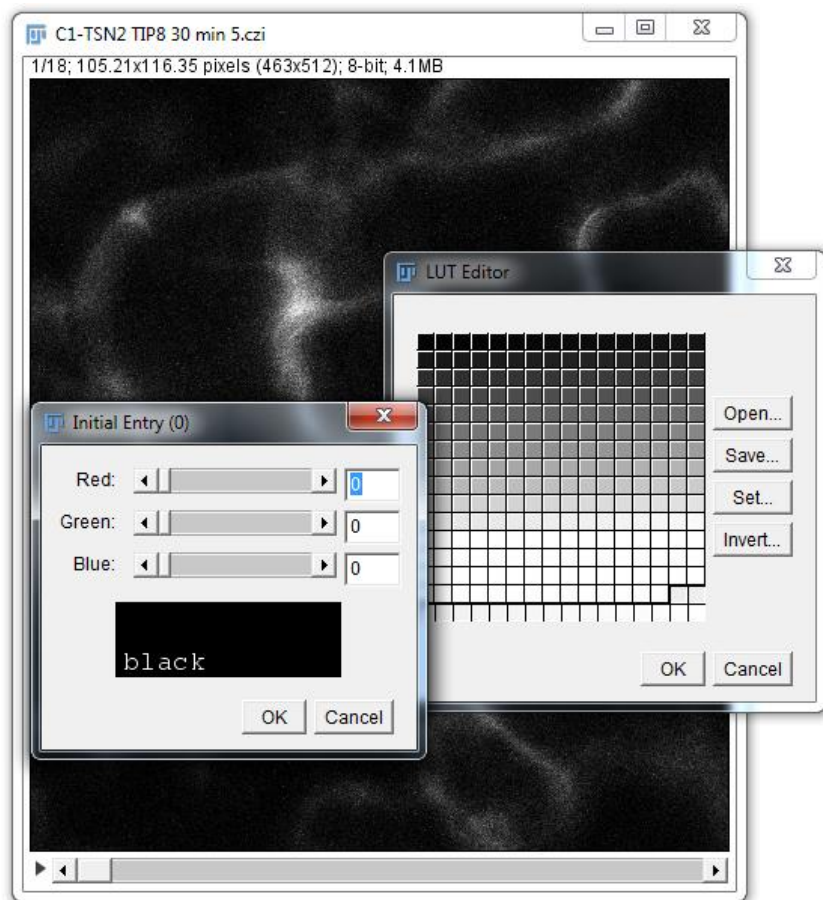
Τί είναι το Look Up Table (LUT)?

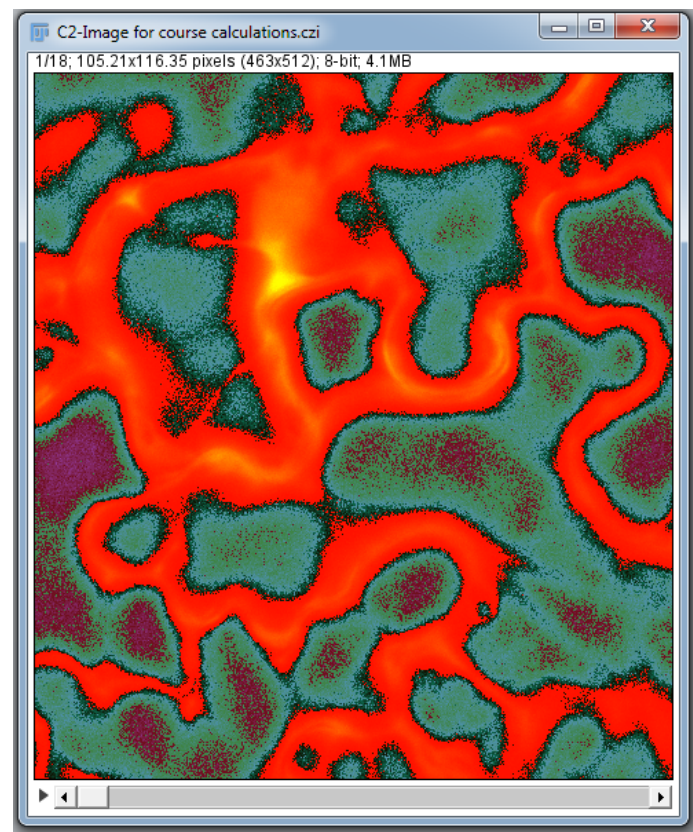
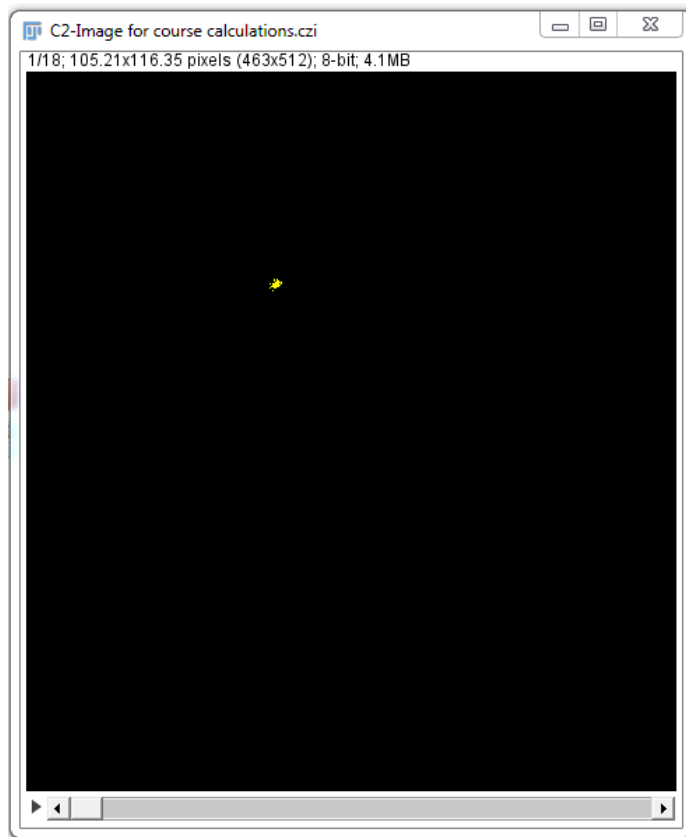


Μπορώ αν επισημάνω συγκεκριμένες εντάσεις



Είναι τα χρώματά μου δυναμικά;





Τύποι αχρωματοψίας



Normal



Protanopia (no red)



Deutanopia (no green)



Tritanopia (no blue)



Protanomaly (low red)



Deuteranomaly (low green)



Tritanomaly (low blue)

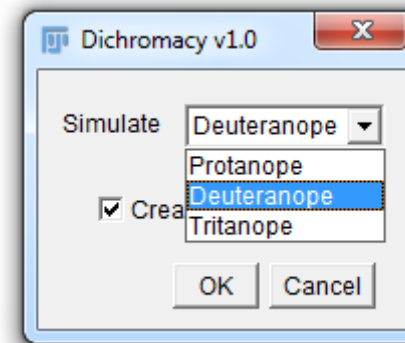
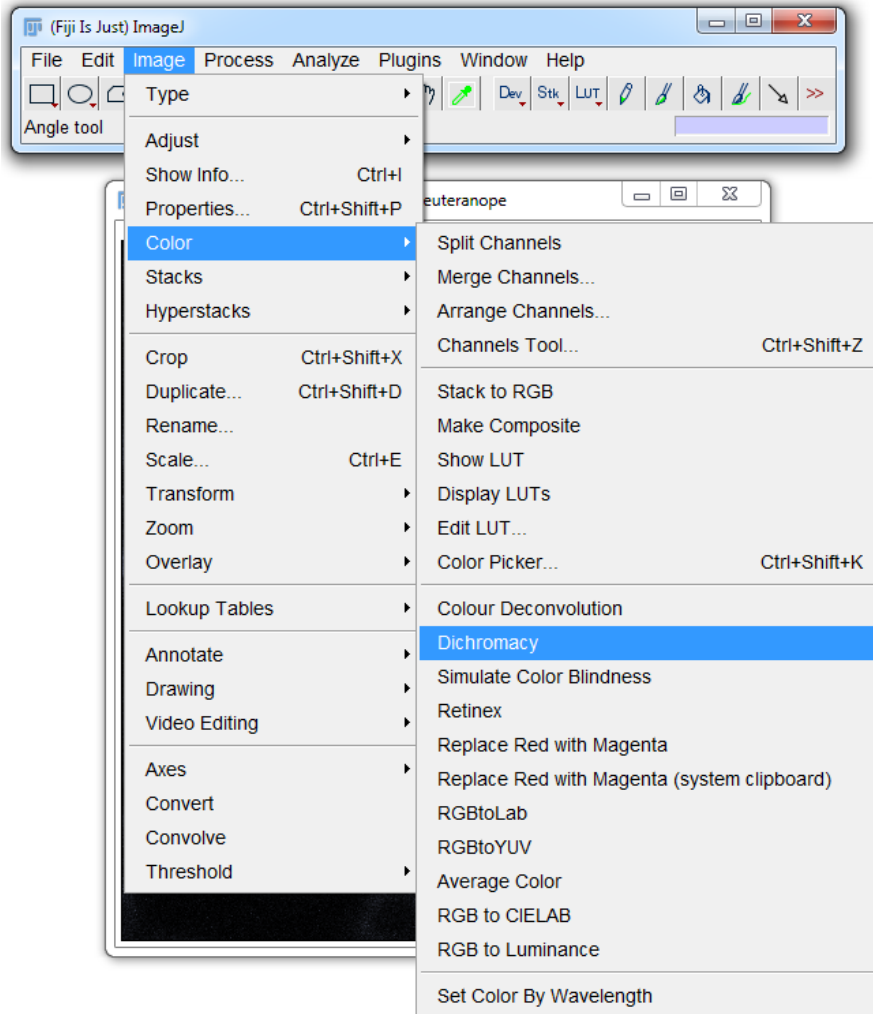


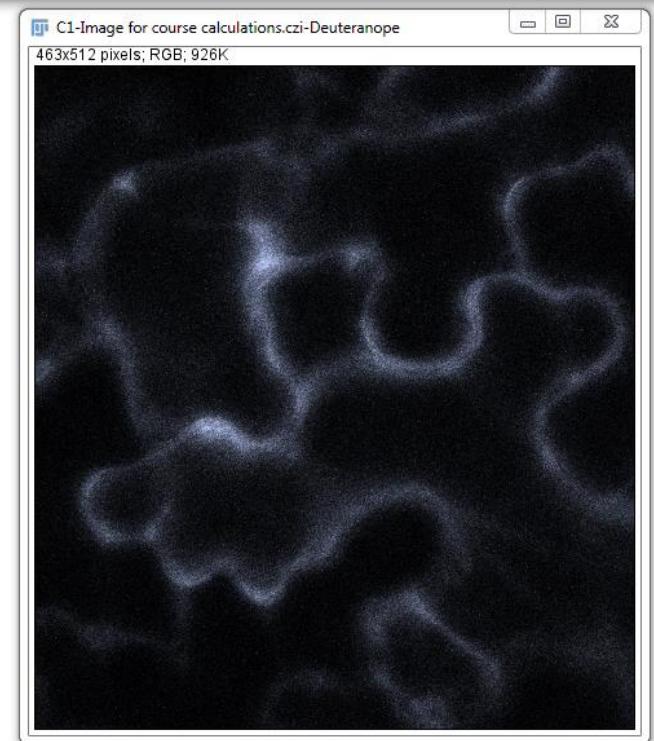
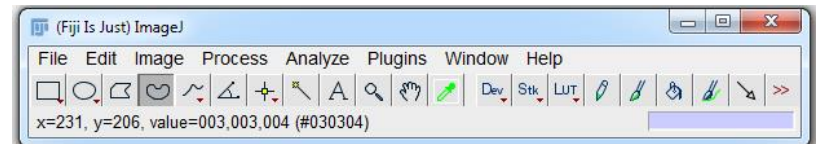
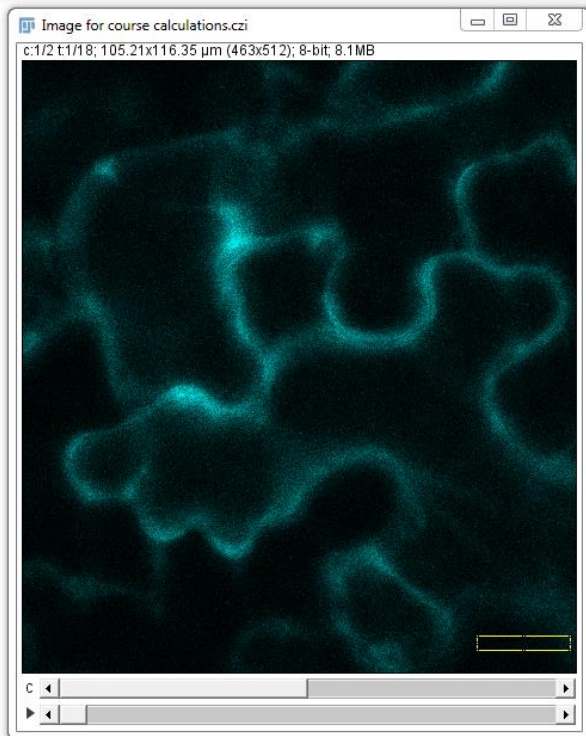
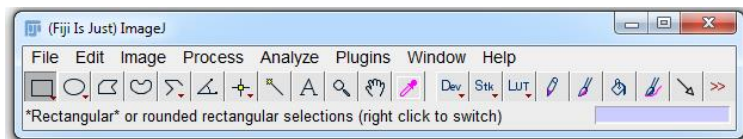
Typical Monochromacy



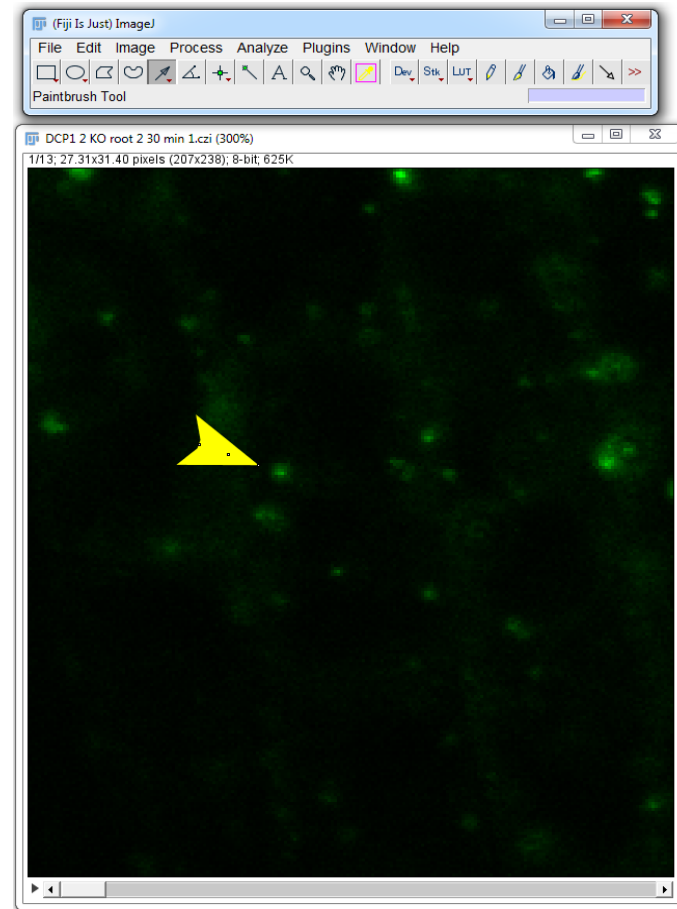
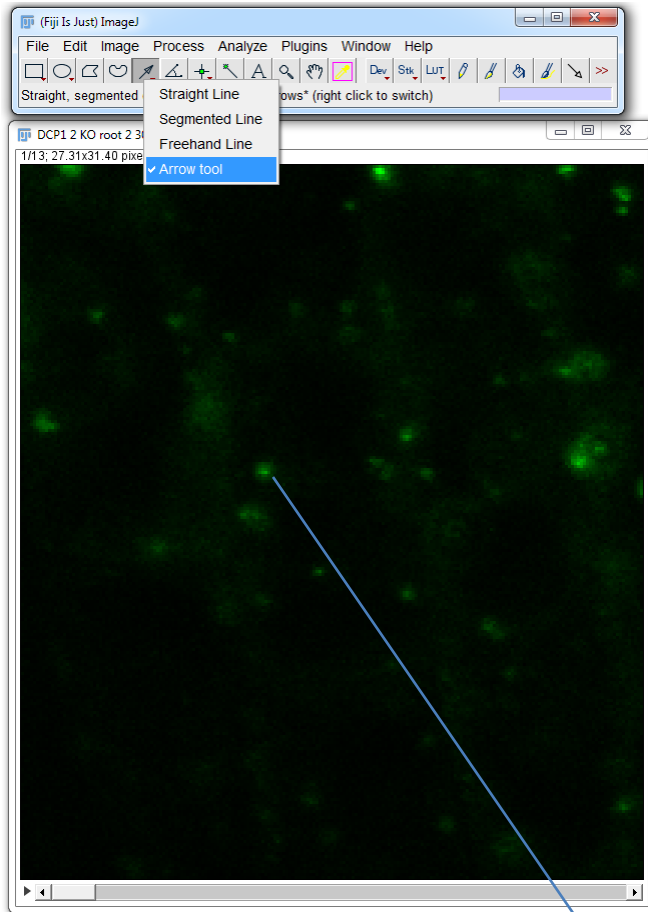
Atypical Monochromacy

Προσομοίωση αχρωματοψίας



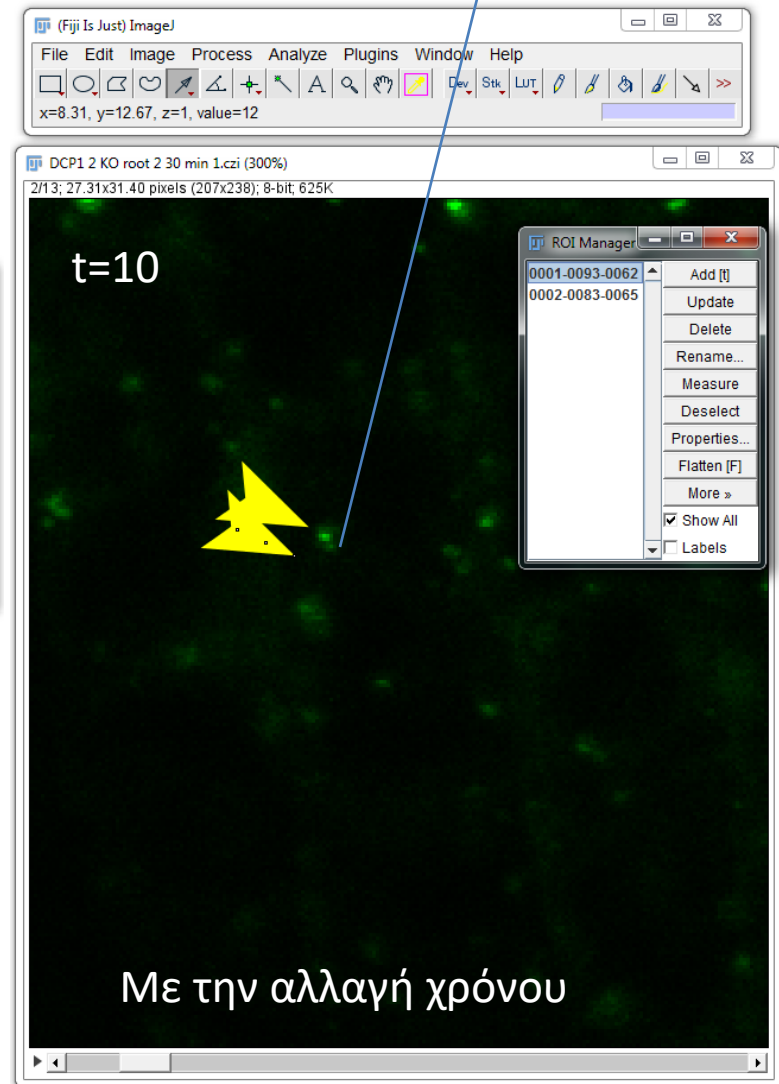
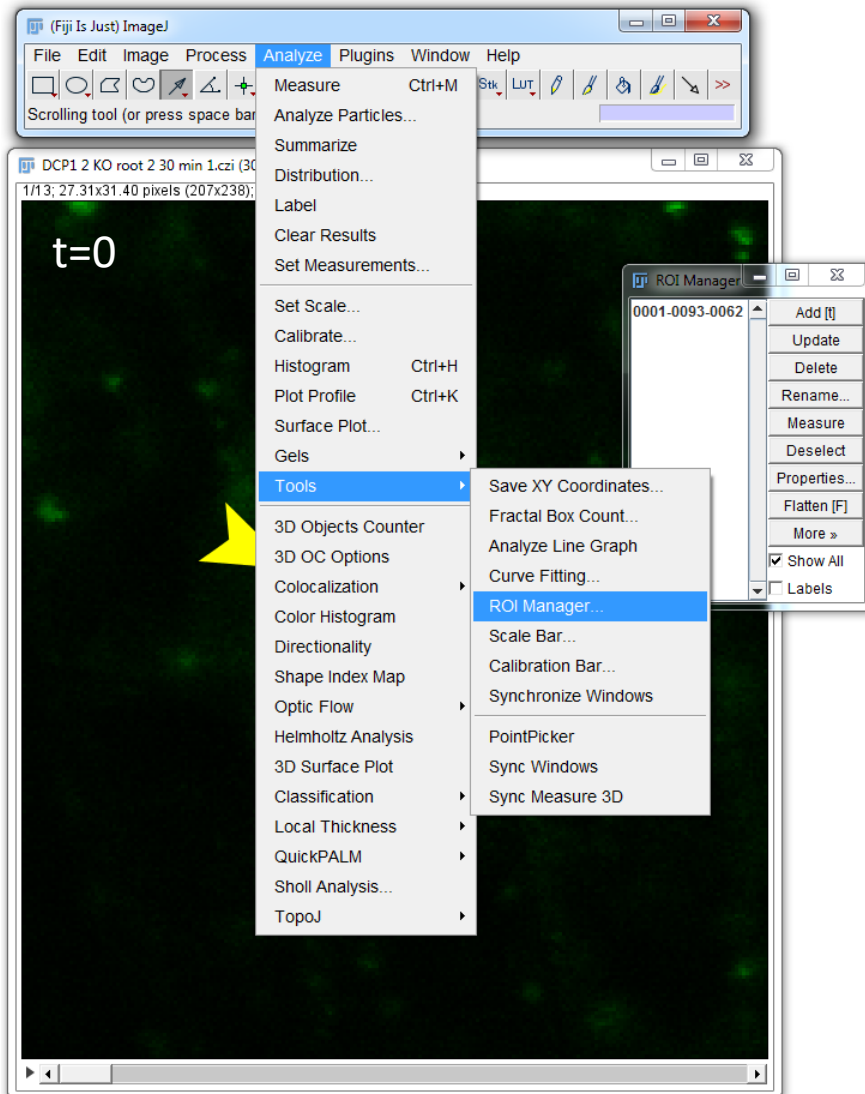


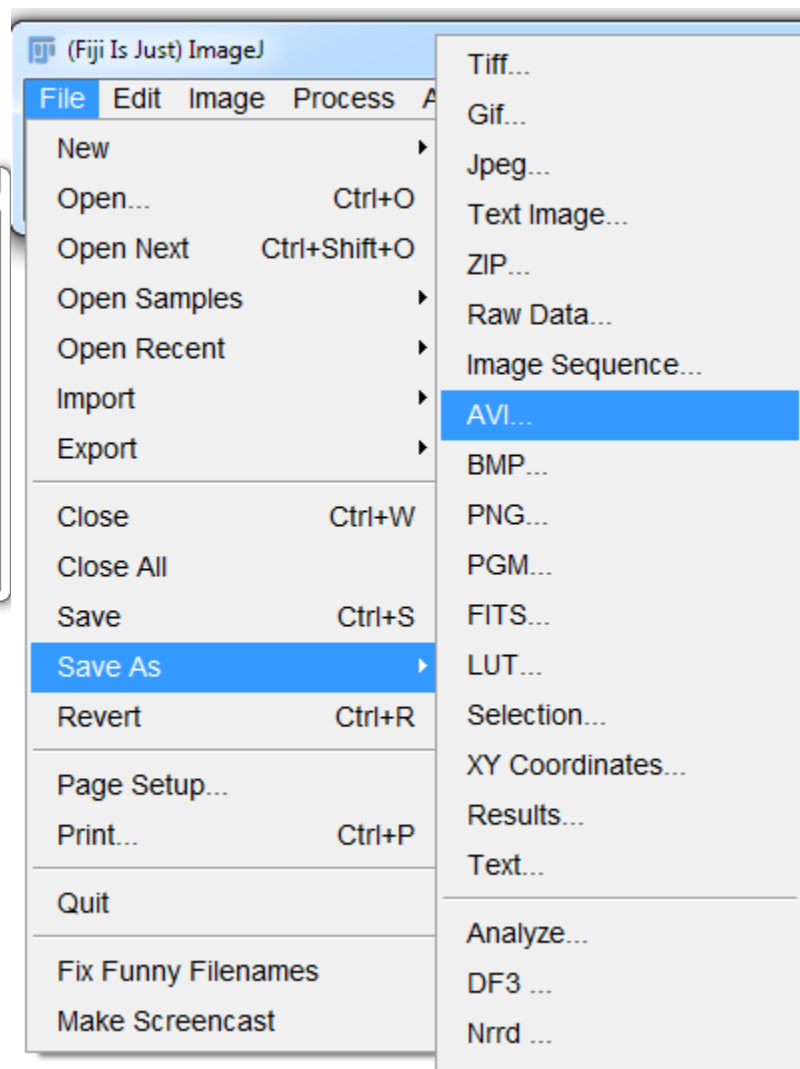
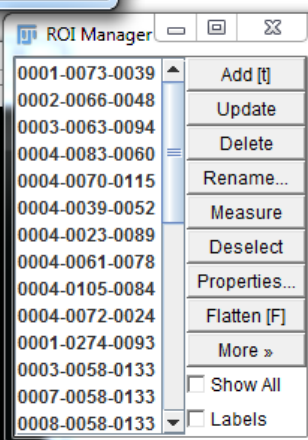
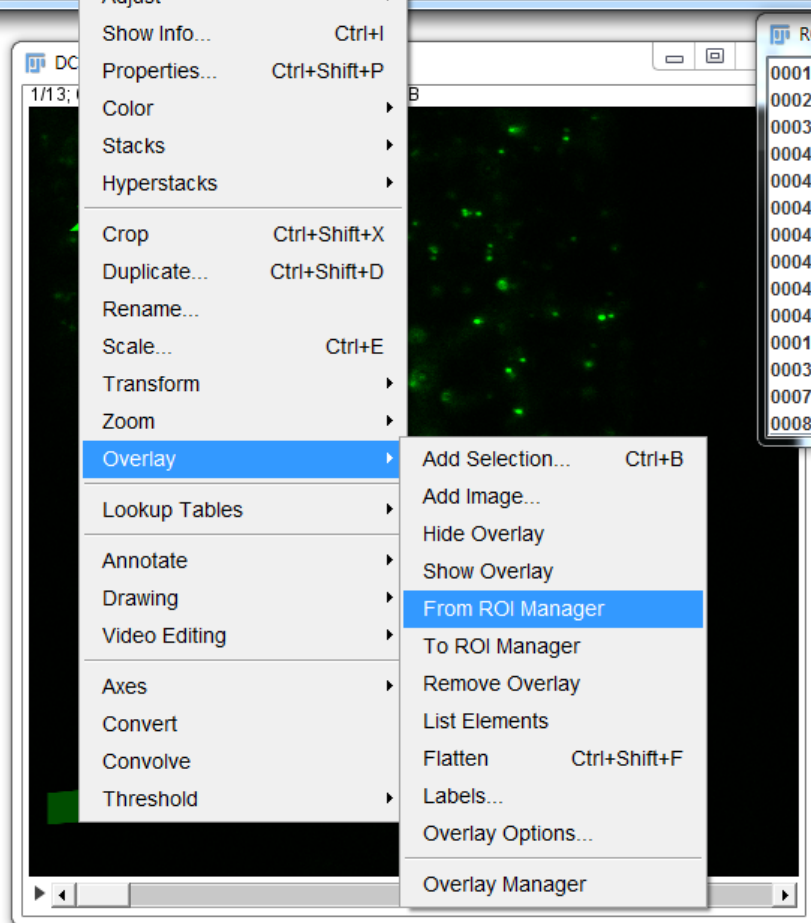
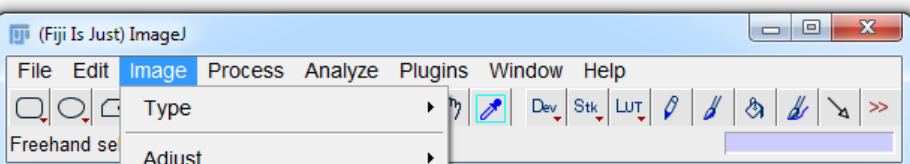
Πώς μπορώ να σχεδιάσω δομές στις εικόνες μου;

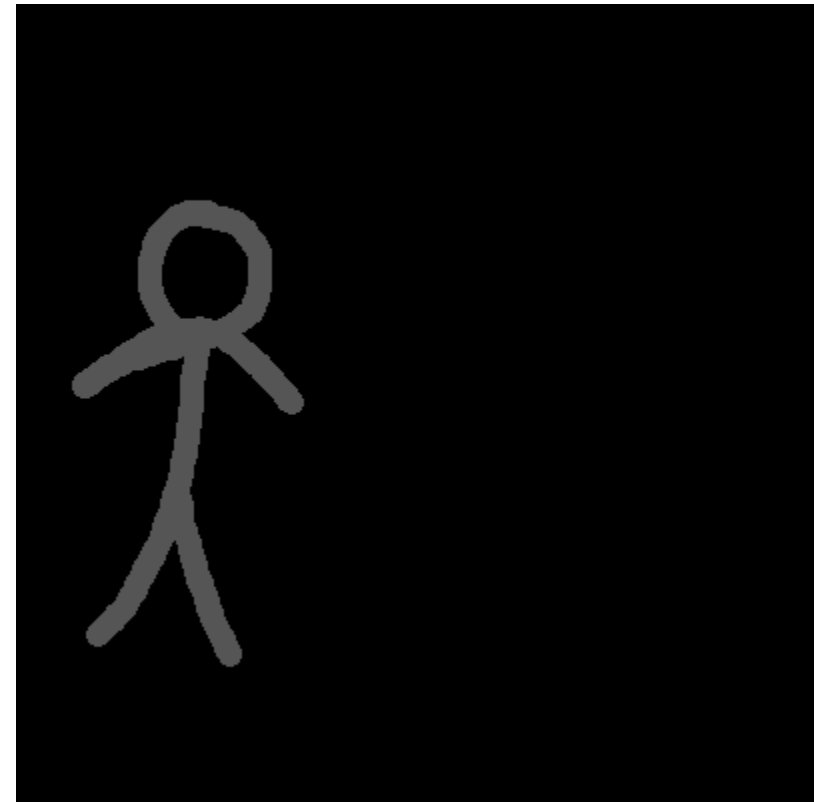
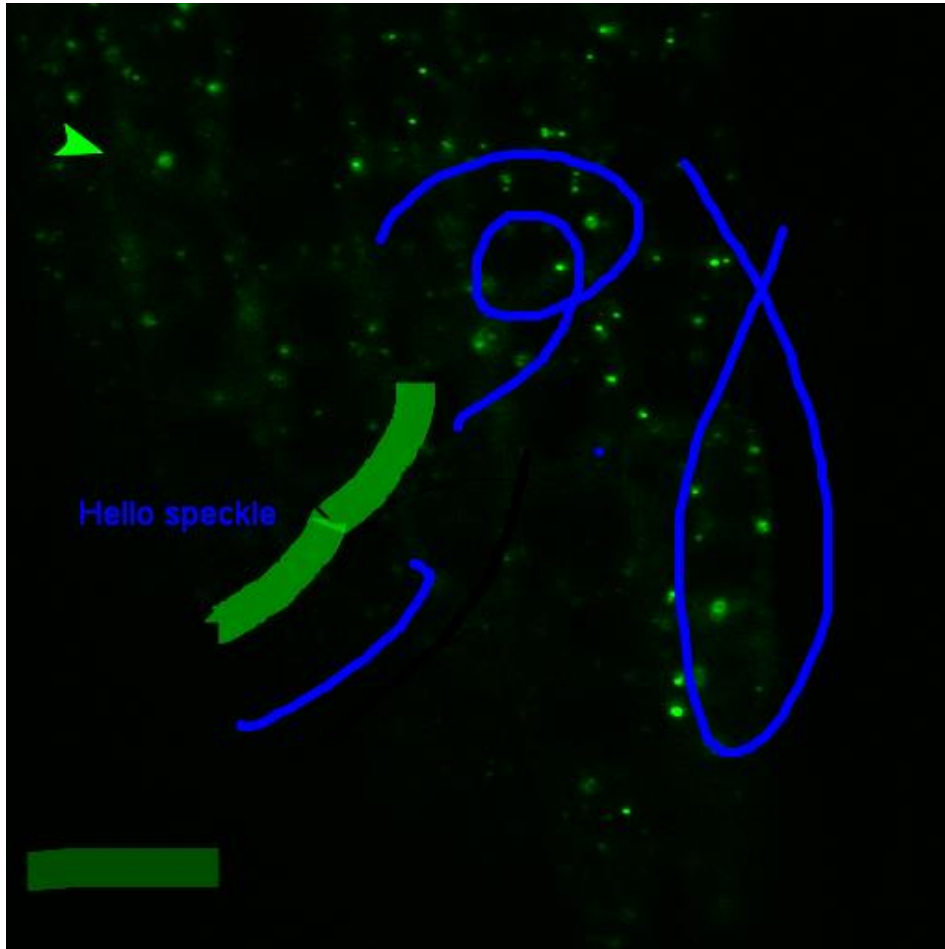


Με ενδιαφέρει αυτό το σημείο

Το σημείο
κινείται...







ΣΥΝΕΝΤΟΠΙΖΟΝΤΑΙ ΟΙ ΔΟΜΕΣ ΜΟΥ;

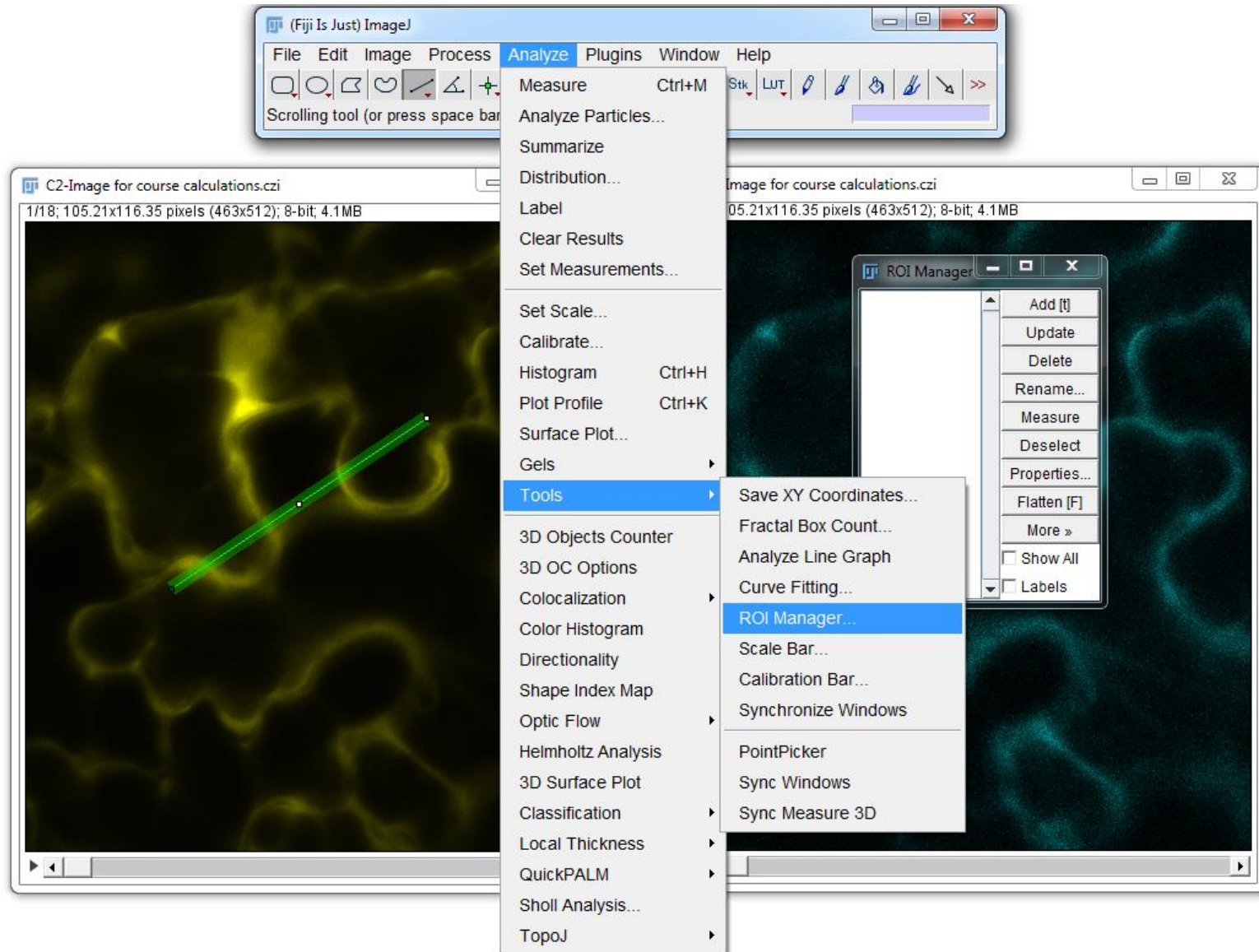
Συσχέτιση μεταξύ χρωμάτων σε δύο κανάλια:

Pearson's correlation coefficient. Απαραίτητη η γραμμική σχέση μεταξύ των χρωμάτων.

Mander's coefficient. Αναλογική με την ένταση σήματος.

Spearman's rank coefficient. Κατατάσσει σε μία σειρά

Διάγραμμα χωρικών εντάσεων



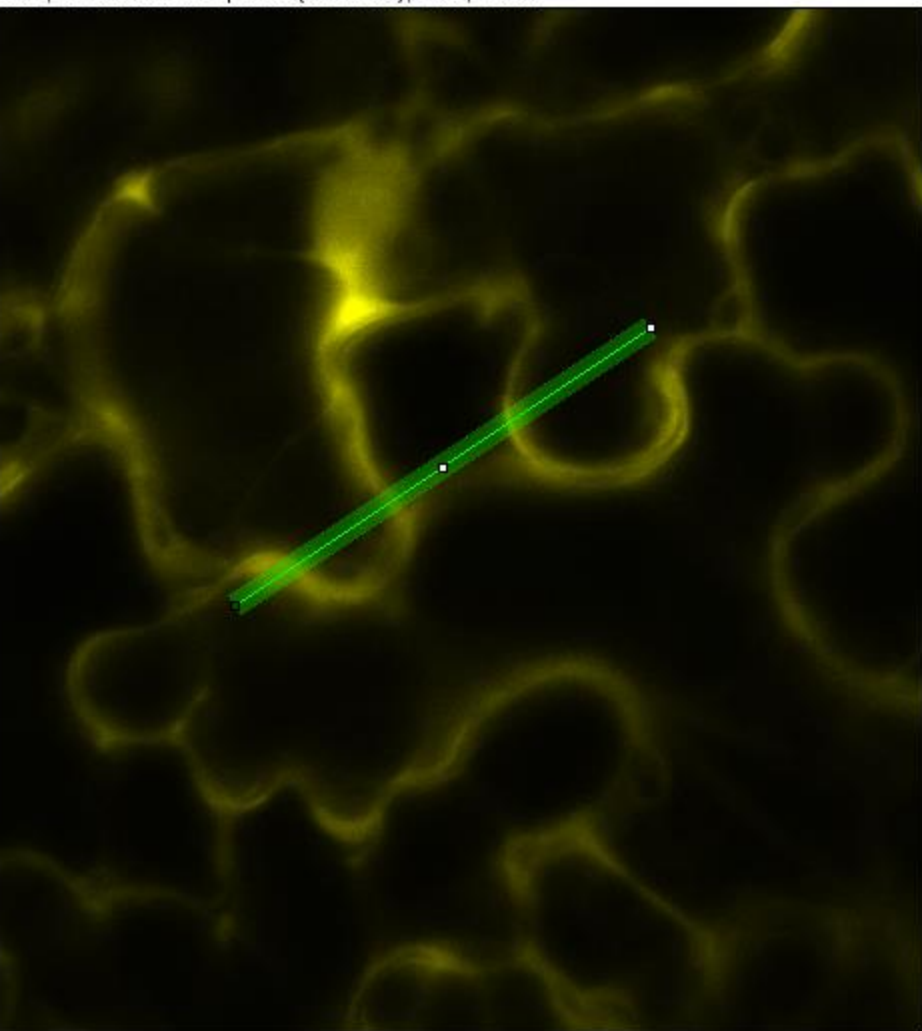
File Edit Image Process Analyze Plugins Window Help



(Fiji Is Just) ImageJ 2.0.0-rc-15/1.49j10; Java 1.6.0_24 [64-bit];

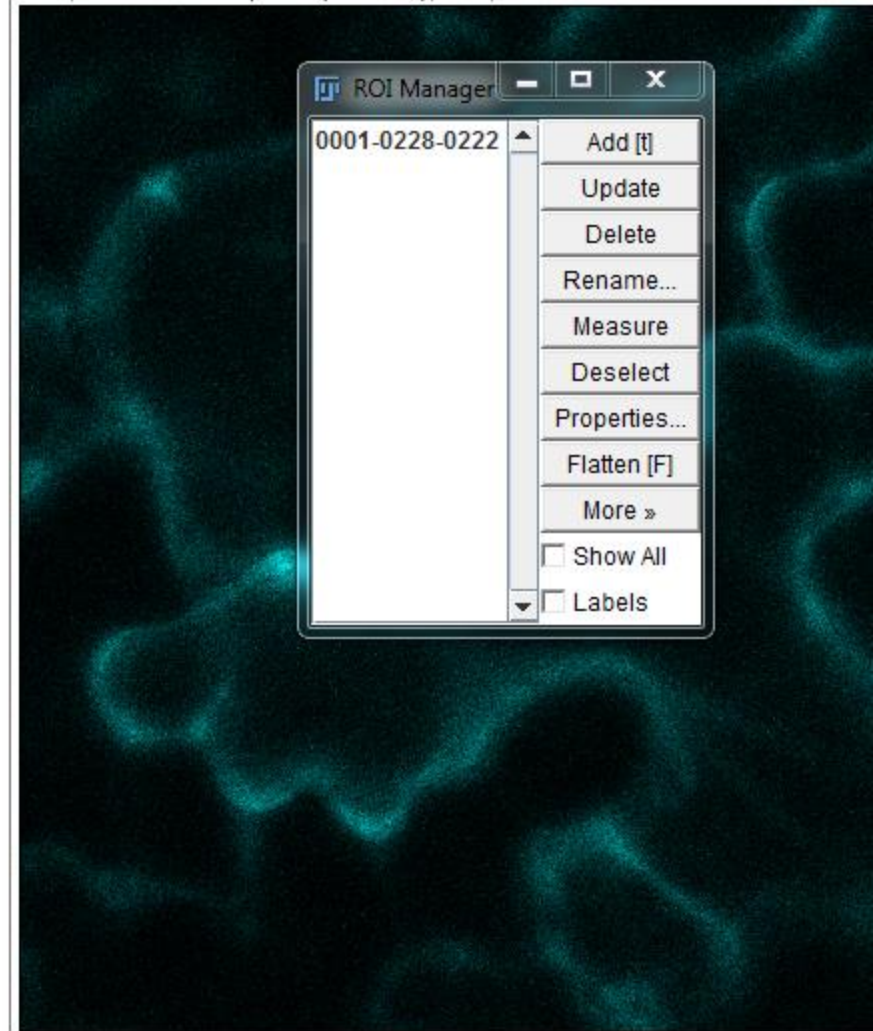
C2-Image for course calculations.czi

1/18; 105.21x116.35 pixels (463x512); 8-bit; 4.1MB

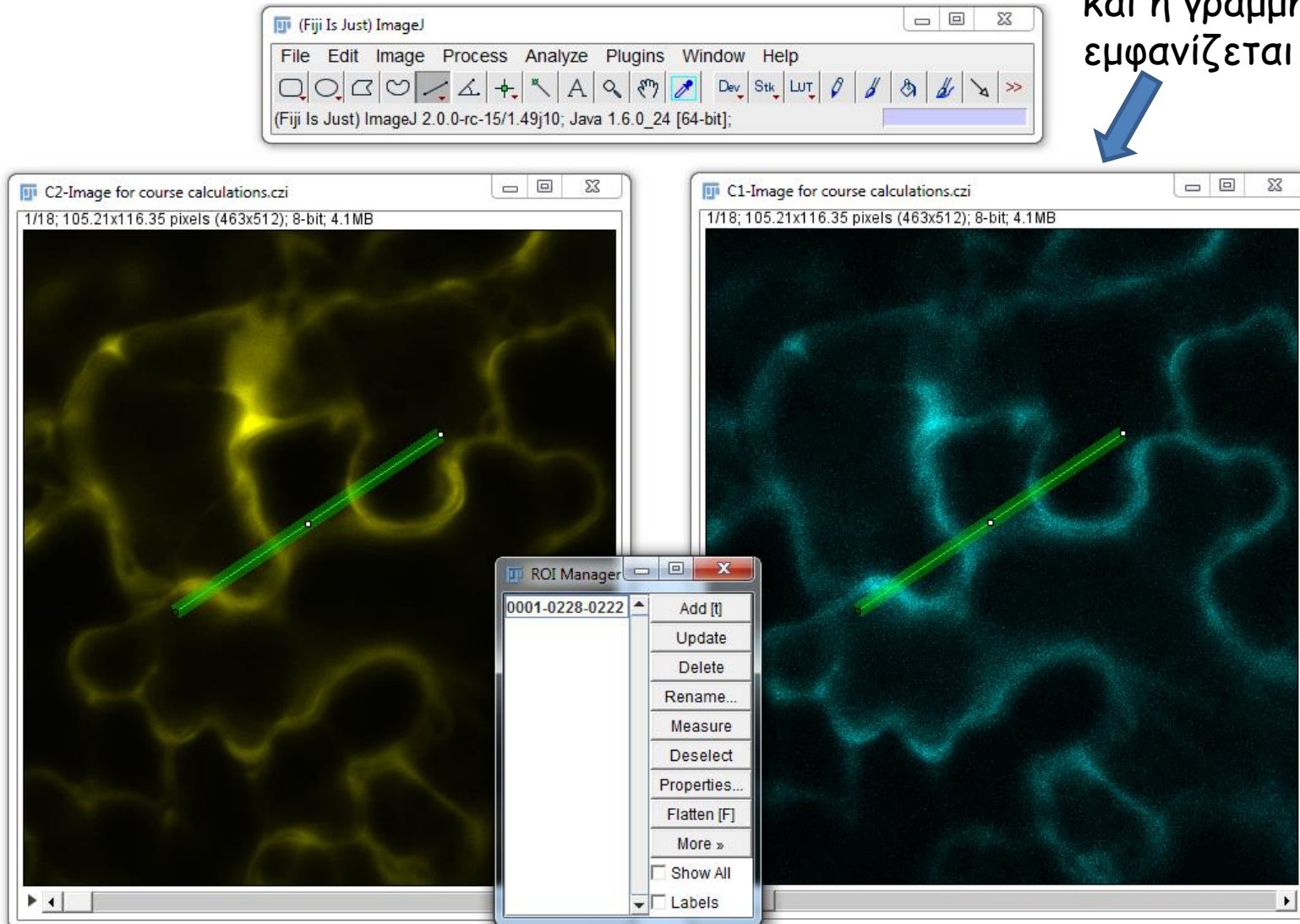


C1-Image for course calculations.czi

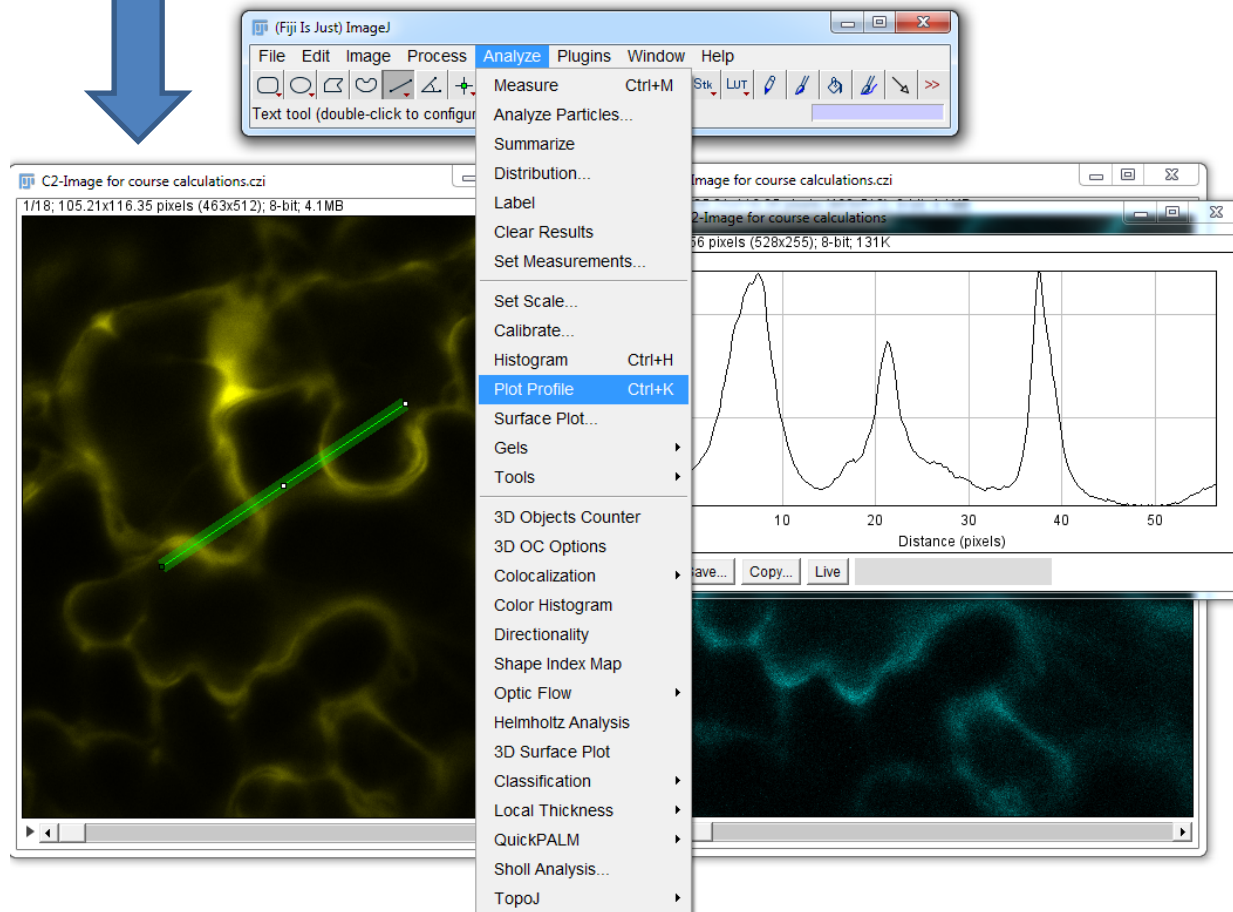
1/18; 105.21x116.35 pixels (463x512); 8-bit; 4.1MB

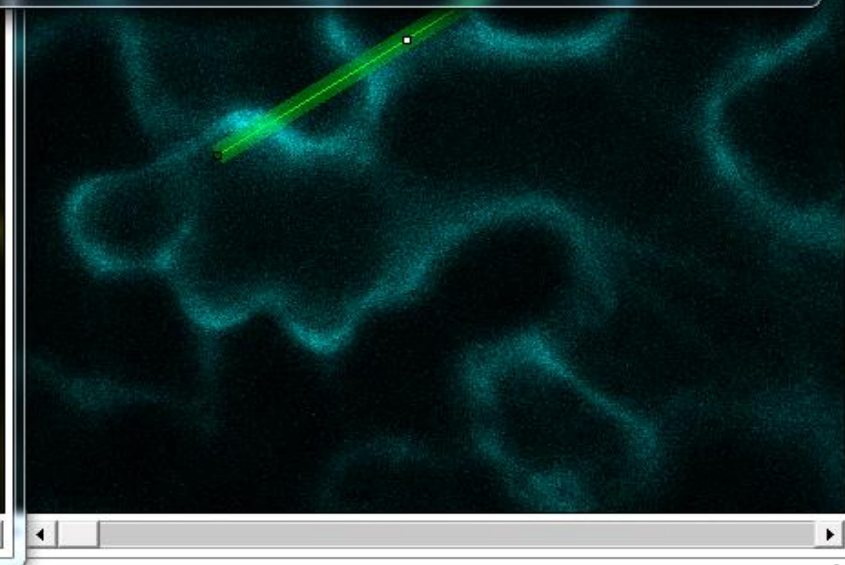
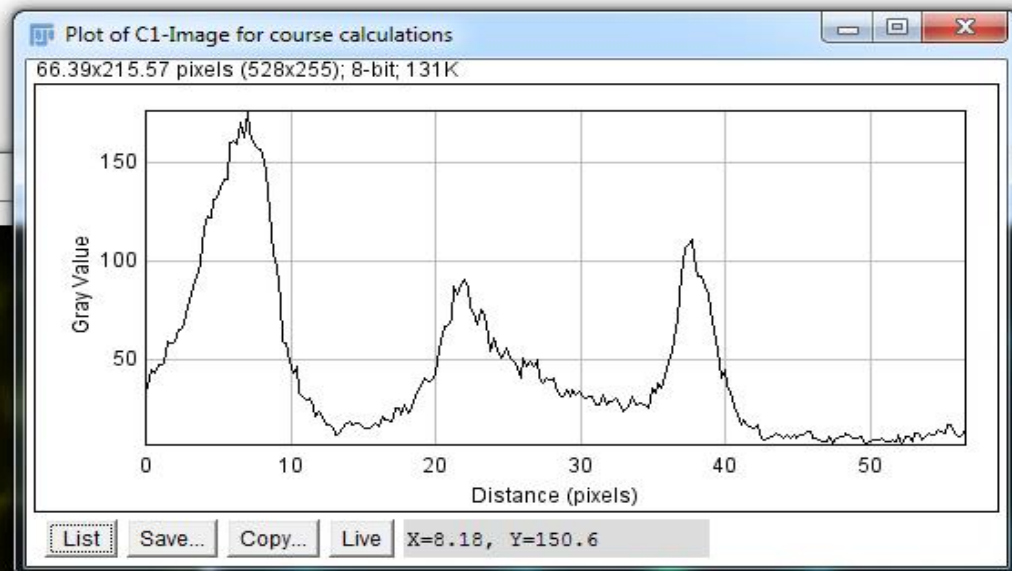
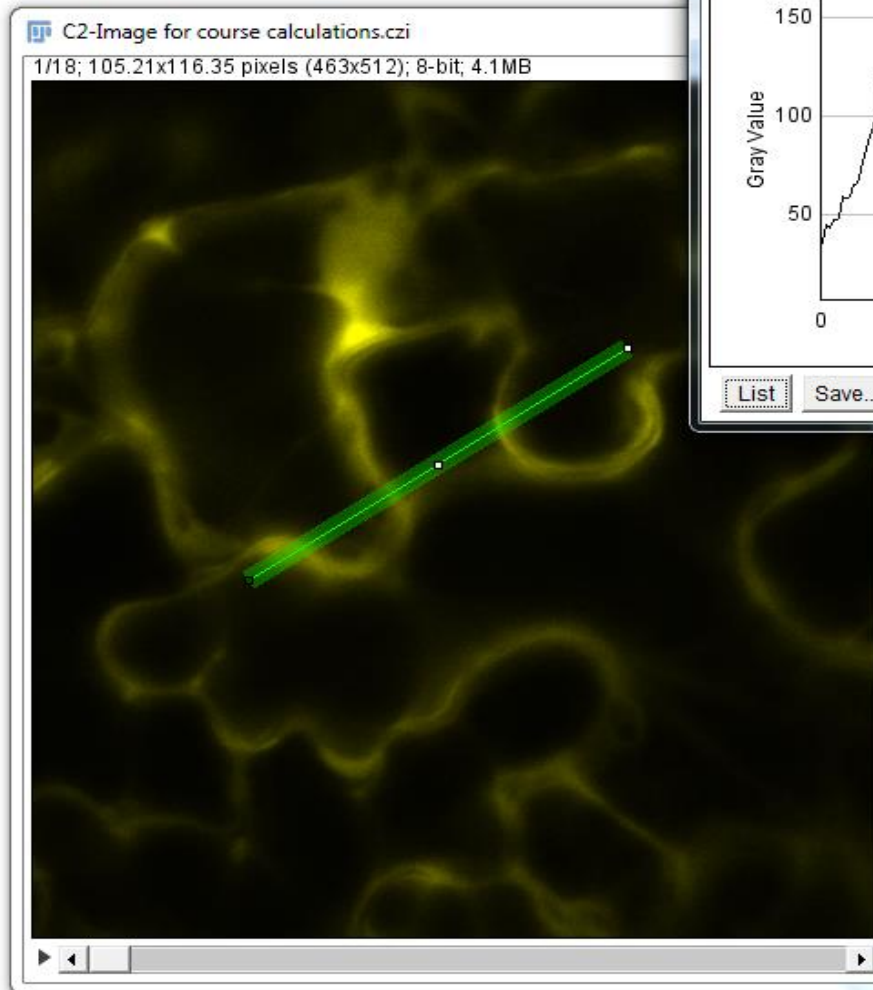
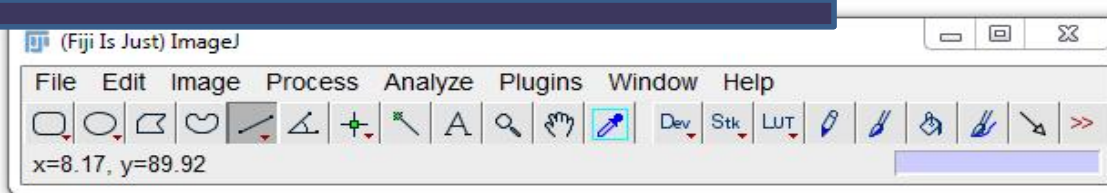


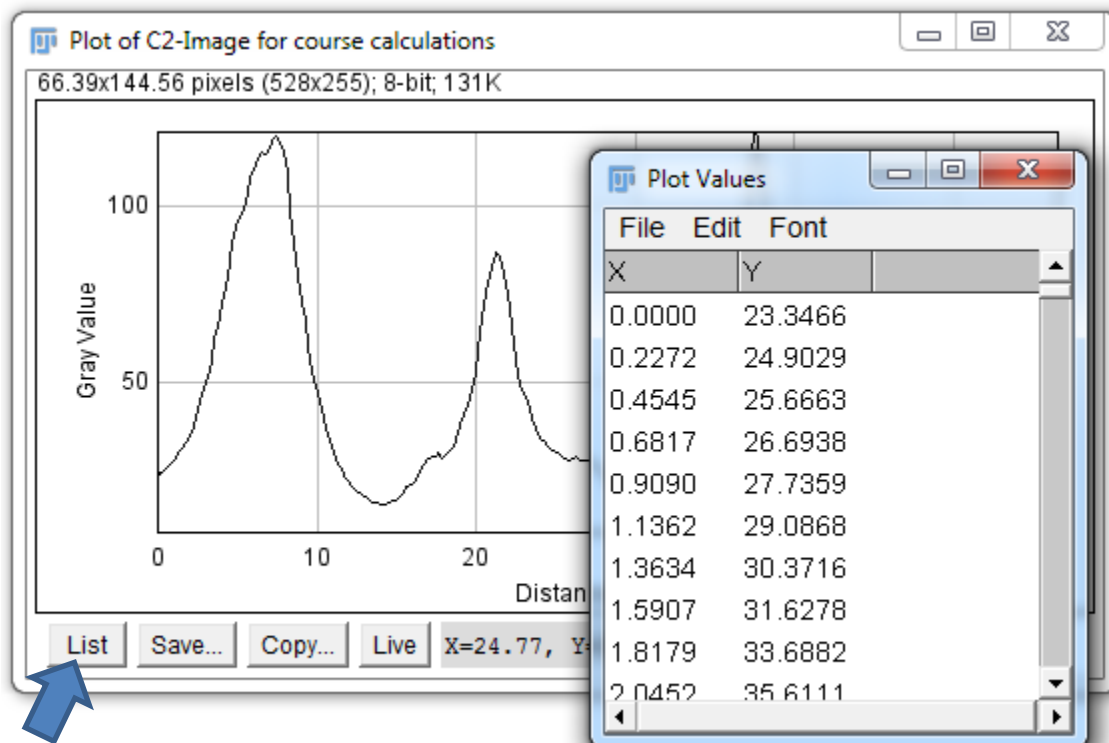
Επιλέγω αυτό το παράθυρο και το ROI από το ROI manager και η γραμμή εμφανίζεται εδώ



Επέλεξε κάθε παράθυρο και
'analyze' -> 'plot profile'

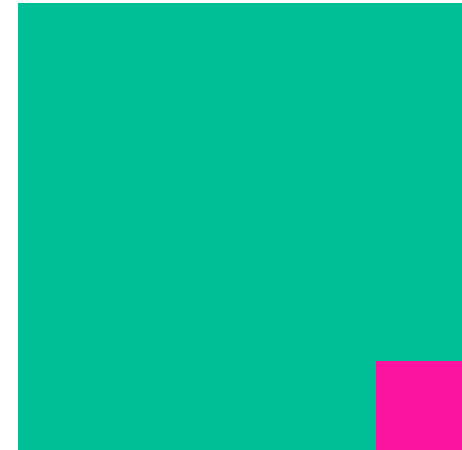
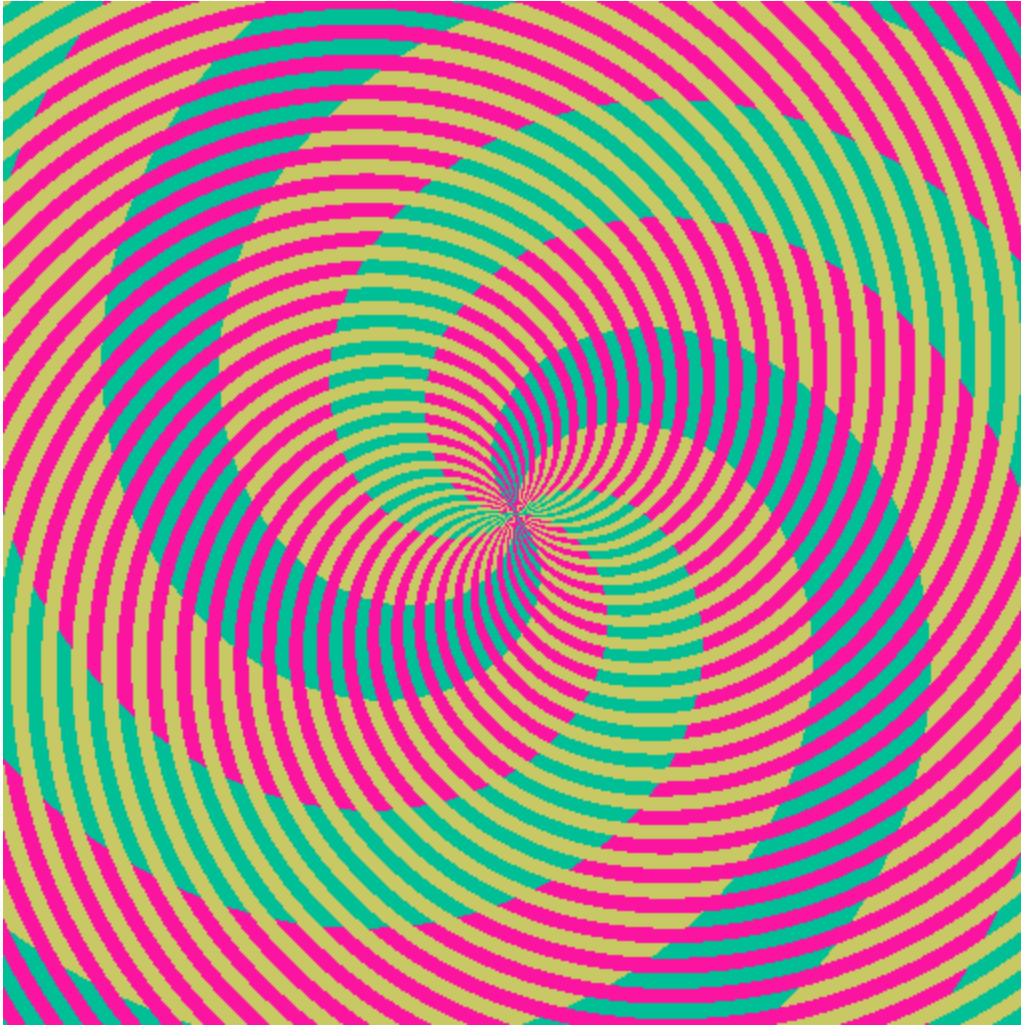


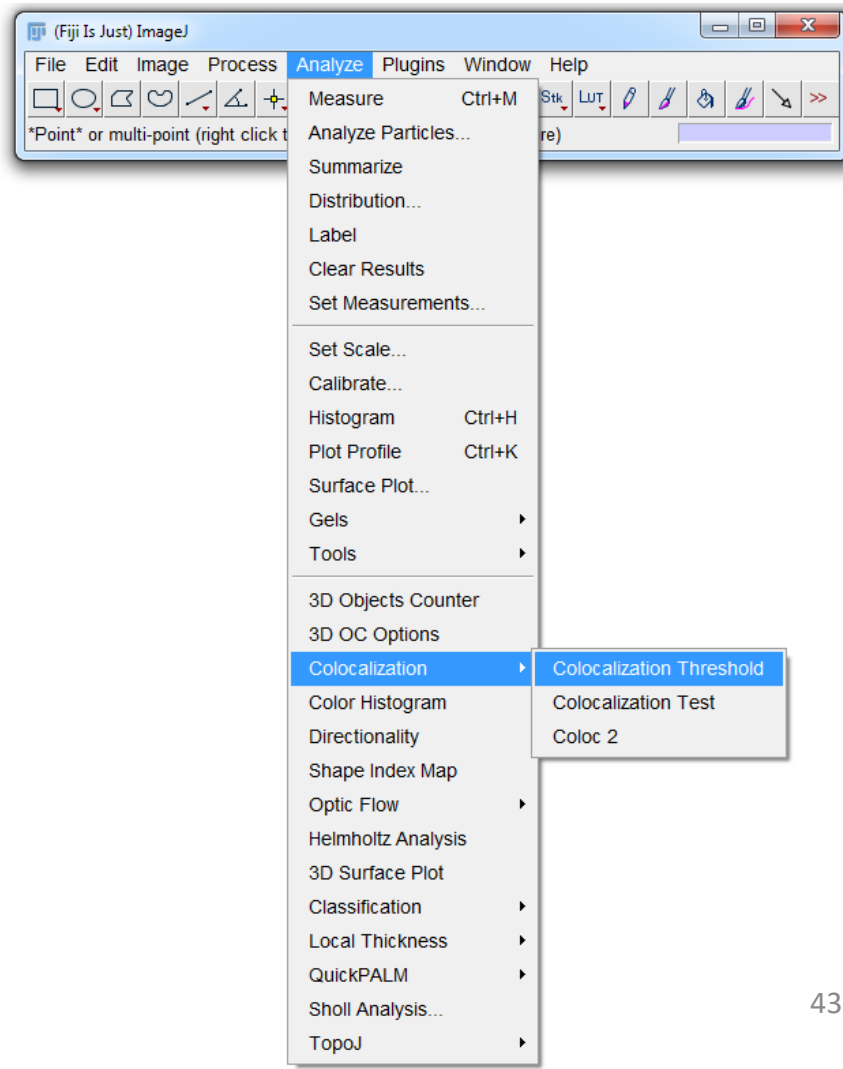
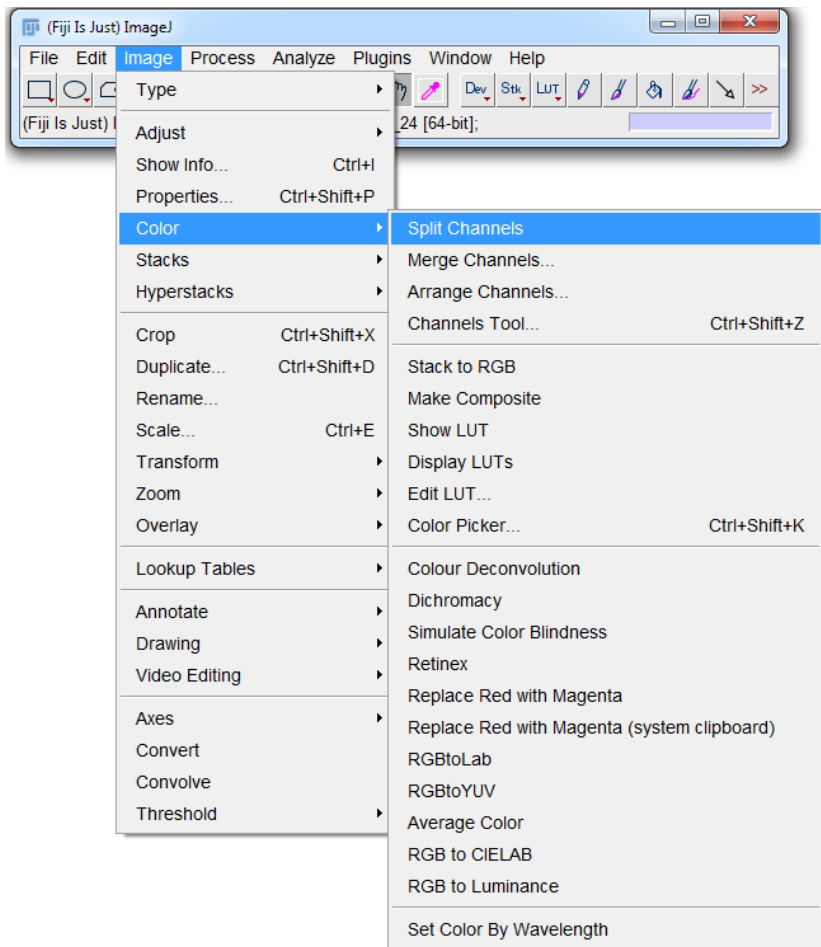


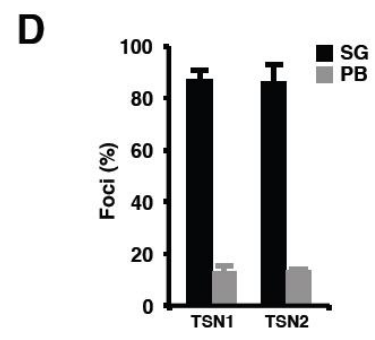
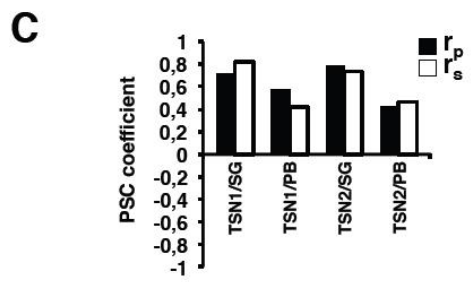
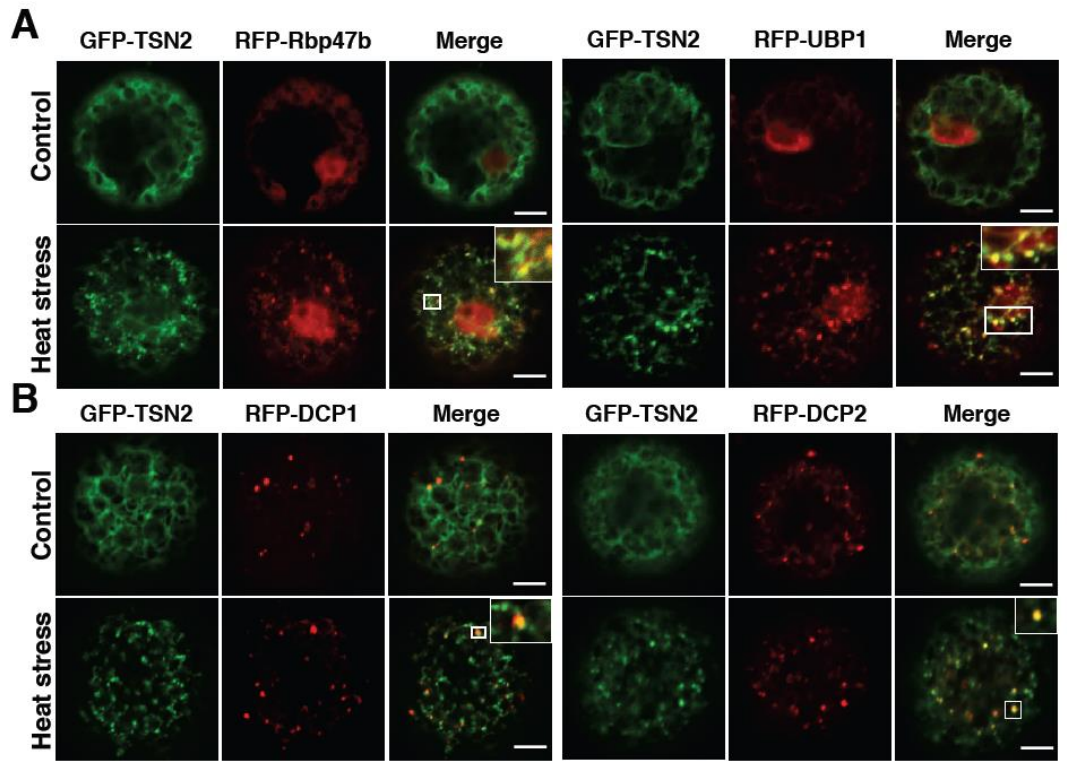
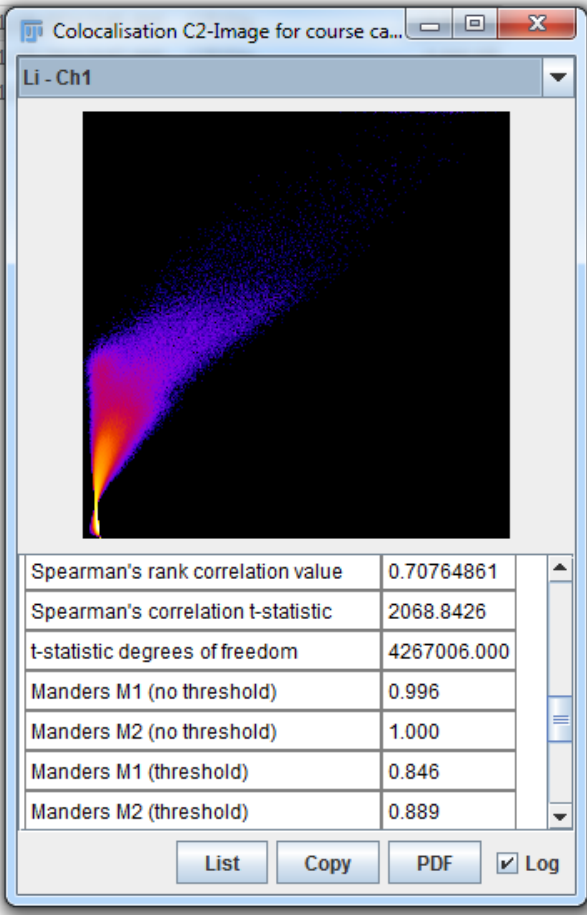


Επέλεξε το list button, πάρε τις τιμές και μετέφερε τις στο Excel

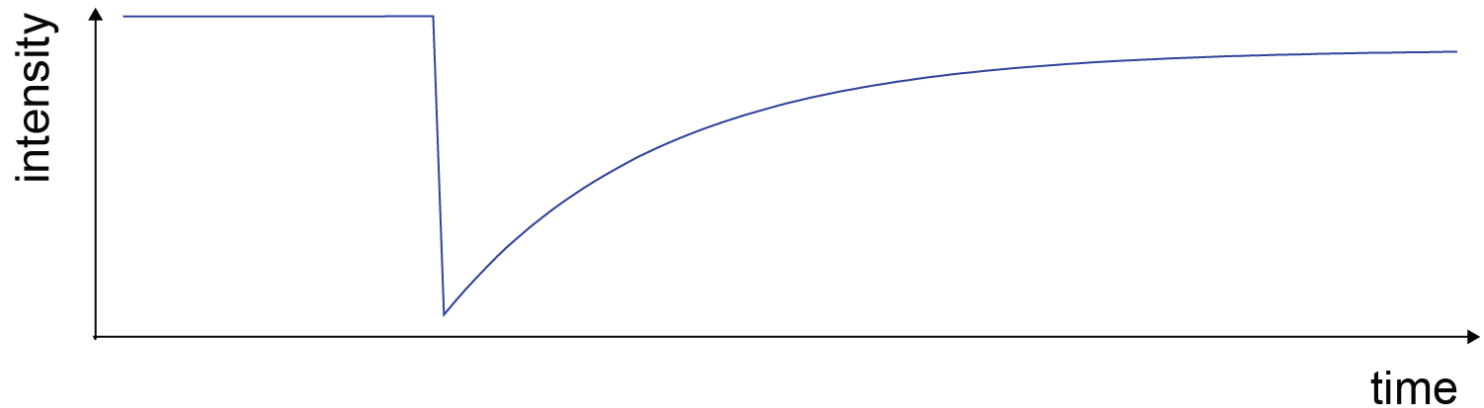
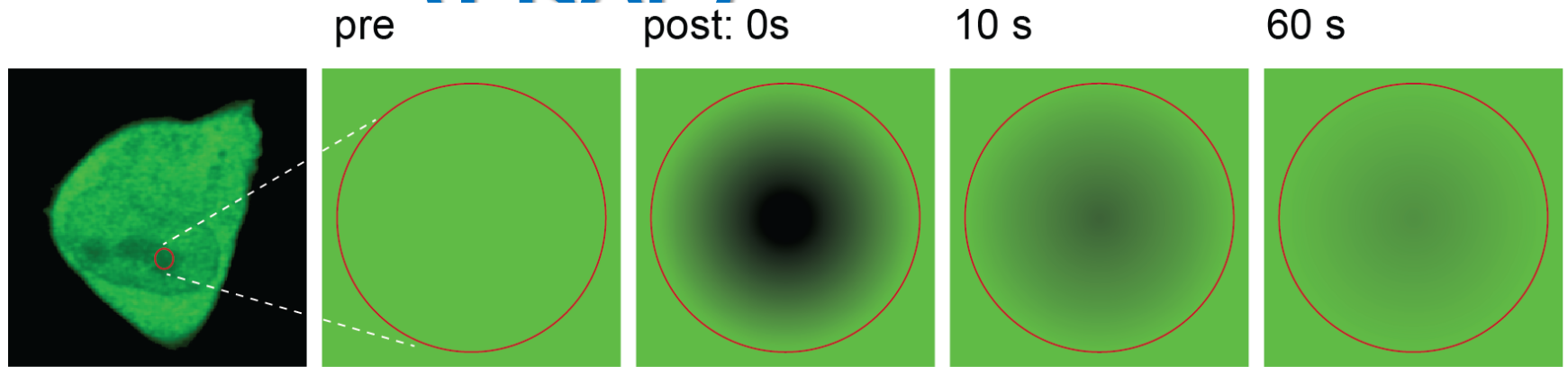
Οκ! Πάμε για μετρήσεις!

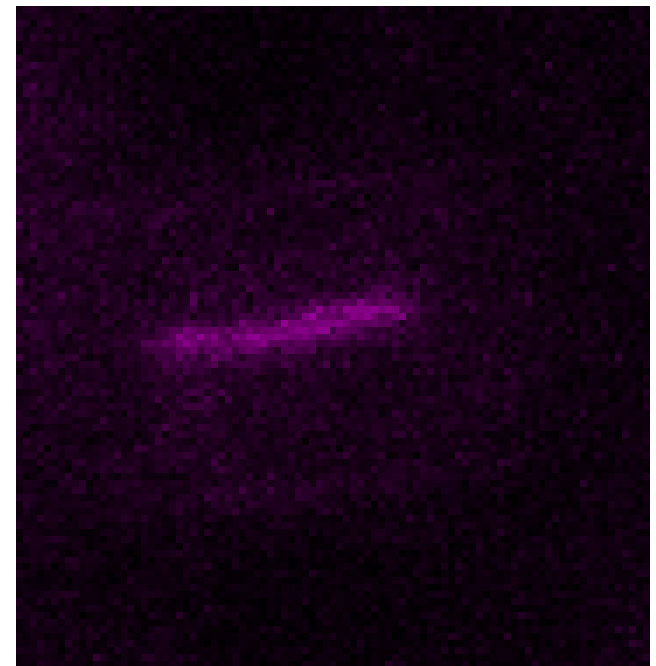
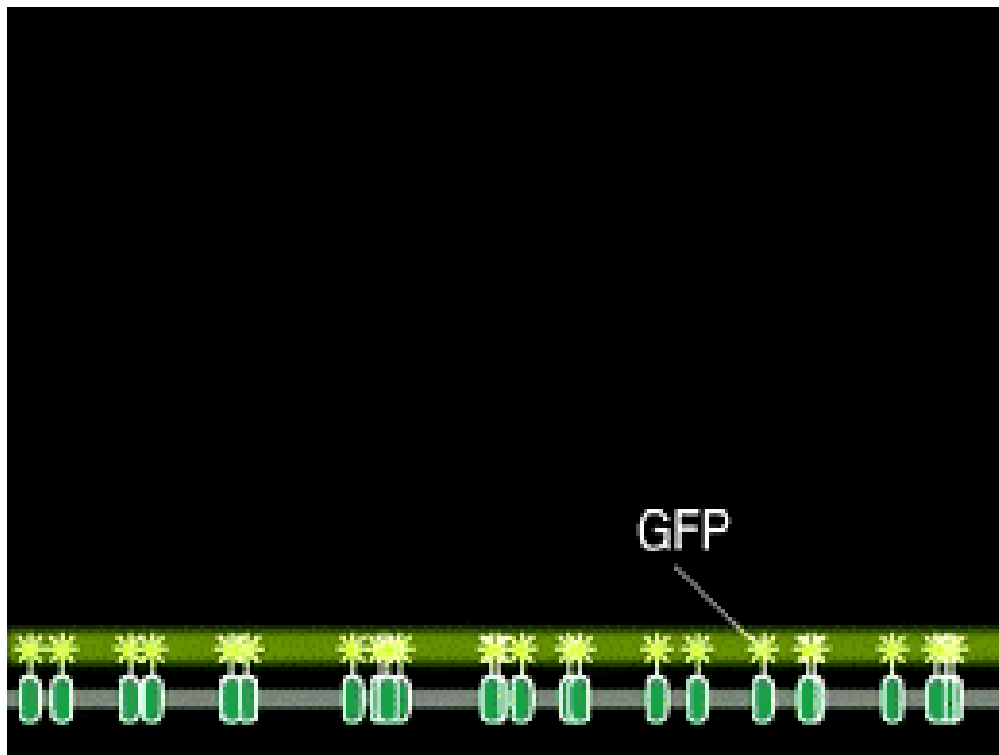






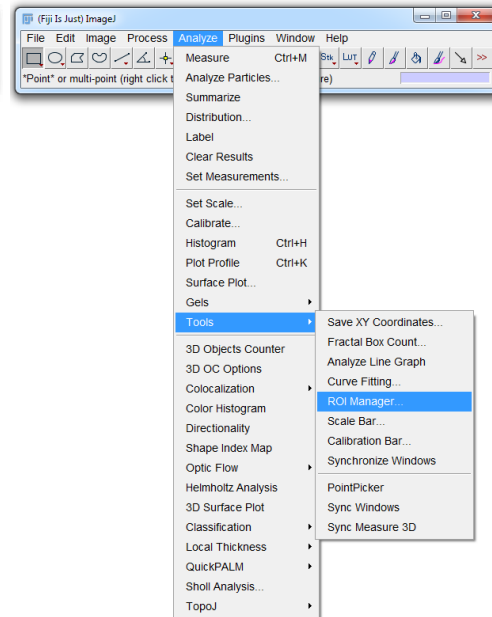
Fluorescence recovery after photo-bleaching (FRAP)



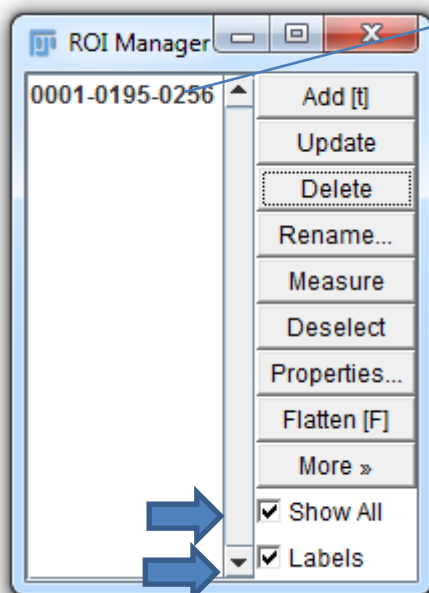


Ανάλυση δεδομένων FRAP

Select the ROI → Use the rectangle tool to select the ROI → Select the ROI manager

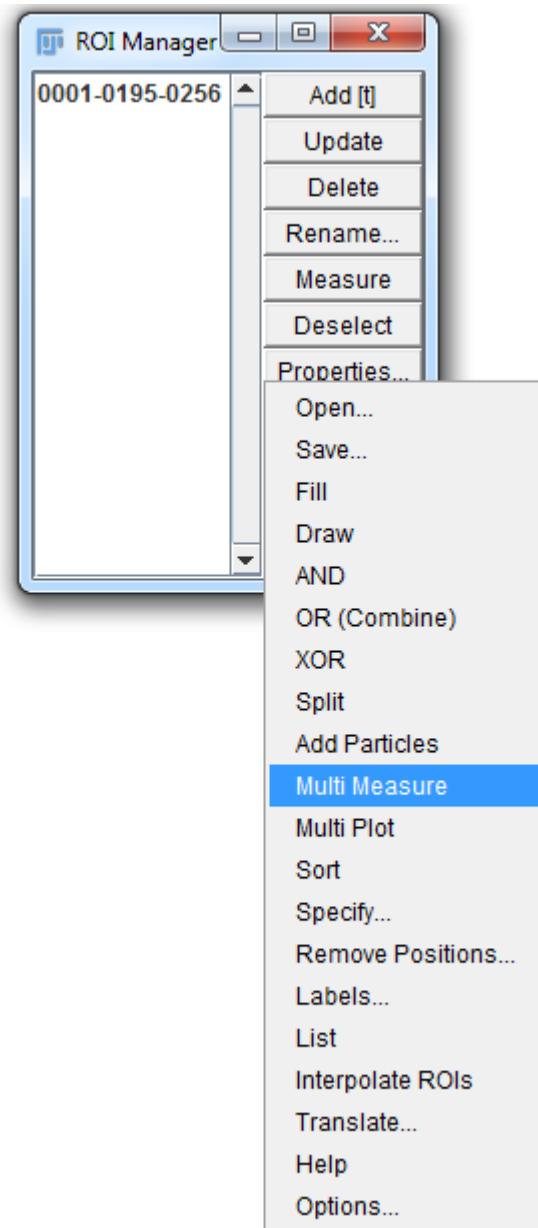


2. 'show all' και 'labels'



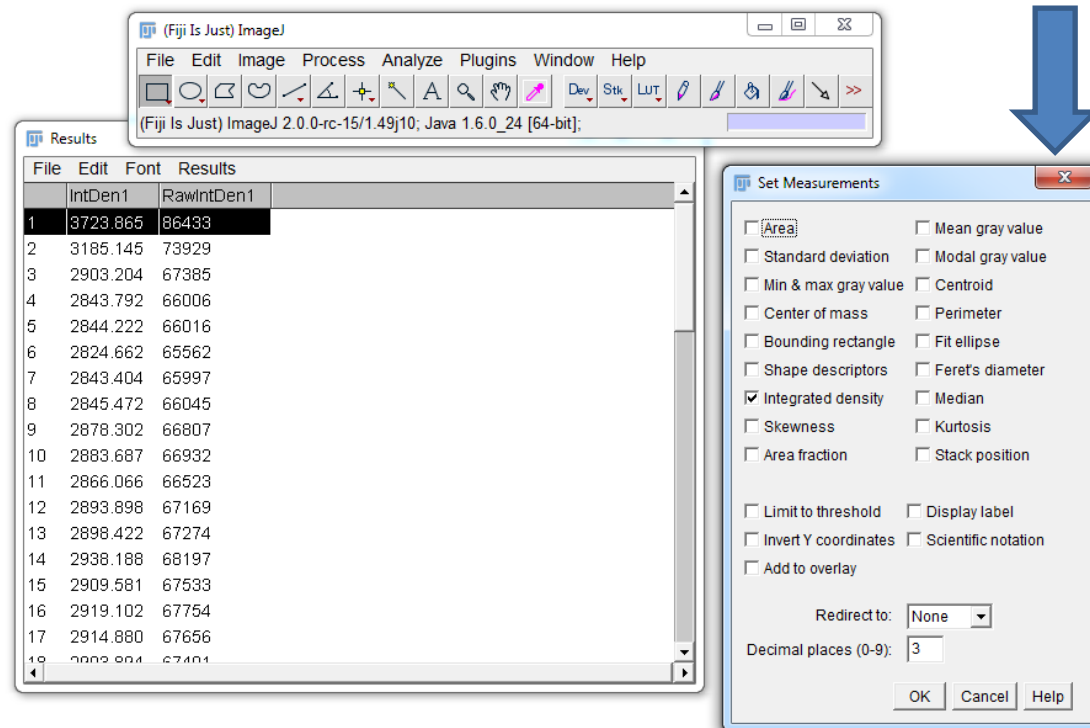
Αυτό
είναι το
ROI

3. επέλεξε 'more' και
'Multi measure'



- 4. καταλήγεις με μετρήσεις εντάσεων

*‘integrated density’ πρέπει να επιλεχθεί
(Results-> Set measurements...)*



- 5. στο Excel

Χρόνος

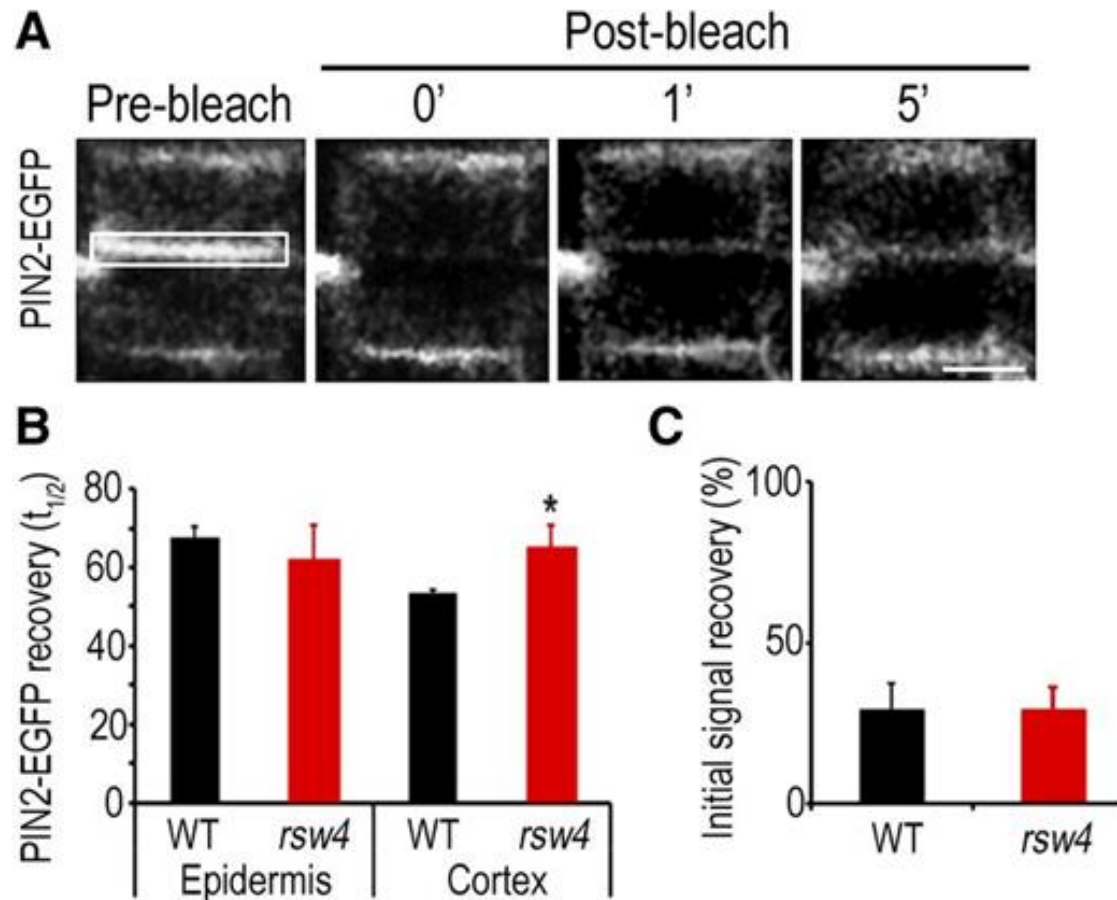


B	C	D	E	F	G	H
WT						
GFP-Ubp						
ROOT1	Value	IntDen	background	Time		
	1	194,068,542,827	46,903,794,809	147,164,748,018		
	2	64,253,054,902	35,820,844,291	28,432,210,611	0	28
	3	75,289,241,916	40,871,302,754	34,417,939,162	3	34
	4	81,892,248,723	42,638,963,217	39,253,285,506	6	39
	5	82,836,871,509	43,901,577,833	38,935,293,676	9	38
	6	85,820,383,084	44,696,557,406	41,123,825,678	12	41
	7	93,050,020,848	47,801,654,091	45,248,366,757	15	45
	8	91,413,298,197	48,690,160,673	42,723,137,524	18	42
	9	88,990,948,675	50,888,045,375	38,102,903,300	21	38
	10	91,114,011,770	50,579,406,246	40,534,605,524	24	40
	11	91,937,049,445	49,662,841,562	42,274,207,883	27	40
	12	89,813,986,350	50,785,165,665	39,028,820,685	30	42
	13	85,988,731,699	53,609,681,325	32,379,050,374	33	45
	14	85,876,499,289	54,470,129,804	31,406,369,485	36	50
	15	87,662,865,153	52,524,768,025	35,138,097,128	39	52

Κανονικοποίηση με
το θόρυβο (my
intensity-
background)

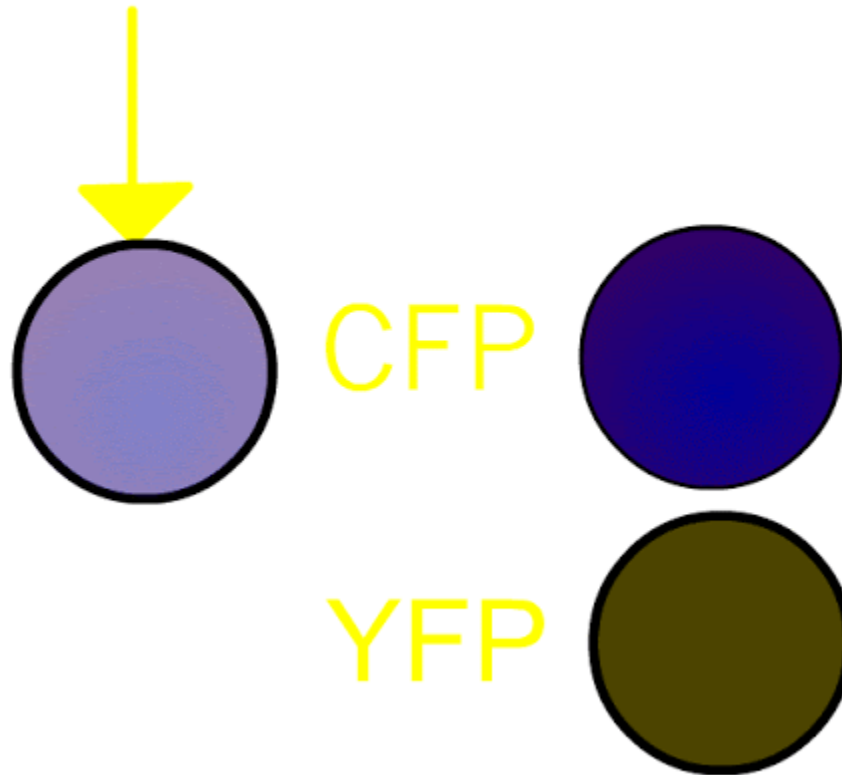


Παρουσίαση FRAP

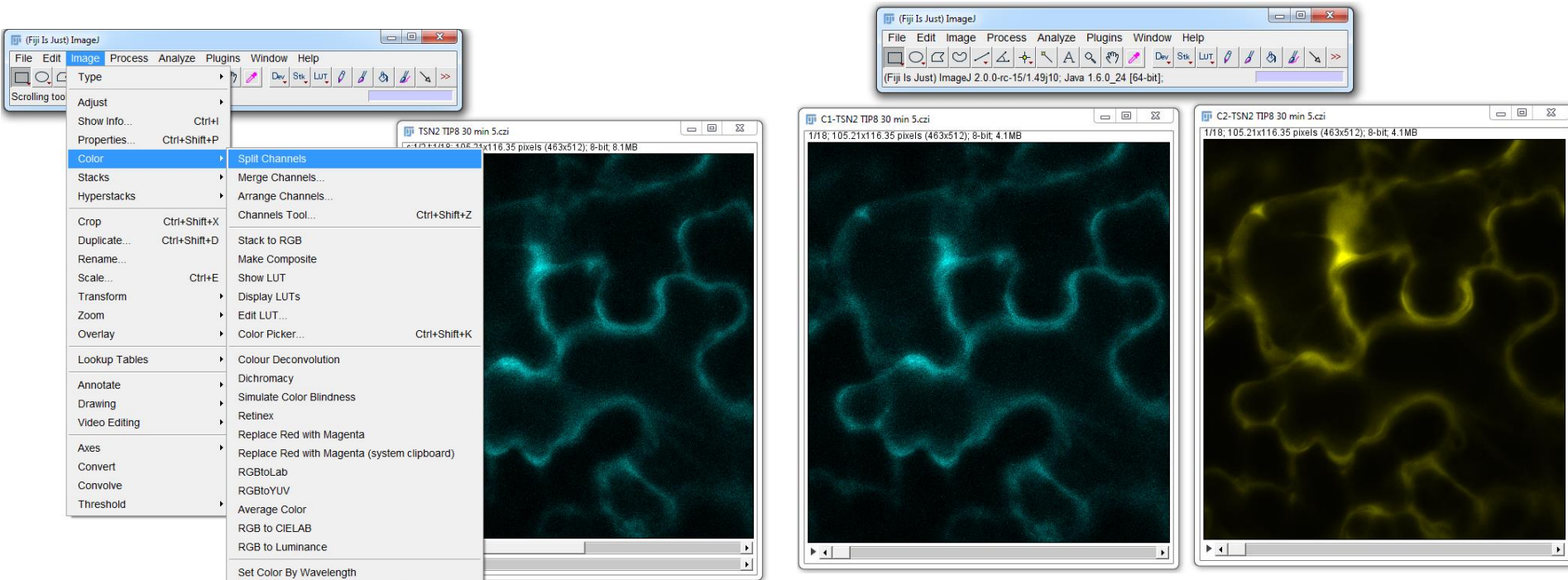


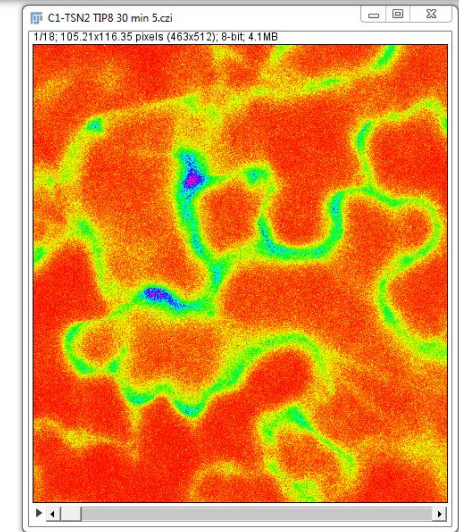
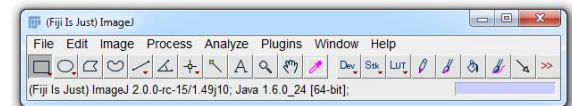
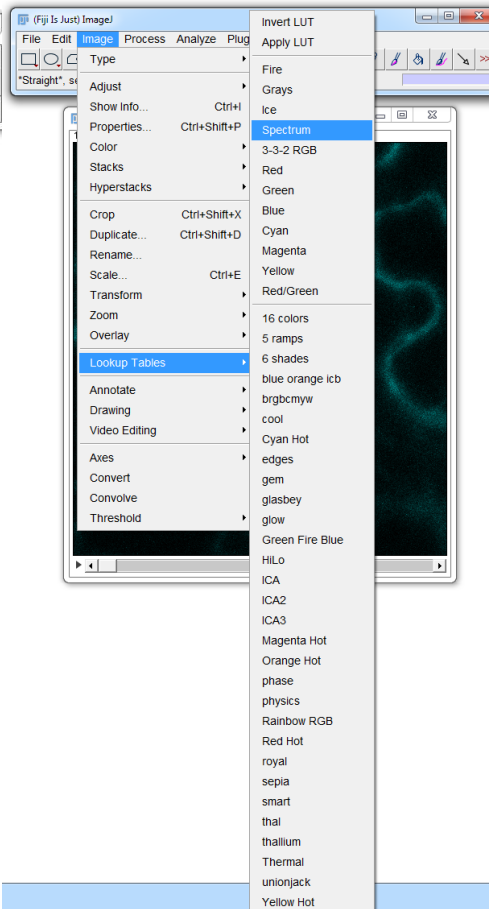
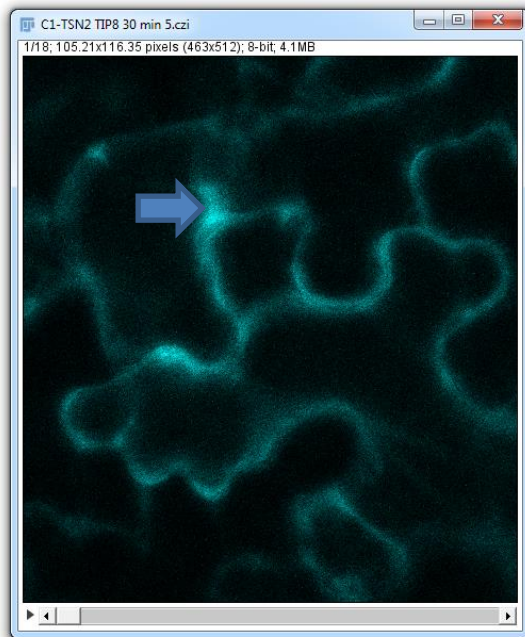
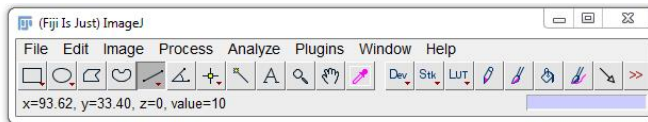
Försters resonance energy transfer (FRET)

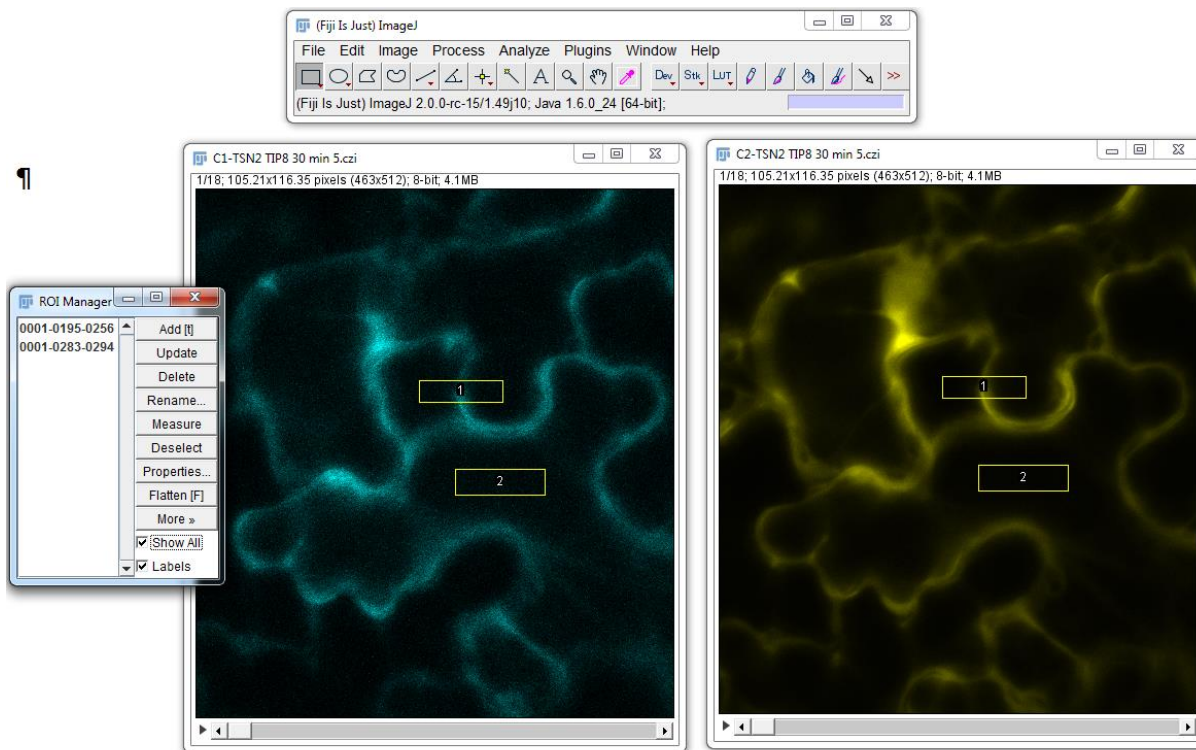
433 nm



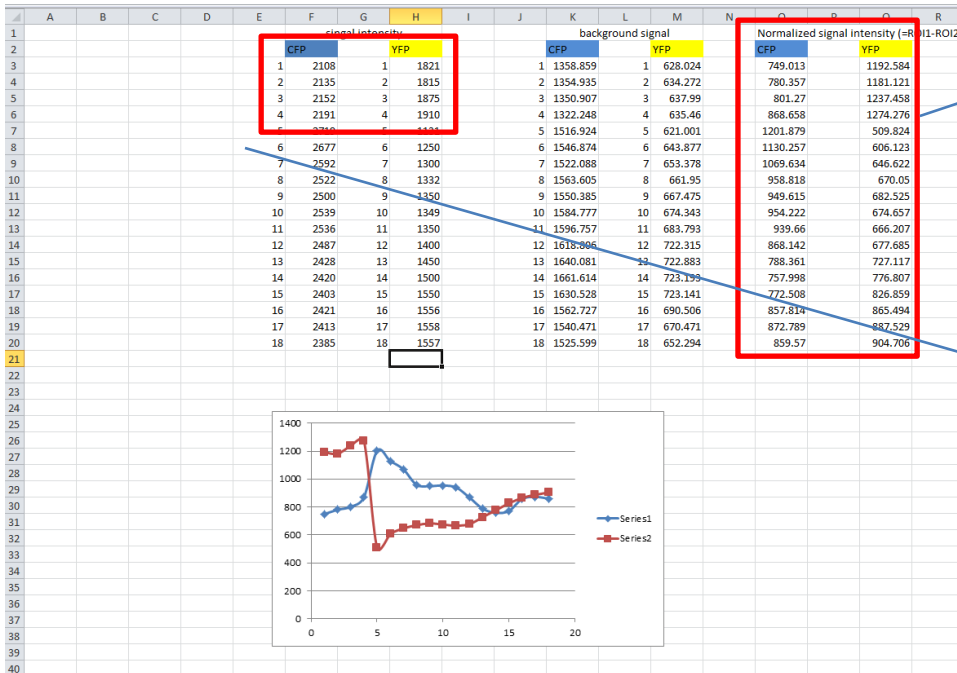
Acceptor photo-bleaching







Όχι σημαντικές διαφορές
στην ένταση



Γράφημα

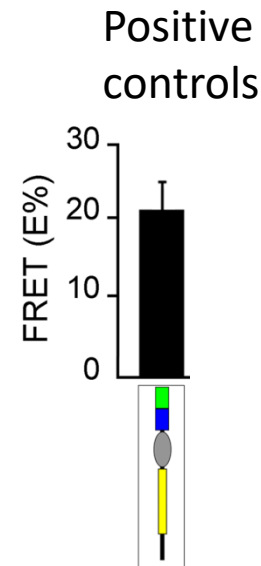
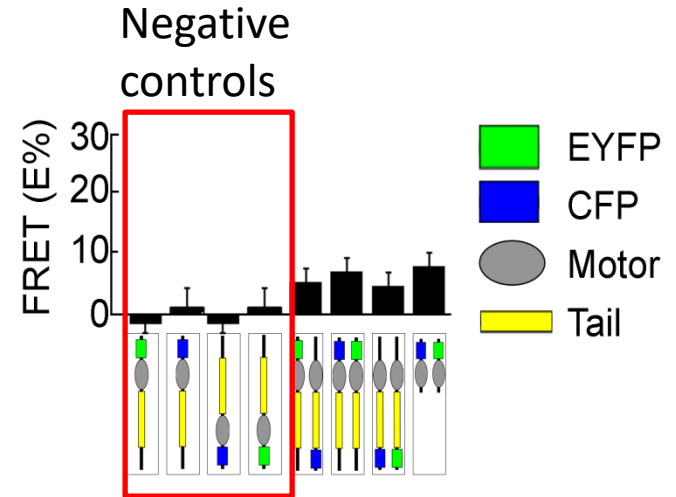
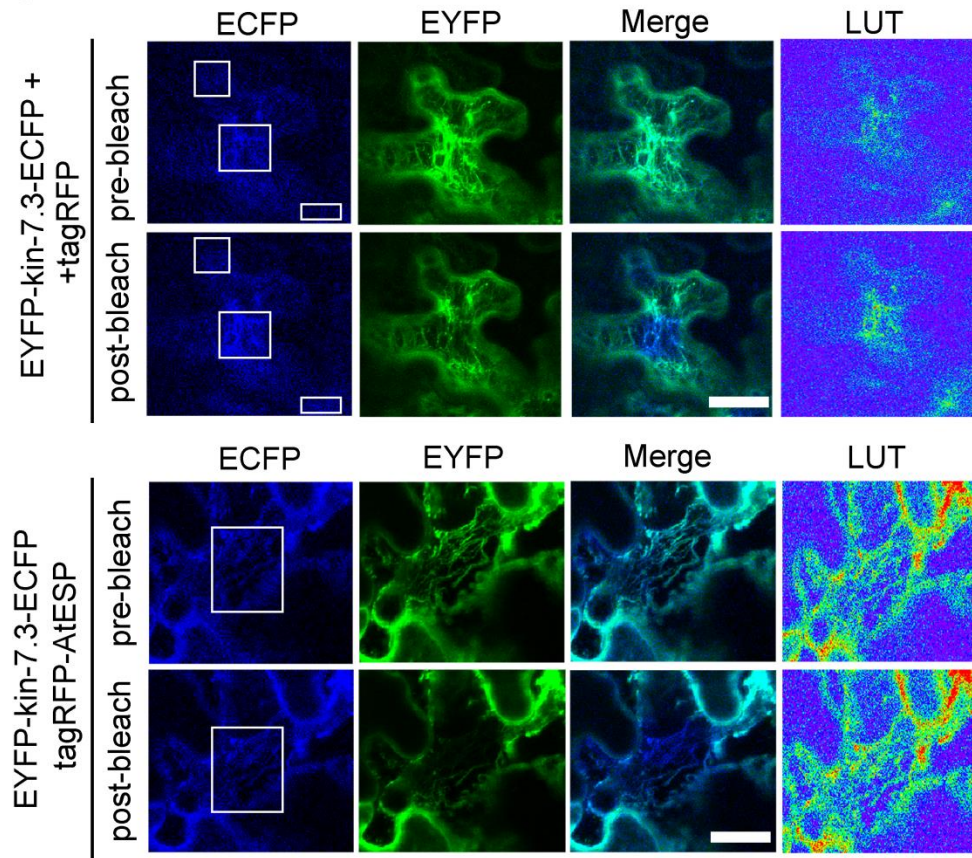
Keep in mind that FRAP takes some time (depending on the iterations). Therefore, I should always check that the increased intensity is not due to 'relaxation' of the CFP

Υπολογισμός απόδοσης κατά FRET

$$Efret(\%) = (\text{Τελική ένταση} - \text{Αρχική ένταση}) / (\text{Τελική ένταση})$$

	A	B	C	D	E	singal intensity		I	J	background signal		N	Normalized signal intensity (=ROI1-ROI2)		T
						CFP	YFP			CFP	YFP		CFP	YFP	
1															
2															
3					1	2108	1 1821		1	1358.859	1 628.024		749.013	1192.584	Εfret(%)= =(O7-O6)/O6
4					2	2135	2 1815		2	1354.935	2 634.272		780.357	1181.121	
5					3	2152	3 1875		3	1350.907	3 637.99		801.27	1237.458	
6					4	2191	4 1910		4	1322.248	4 635.46		868.658	1274.276	
7					5	2719	5 1131		5	1516.924	5 621.001		1201.879	509.824	
8					6	2677	6 1250		6	1546.874	6 643.877		1130.257	606.123	
9					7	2592	7 1300		7	1522.088	7 653.378		1069.634	646.622	
10					8	2522	8 1332		8	1563.605	8 661.95		958.818	670.05	
11					9	2500	9 1350		9	1550.385	9 667.475		949.615	682.525	
12					10	2539	10 1349		10	1584.777	10 674.343		954.222	674.657	
13					11	2536	11 1350		11	1596.757	11 683.793		939.66	666.207	
14					12	2487	12 1400		12	1618.806	12 722.315		868.142	677.685	
15					13	2428	13 1450		13	1640.081	13 722.883		788.361	727.117	
16					14	2420	14 1500		14	1661.614	14 723.193		757.998	776.807	
17					15	2403	15 1550		15	1630.528	15 723.141		772.508	826.859	
18					16	2421	16 1556		16	1562.727	16 690.506		857.814	865.494	
19					17	2413	17 1558		17	1540.471	17 670.471		872.789	887.529	
20					18	2385	18 1557		18	1525.599	18 652.294		859.57	904.706	
21															

Πώς παρουσιάζω δεδομένα FRET



Επιπλέον διάβασμα

- <http://fiji.sc/Cookbook>
- [http://fiji.sc/Time Stamper](http://fiji.sc/Time%20Stamper)
- [http://fiji.sc/Annotating Images](http://fiji.sc/Annotating%20Images)
- <http://occm.otago.ac.nz/resources/Making-a-Look-Up-Table---LUT.pdf>
- [http://fiji.sc/Image Intensity Processing](http://fiji.sc/Image%20Intensity%20Processing)
- Abcissa software <http://rbruehl.macbay.de/>
- Θεωρία FRAP (Balinski paper)
<http://jcs.biologists.org/content/114/21/3885.full.pdf+html>