

Mshot

Digital Imaging Analysis System

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Fluorescence imaging functions

Routine functions

Histogram

Merge channels

Shifting correction

Featured function

Dynamic multi-image merge

Other functions

Split RGB channel

Graying

Line profile

Quicky RGB dye

Coming functions: Auto image splicing ,Auto extend depth of field

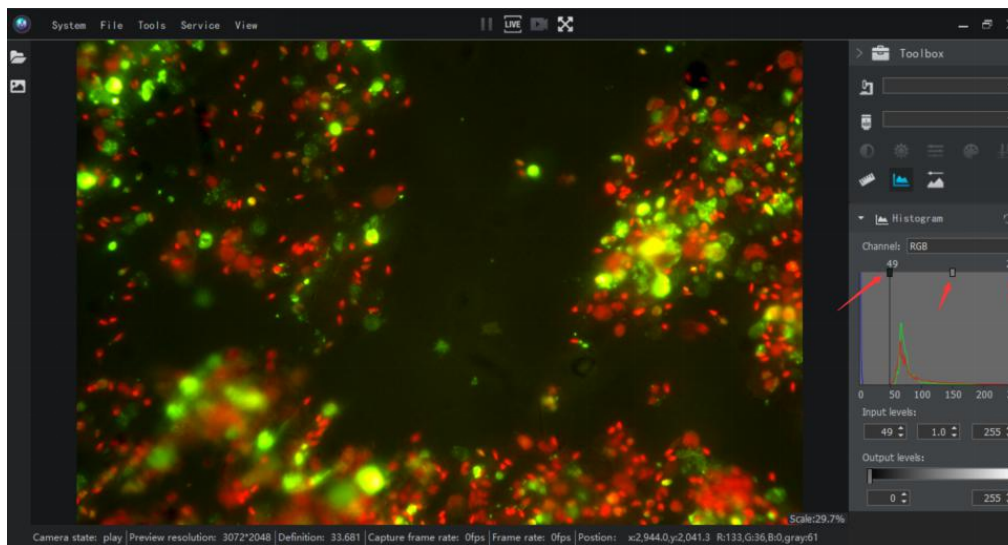
How to get a good fluorescence image?

1. Open software - 2.double click the fluorescence image - 3. Choose 'Histogram' Tool

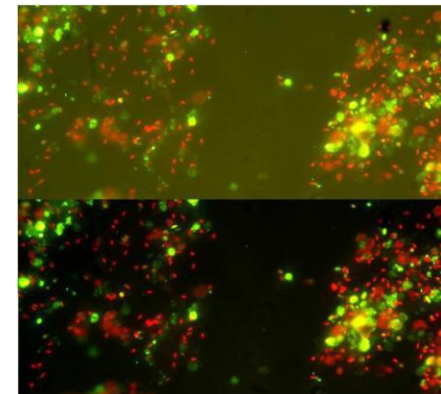
Adjust 'Min. value' from 0 to bigger - get darker background

Adjust 'Max. value' from 255 to smaller - get brighter fluorescence signal

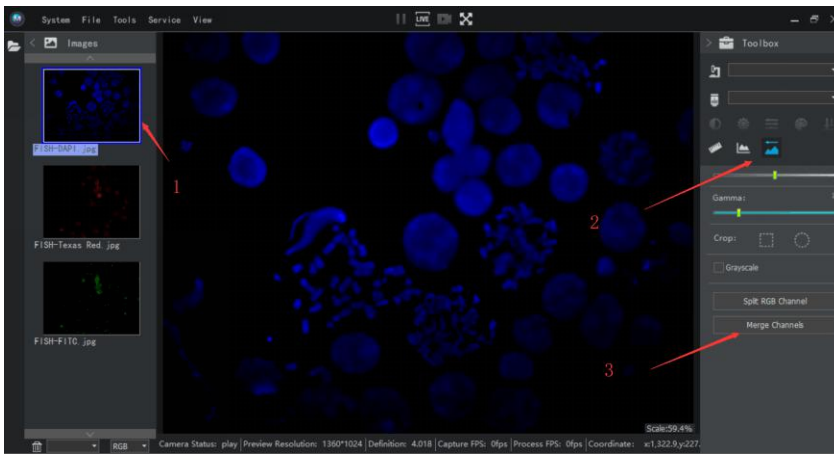
(Midline bard does not need to change in general, input level does not need to change)



Before

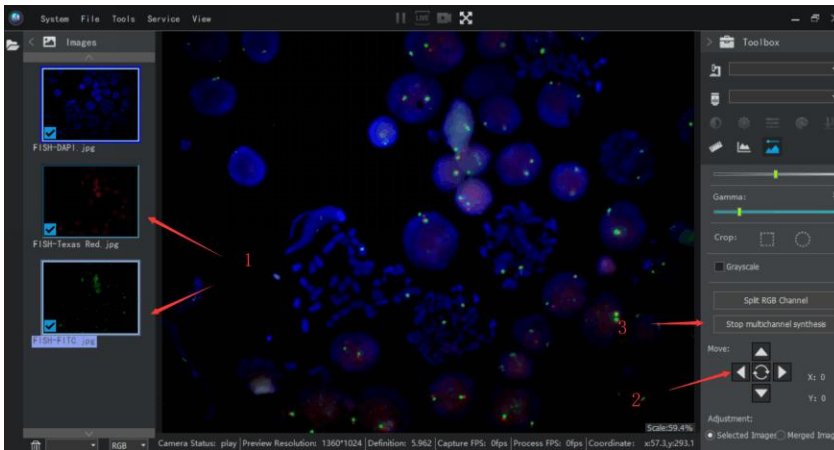


After



Merge channels

1. Choose image - 2. Choose 'Static image processing' - 3. Choose 'Merge Channels'

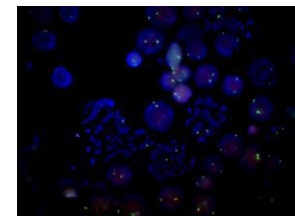
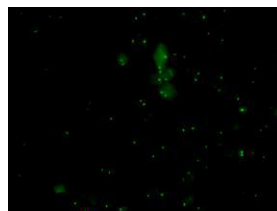
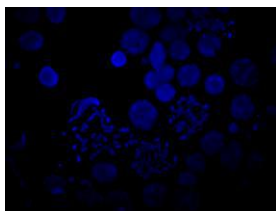


Merge channels

1. Add images - 2. shifting correction (if needed) - 3. Stop merge to save image.

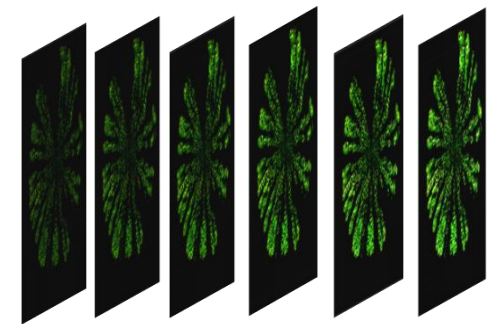
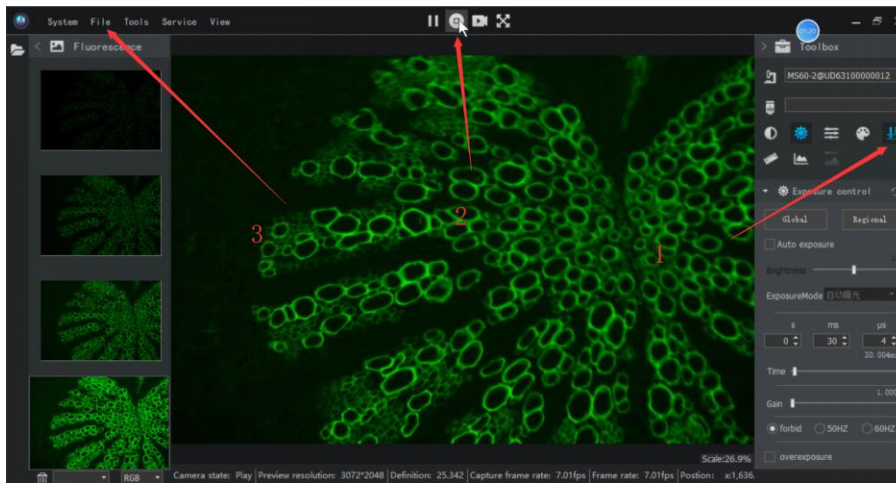
*Max. 5 images in total,

*Merging of phase contrast, bright field with fluorescence are workable



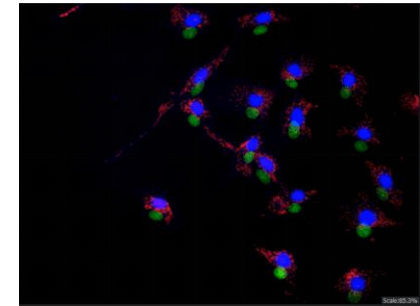
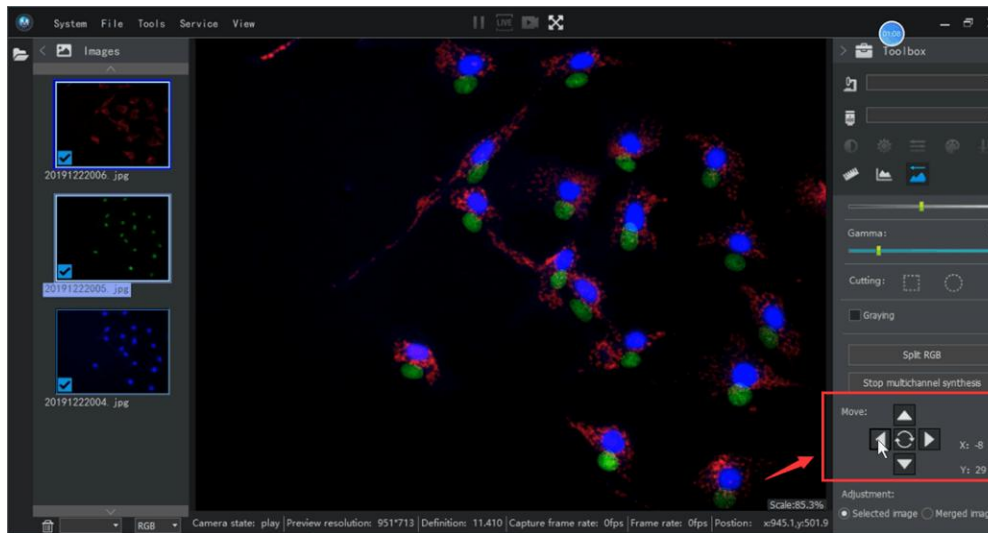
Featured function: Dynamic multi-Images merge (Turn weak fluorescence to bright fluorescence)

- In time merging max. 7 different exposure images to one better image, reduce image noise image.
- 1. Choose ' Dynamic multi-image merge' tool - 2. Click ' Capture' button - 3. Open ' File' Save well done image

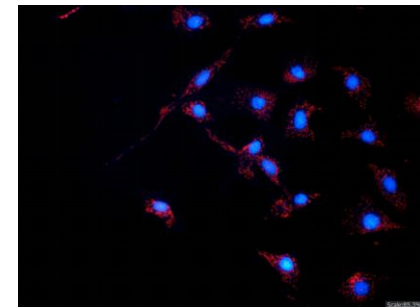


Shifting correction

- Different fluorescence dye images of one specimen might be out of original position because external move and microscope quality, we call it shifting, the tool can move any image position you want to correct shifting.



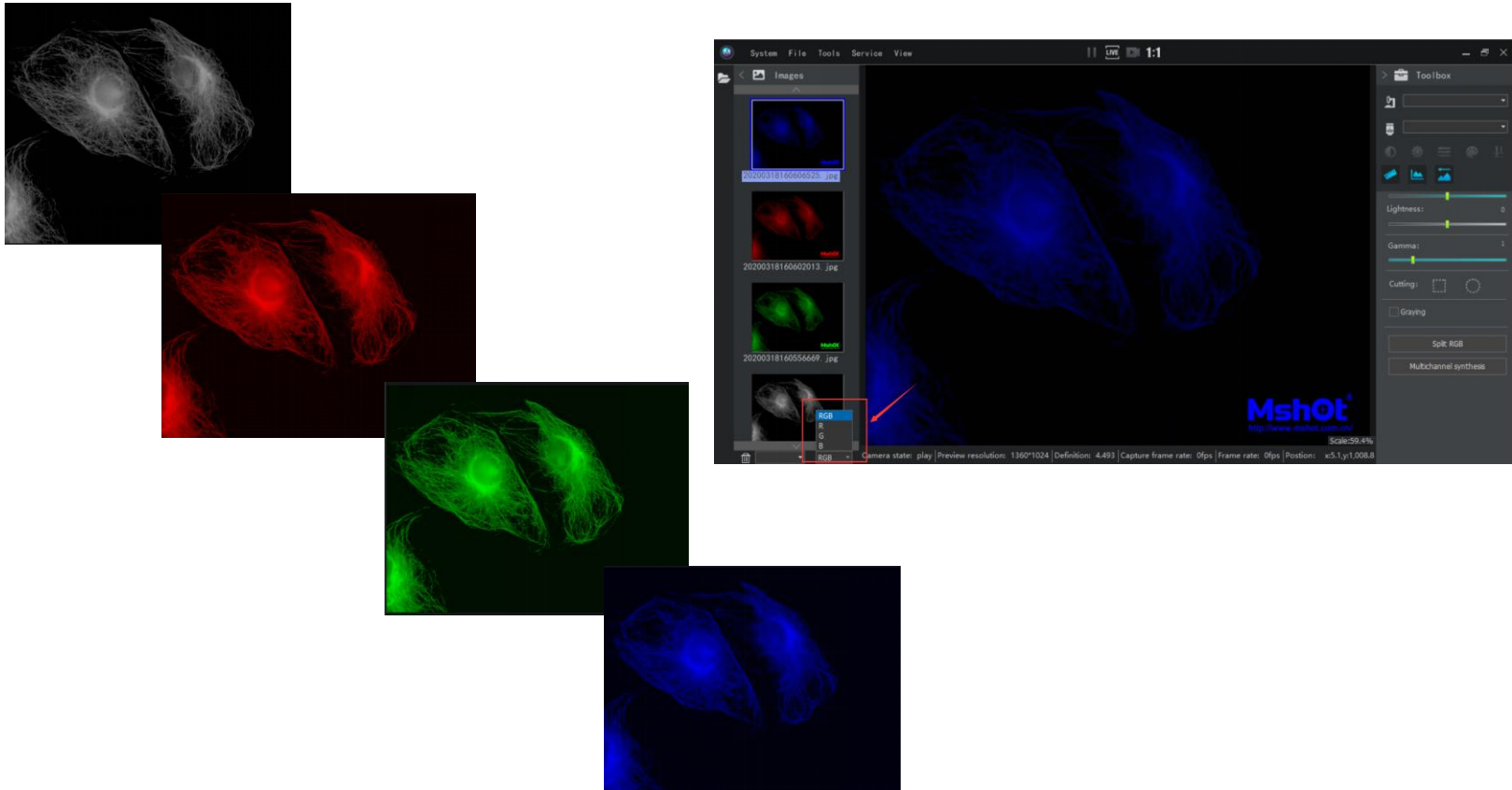
Before



After

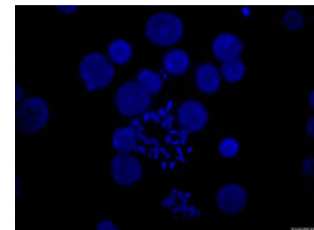
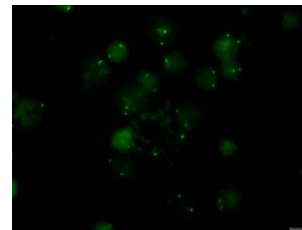
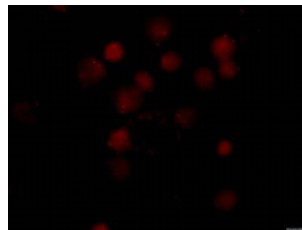
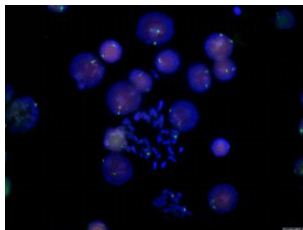
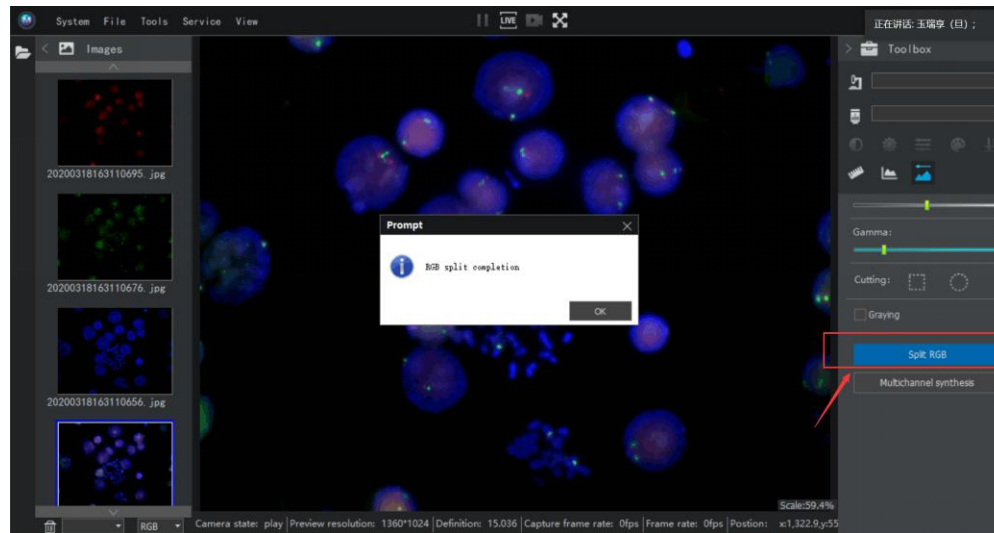
Quickly dye

- Just choose R/G/B channel to dye the monochrome fluorescence image for quickly observation.



Split RGB

- One-push split a multi-channel fluorescence image into single channel images by Red, Green and Blue to quickly separate different fluorescence signal.



Line profile

- Show light intensity of different specimen area.

