

***MEDIA  
CYBERNETICS, INC.***

***User Manual  
Version X3***

***From Images to Answers***

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# Installation, Transferring a License, Uninstalling, and Hardware Requirements

## Sales Information

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## Software and Hardware Requirements - Windows Systems

AQX supports the following operating systems:

- Windows XP 32-bit
- Windows 7 32- and 64-bit

You will also need:

- **Minimum Requirements:**
  - Processor: Intel dual core processor
  - RAM: 3 GB Memory
  - Free Disk Space: 2GB on installation driver + free space for images (20GB+)
  - OS: Windows XP (32) or 7 (32 or 64)
  - Graphics Card: NVIDIA Graphics Card (for 5D Viewer performance)
- **Recommend Requirements:**
  - Processor: Intel quad core 64 bit processor (current recommended model,

Core i7 2600)

- RAM: 8 GB Memory
  - Free Disk Space: 2GB on installation drive + dedicated data drive (500GB+)
  - OS: Windows 7 (64 bit)
  - Graphics Card: NVIDIA Quadro 600
- **Super Number Cruncher:**
  - Processor: Dual Intel quad core 64 bit processors (current recommended model, 2X Xeon 5600 series)
  - RAM: 48GB or higher

- Free Disk Space: 2GB on installation drive and one or more dedicated SATA 6Gb/s data drives (2TB+)
- OS: Windows 7 (64 bit)
- Graphics Card: NVIDIA Quadro 4000

Larger memory sizes are recommended for improved performance, keeping in mind that the operating system, the window interface and potential tasks of other concurrent tasks make strong demands on memory resources.

The sample 3D images that are supplied with the software require approximately 350MB of disk storage. It is recommended that due to the inherently large size of 3D images, the images be stored on large volume expandable media such as CD-ROMs, rewritable CDs or DVDs. For routine use, a disk drive of capacity 2GB or larger is recommended.

The Volume Projection - Hardware selection in AutoQuant 5D Viewer requires a video card which supports OpenGL extensions. It is recommended that an ATI Radeon 9600 series or higher be used for the 3D Visualization. Following is a non-inclusive list of consumer level video cards which support OpenGL coding:

- ATI Radeon 8500, 9000, 9700
- Diamond FireGL1
- nVidia GeForce 256, GeForce 2, GeForce 2 MX, GeForce 3, GeForce 4, all Quadro lines
- 3D Labs Oxygen GVX1 and Wildcat family
- Matrox Parhelia

## Software Installation

Note: Installing AQX on Windows Vista from a guest account is **not** supported. Install from an account with Administrator privileges, or as a restricted user.

To install complete the following procedure:

- (1) Unplug all USB dongles, then insert the software CD-ROM into your CD-ROM drive.
- (2) If the software asks if you want to install Microsoft .NET framework 2.0, select *Yes*. If it does not ask this, it is because it is already installed on the computer.
- (3) Follow the Installation Wizard's onscreen prompts to install AutoQuant X/5D Viewer.
- (4) When the Installation Wizard states that AutoQuant X/5D Viewer has been installed, click *Finish*.
- (5) A series of HASP device driver dialogs will pop up informing you that it is uninstalling and reinstalling HASP drivers. Click *OK* through them. This will ensure that the dongle drivers are up to date.



(6) Plug in the USB dongle and wait for Microsoft Windows to notify you that the new hardware is ready to use.

(8) Double click the AutoQuant X3 icon on the desktop or in the Start menu.

*Note: If AutoQuant X3 does not automatically start the Installation Wizard, browse to the CD-ROM drive and double click setup.bat.*

You can find the installed folder by looking in the following directories:

**C:\Program Files\MediaCybernetics\AutoQuant X3 .**

## **Dongle Installation and Requesting a Permanent License**

All Media products will run for one month following their initial installation on your computer. During this time you will need to contact Media for a permanent license so that you can continue using your software beyond the initial trial period. Once you have purchased a permanent license, you will receive a CD and dongle, at which point you can follow the instructions below.

### **Updating the Dongle**

If you have purchased additional functionality, or an extension to your maintenance or license agreement, you will need to update your dongle to reflect the changes in your license.

- (1) Click Start->Programs->Media Cybernetics->AutoQuant X3->Dongle Update.
- (2) You will be presented with the dongle update screen.
- (3) Contact Media Cybernetics Inc., Provide the User Registration ID and Dongle ID shown in the Dongle Update Utility dialog.
  - By Email: [customerservice@mediacy.com](mailto:customerservice@mediacy.com)
  - By phone: 301-495-3305
- (4) If the change that must be made has been approved, you will receive an update code in the form of a long string of letters and numbers. Enter this string into the Dongle Update Code text box and click Update Dongle.
- (5) Upon successful update, you will receive confirmation, and the update will take effect the next time you run your AutoQuant product.

## **Starting the AutoQuant Software - Windows Systems**

To start:

- (1) Choose the Start button at the lower left-hand corner of your screen and select Programs.

- (2) In the Media Cybernetics Folder, select the name of the application.

## **AutoUpdate**

Media's products incorporate an AutoUpdate feature, which will once a month connect to the internet to search for product updates automatically. If you wish to search for an update without waiting for the default time to pass, you can select Search for Update from the Help menu. See the Help chapter for more information on the AutoUpdate feature.

## Uninstalling the Software

Go to Add/Remove Programs under the Control Panel of your Operating System. The Add/Remove Programs box will list all currently installed programs. Find and click on the name of the program you wish to remove. Once the name of the program you want to remove is highlighted, click on the Change/Remove button. The InstallShield Wizard will remove the program from your computer.

## How to Use This Manual

The Tutorials of the manual go through every menu item in the Menu Bar.

Depending on the product you have purchased (*AutoQuant*, *AutoQuant 5D Viewer*, *FRET*, *Ratiometrics*, *etc.*) some sections may not apply to you (this will be clearly indicated at the beginning of the corresponding tutorial).

# Guidelines for Collecting 3D Image Datasets

## Overview

For transmitted-light brightfield and widefield-fluorescence microscopy, the main considerations are the following:

- Collection of the optical sections
- Collection of bias frames
- Collection of the flat field frame
- Vibration control

It is important to follow the guidelines outlined in the following sections. The adage of “garbage-in garbage-out” is true for following these guidelines. Even though *AutoQuant* is relatively robust against non-adherence to any of these guidelines, with each guideline that is not followed, and depending upon how severely the guideline is broken, the microscopist increases his/her chances of having unsatisfactory experimental results.

It is best to follow each and every guideline, to the letter, wherever possible. Do not violate a guideline for the sake of convenience. Break only those guidelines for which you have no choice but to break, because of restrictions caused by your experimental conditions. Remember that with each guideline broken, the risk of having unsatisfactory results increases.

## Optics Alignment

Before beginning any procedure, be sure that the entire optical train is well aligned. Follow the instructions provided by your microscope manufacturer on proper adjustment and alignment of the optical elements. This should be performed daily, prior to each experiment, and should be re-checked throughout the day if the microscope is receiving heavy usage. Make certain of the following:

-The fluorescence excitation lamp is properly centered and focused (if fluorescence is being used).

-The trans-illumination lamp is properly centered and focused (if transmitted-light brightfield is being used).

-The condenser is properly centered and focused (if transmitted-light brightfield is being used).

-For transmitted-light brightfield: Adjust the condenser iris diaphragm (numerical aperture, NA) according to the following:

For 3D digital imaging, as performed here, it is best to have the transmitted-light be as incoherent as possible. This implies opening the NA, as far as possible.

For direct eye viewing, on the other hand, it is best to eliminate scattered and stray light in order to improve contrast, and this implies closing the NA to a point just below the NA of the objective lens. **DO NOT DO THIS!** As a rule of thumb, it is recommended to simply open your condenser NA as far as it will go. If doing so makes viewing the sample overly difficult, then lower the condenser NA, until it is at about 1.2 (20% higher) of the objective lens NA. If you are using an oil objective lens having an NA that is greater than 1.0, and using a condenser that allows usage of oil, then you will need to do so (i.e., you will need to place a drop of oil on the condenser) in order to ensure this condition. If you are using an oil objective lens for which the NA is higher than the highest available NA of the condenser (for example, a 1.4 NA objective lens and a 1.25 NA condenser), then use the highest available NA that your condenser will allow and use a drop of oil on the condenser (assuming your condenser allows you to use oil with it).

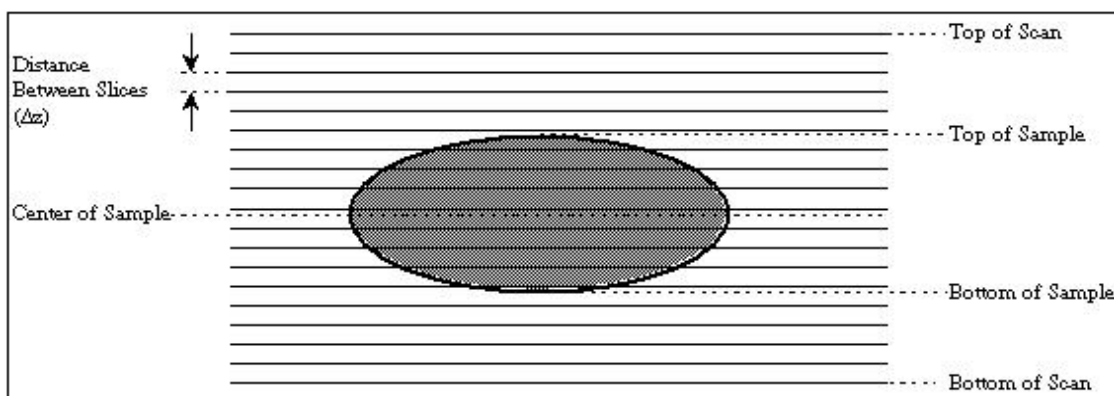
## Collecting Optical Sections

### Setting the Top and Bottom of the Sample, and of the Scan

You need to select the top and bottom slice of the sample. By performing a through-focus operation, with either direct-eye or video camera viewing, you should judge where the top and bottom of the sample are.

*Note: The top and bottom of the scan are not necessarily the same as the top and bottom of the sample.*

The rule of thumb for selecting the top and bottom of the scan is described below, using the terms illustrated in Figure 1 below.



**Figure 1:** Shows the geometry, and illustrates the terms used for 3D widefield dataset collection.

*Note: The actual axial distance that is scanned is about twice that of the sample. The top and bottom are over-scanned by approximately half the thickness of the sample. The above rule of thumb*

*provides the best results. If you have limited disk or memory space, then you may not be able to follow this rule of thumb. In that case, scan the depth of the sample, plus any additional amount allowed by your free hard disk space.*

## Setting the Top and Bottom of the Scan

For best results, the rule of thumb in setting the axial distance between slices is to have this distance equal to the Rayleigh depth of field (DOF) as given by the following equation:

$$\Delta Z_{\text{slice}} = \text{DOF} = \frac{\lambda}{4\eta \sin^2(0.5 \sin^{-1}[\text{NA}/\eta])}$$

where  $\lambda$  is the wavelength (if unknown, use a value of 0.5 micrometers),  $\eta$  is the refractive index of the immersion medium (1.0 for air, 1.33 for water, 1.515 for oil) and NA is the numerical aperture of the objective lens.

The total number of slices required, then, will be equal to the thickness of the scan, selected as described earlier, divided by the chosen axial distance between slices. If you have limited disk storage space, you may not be able to follow the above rule of thumb. In that case, you may set your axial distance between slices equal to a number larger than the depth of field. ***AutoQuant*** will still work in that case. However, as a trade-off, you can expect to experience some degradation in image quality. The extent of the degradation will depend upon the axial sampling distance chosen relative to the DOF.

Deblurring is best carried out when the datasets are seemingly oversampled. In other words, best results are achieved when the pixel or optical section spacing is finer than what would be used normally, without deblurring. In-plane deblurring is most successful when the sample spacing is finer than 1/2 the Rayleigh width. If in-plane deblurring is desired, sample at 0.1  $\mu\text{m}$ , or finer, when the 1/2 Rayleigh width is  $\sim 0.25 \mu\text{m}$ . Axial deblurring is substantial with widefield optics regardless of the z sampling. The rule of thumb for confocal datasets, without deblurring, is to have the optical sections spaced according to the confocal spot size along z, which is the resolution element along the z-direction calculated by the following equation [Webb et al, 1995]:

$$\Delta Z = \frac{(1.4\lambda\eta)}{\text{NA}^2}$$

where  $\lambda$  is the wavelength in microns ( $\mu\text{m}$ ),  $\eta$  is the refractive index of the immersion medium, and NA is the numerical aperture of the objective lens. If  $\Delta z$  is 0.5  $\mu\text{m}$ , sample at 0.5  $\mu\text{m}$  or finer per slice. The noise suppression mentioned above requires this type of fine sampling.

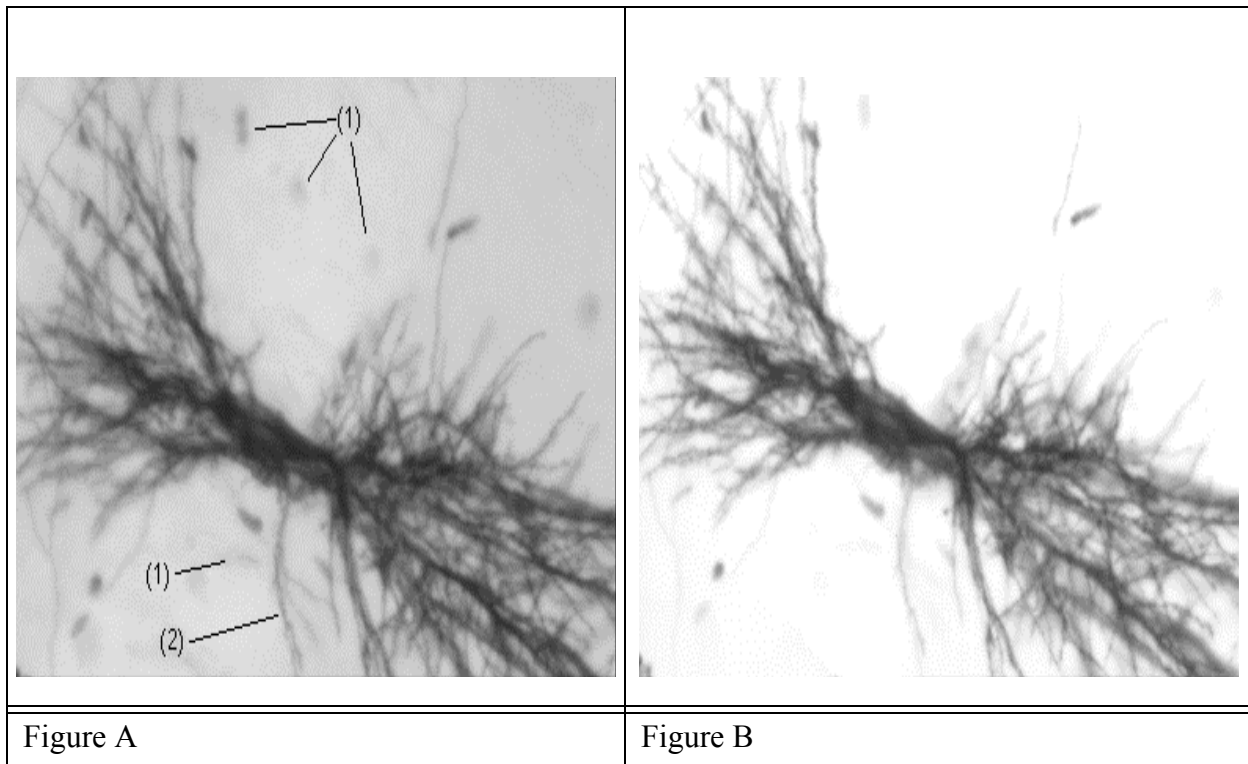
## Setting the Exposure, Gain and Offset

### Cooled CCD Cameras

With cooled CCD cameras, the exposure time is usually adjustable. Consult the manufacturer's manual to determine how to change the exposure time. To set the exposure time for collecting optical sections, use the following procedure.

Focus onto a point near the center of the sample (see Figure (1)). Adjust the illumination light to a setting that you desire. If you are using a transmitted-light brightfield microscope, adjust the light so that you can view the sample comfortably through the eyepiece. Next, try an exposure time setting. Usually, if you have done this before, you can start with a setting that has worked well for you in the past. If you are doing this for the first time, or if an approximate setting is not known, start with an exposure time around 0.05 seconds. Grab and view a single frame at this setting. If the picture on your screen appears too dark, increase the exposure time and repeat the frame grabbing and viewing procedures. Be careful not to saturate (make too bright) any portions of the image. Saturated regions can usually be identified as bright, flat, white regions in the picture, as seen in Figure (B).

**Figure 1-1**



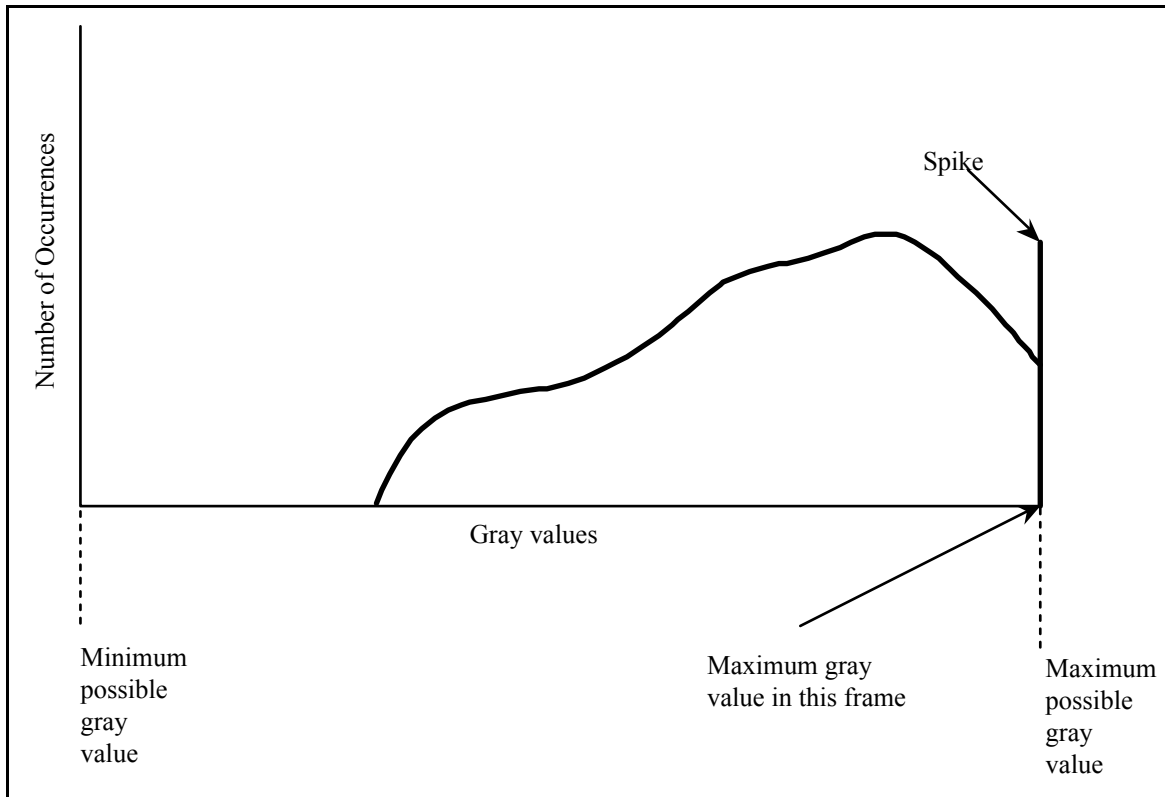
**Figure (A)** A successful, unsaturated image

**Figure (B)** A saturated image of the same sample. Note the obvious bright, flat, white background. Note that the highlighted background features (marked '1') in Figure A are erroneously



eliminated in the saturated image. Note that other features of the sample (marked '2') are eroded in the saturated image.

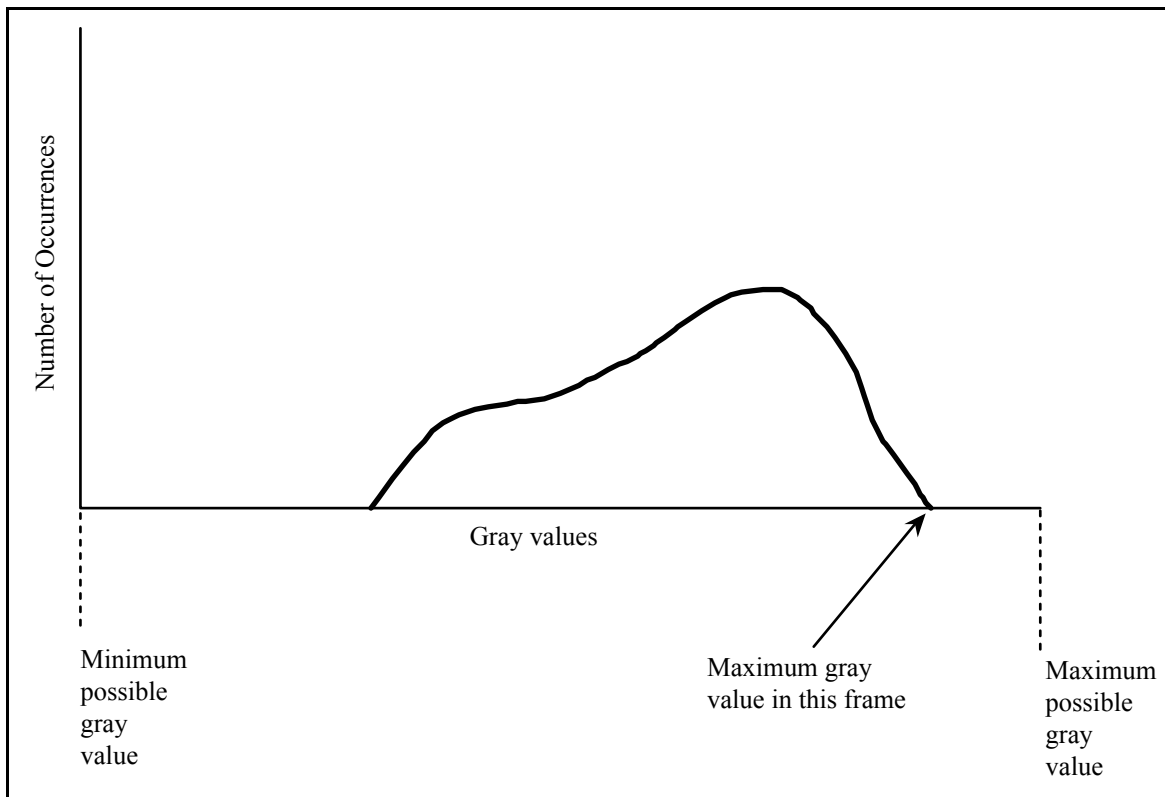
A reliable method for detecting saturation is to compute and display the image intensity histogram. A straight vertical spike/line at the end of the abscissa (see illustration in Figure C) is an indication of saturation, which is to be avoided.



**Figure C:** Histogram of gray values in an image frame that has saturated pixels. The maximum gray value of the frame and the maximum possible gray value coincide. Typically, though not always, a spike will appear in the histogram, as shown above.

If you observe saturated pixels, lower the exposure time and repeat the grabbing and viewing procedure.

As a rule of thumb, adjust the exposure time so that the brightest pixel is at approximately 75% of the well capacity of the CCD camera. This condition is often detectable (depending upon the sample), again, by using a histogram. The 75% well capacity condition is detected by noting that the highest abscissa value having a non-zero histogram value is at 75% of the range of the abscissa. An example of this is shown in Figure D below.



**Figure D:** Histogram of gray values in an image frame with no saturation. Adjust the camera/frame-grabber gain and/or illumination level such that the maximum gray value in the image frame is between 75% to 85% of the maximum-possible gray value. For instance, for an 8-bit camera (most RS-170 camera/frame grabbers), the maximum gray value is 255. For a 12-bit cooled CCD camera, the maximum possible gray value is 4095.

If your cooled CCD camera provides gain adjustments or frame averaging, follow the manufacturer's directions to achieve the above-described intensity histogram profile.

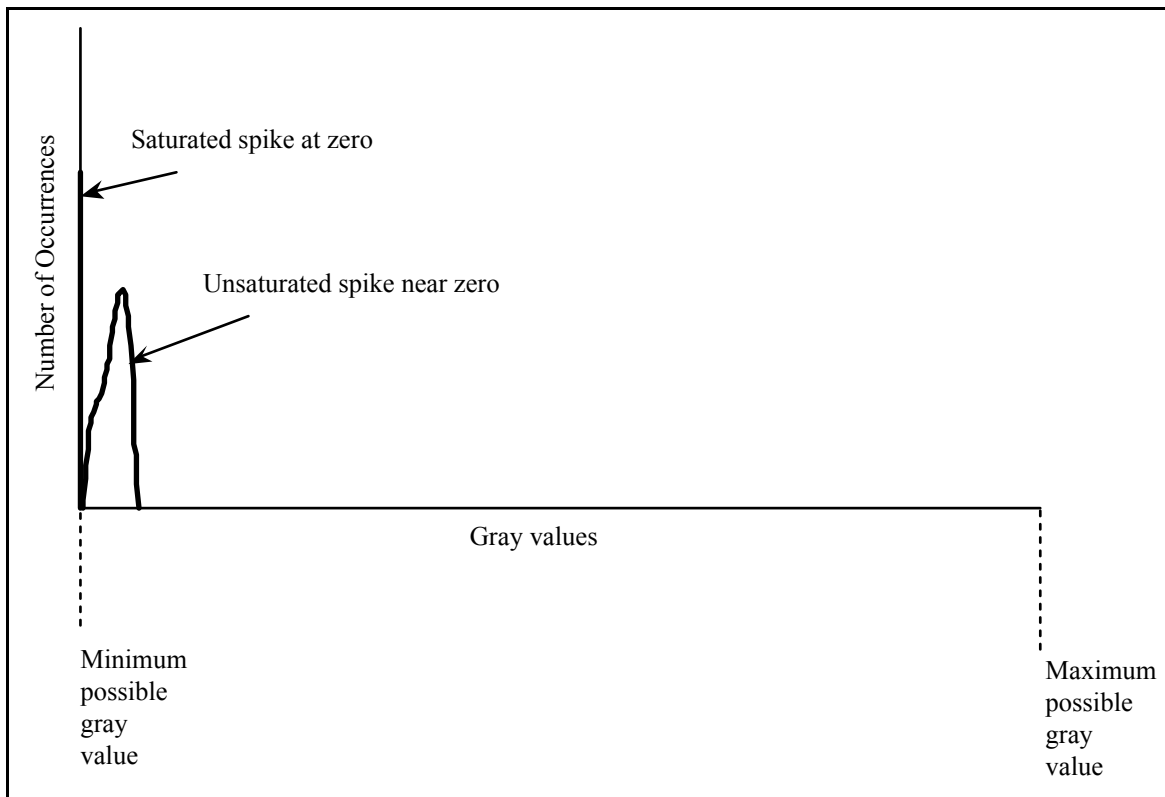
## RS-170 Cameras

Many RS-170 cameras provide some control over the video **gain**. Some cameras provide a gain adjustment device on the camera itself while others provide this adjustment as part of a camera electronic control box. Often, frame grabber cards provide a software selectable gain factor. Finally, the lamp brightness control and the neutral density excitation filter control provide yet two more mechanisms for selecting the gain. Please look through the manuals provided by your microscope, camera and frame grabber manufacturers to determine which if these options are available to you, and methods to access them.

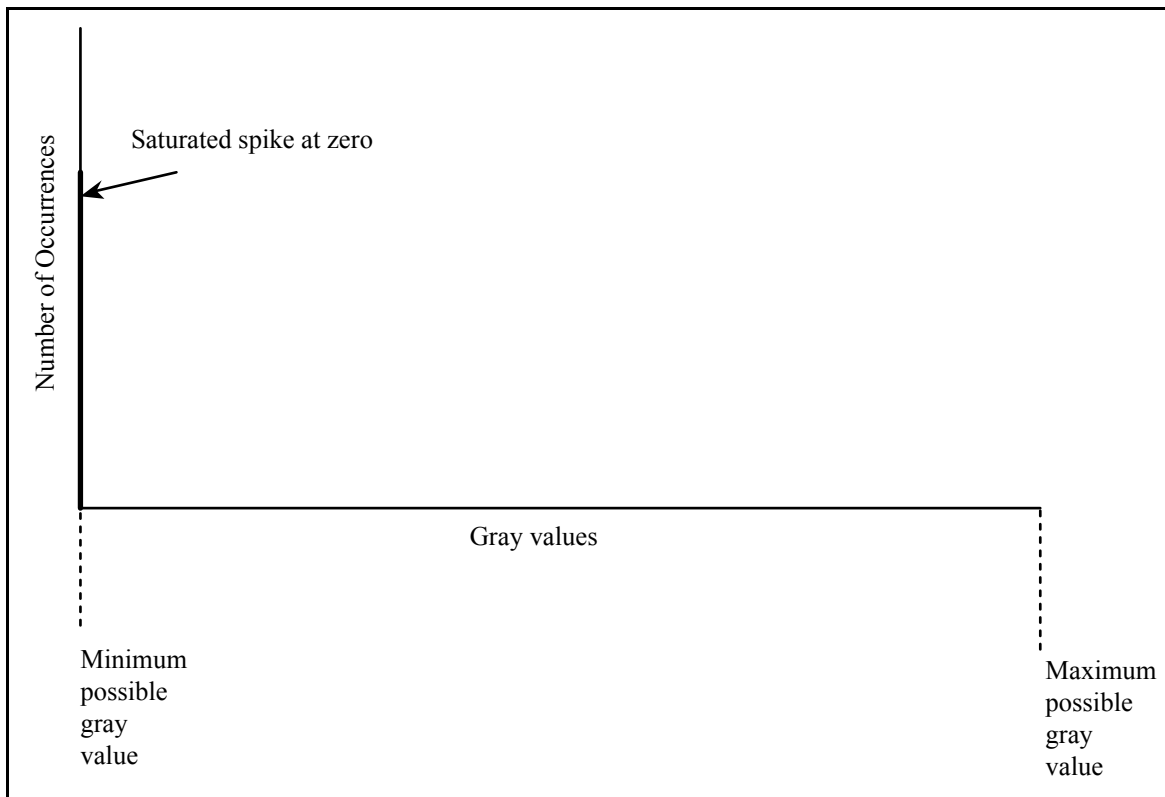
Similarly, cameras and/or frame grabber cards provide an **offset** (black level) adjustment mechanism. Again, please refer the manufacturers' manuals on methods to access the offset adjustment.

Once you have determined the above information, the following procedure is recommended for setting these adjustments.

- (1) Set the illumination light intensity (for transmitted light brightfield) or the neutral density filter (for fluorescence) first. Do this by direct eye viewing and selecting the intensity so you can comfortably view the sample through the eyepiece. If you are trying to maintain a very low light level, to minimize photobleaching with a fluorescence sample, this might not be possible. Instead you may have to simply set the light level to a suitably low level which cannot be seen by eye.
- (2) Set the offset value on the video camera and/or frame grabber. Ideally, you want the offset to be adjusted so that a zero light level corresponds to a zero value on a histogram of gray values. Block all light to the camera. This may be accomplished either by turning the lamp power switch off (for transmitted-light brightfield) or by closing the excitation shutter (for fluorescence). It may also be accomplished by sliding the position of your dichroic mirror filter set. Next, adjust the offset, or black level position(s) on your camera and/or frame grabber card. If possible, to keep matters simple, adjust only the setting of the camera and leave the rest of the system alone. Make this adjustment so that you see a spike in the gray-level histogram right at the zero (left most) value of the abscissa, as shown in Figure E. Avoid a completely saturated histogram as shown in Figure F. This is identified by having only a single spike at the abscissa value of zero and having no spike of finite width.
- (3) Set the gain. As mentioned, you may have essentially three independent ways to control the gain (camera, frame grabber, and light intensity/neutral density filter). To keep matters simple, try adjusting only one of the gains. If fluorescence is being used with an intensified camera, we recommend adjusting only the camera gain, keeping the excitation light as low as possible. If transmitted-light brightfield is being used with an ordinary video-rate CCD camera, such as an RS-170 camera, we recommend adjusting only the illumination light level. Open the light to the camera. Adjust the gain (or illumination level) setting until the highest gray level (abscissa) on the histogram, which has a non-zero value (ordinate), is at about 75% of the range of the abscissa, as shown in Figure D. Avoid/minimize saturated pixels using the procedures described earlier in this section.



**Figure E:** Histogram condition for setting the offset (black level) of the camera/frame grabber. Look for an unsaturated spike at or near zero. There may also be a saturated spike at zero.



**Figure F:** A zero-saturated histogram. This condition is to be avoided when setting the offset (black level).

## Performing the Axial Scan

Begin grabbing images at each optical section. Start by focusing onto one end of the sample. Grab a frame. Do this with frame averaging if this feature is available on your setup. This is especially important if the datasets are noisy due to low light levels. It is recommended that you average 255 frames if possible. Refocus the microscope to the next adjacent frame and grab the next image, and so on. Refer to file formats that are compatible with *AutoQuant*. If a TIFF format is used, for instance as with some standard frame grabber cards, store each frame in a file, in sequentially numbered files with a numeric suffix, i.e., files with names of the form IMAGE.1, IMAGE.2, and so forth. If a raw 16-bit integer format is used, for instance as with some of the cooled CCD cameras, then store all of the frames in sequential order and contiguously in the same file having an extension .deb. Repeat this image-grabbing and storing sequence until you have scanned the entire sample. Scan the entire depth that was chosen according to Figure 1.

After collecting the datasets, check the computer's directory for the file names that were chosen, as a diagnostic check to be certain that all of the datasets were saved. Check for their expected file sizes in numbers of bytes. For instance, if a raw 16 bit binary format is used, as with a cooled CCD camera, and if the image size is 1024 by 1024 with 85 slices, then the number of bytes shown in the directory for this file ought to be  $1024 \times 1024 \times 85 \times 2 = 178257920$ .

Be careful to move the plane of focus the exact same distance for each intermediate frame. Check the specifications of your microscope and identify the amount of movement effected by a revolution of the fine focus knob (i.e., the micrometers per revolution). For example, for the Olympus BH-2, the movement is 200 micrometers per full revolution of the fine focus knob. These are well indicated by graduations on the knob, with the knob having 100 graduations. Thus, the stage will move 2 micrometers for every graduation that the knob is turned.

It is important to have a reliable axial scanning mechanism, with some way of verifying the step sizes. The accuracy of the *average* step size is critical. This should be within 5% of the specified step size. The accuracy of *individual* step sizes is not nearly as critical. As a rule of thumb, these ought to be within 20% of the specified step size, but their *average value* (to emphasize the point) needs to be within 5%. For example, suppose that you specify that the step sizes are 0.2 micrometers, and that the scan will be 50 planes deep. Then, by using a position gauge on the stage or by monitoring the focus-knob graduations, you should record and verify that the stage moved within 5% of the 10 micrometers (between 9.5 and 10.5 micrometers). With a high-resolution gauge having a resolution of 0.01 micrometers you may be able to verify the individual step sizes. However, with lesser expensive gauges, having a resolution of 0.1 micrometers, it will be sufficient to verify the larger step sizes, of 1 micrometer or more. By specifying step sizes of 1 micrometer and higher, verify that these steps are indeed within 20% of the 1 micrometer (within 0.8 and 1.2 micrometers). However, make sure that the deviation from this step size is randomly dispersed about the 1 micrometers so that the *average* step size is still within 5% of the 1 micrometers.

A position measurement device, such as a Mitutoyo (Aurora, Illinois, USA, 708-978-5385) MU-Checker and digital Minich checker having resolutions of 0.1 micrometers, should be used to verify the average step size. Heidenhain sells a number of position gauges as well. A resolution in the measurement of 0.1 micrometers is sufficient, because it is the average of the step sizes that is most critical value. One of the common malfunctions that can occur in dataset collection is gear slippage. The slippage of the microscope gears or slippage of any linkage between the motor controller and the microscope can be a critical error in dataset collection. A position measurement device will check for this error. If a positioning gauge is not available, then the next best way of checking the positions is to record the position of the graduations on the fine-focus controller. Each graduation specifies a certain number of microns. A careful recording of these graduation positions will allow you to identify slippage in the motor controller and in the linkage between the motor and the microscope, but it will not allow you to determine if there is slippage in the microscope gears themselves.

Check with your microscope's manufacturer to determine what type of gear mechanism is used, and if this gearing may be prone to slippage. Regardless of the design of this gearing, any microscope may have problems, since any gearing mechanism may become stripped, and easily so if it experiences harsh usage such as in educational environments. Some microscope models, have ball bearing friction linkages between the focus control and the stage, and thereby are prone to slippage under normal conditions.

There are a number of manufacturers that sell turnkey 3D dataset collection systems, which includes both the needed hardware and software, for widefield microscopy. These products carry

out the above instructions automatically. Some of the manufacturers to check are listed here (Universal Imaging Corporation, Downingtown, PA; Compix, Cranberry Township, PA; MediaCybernetics, Bethesda, MD; Nikon, Melville, NY; Leica, Bannockburn, IL).

## Avoiding Backlash and Hysteresis

While stepping the focus, always scan “against the load.” If you have a stage-focusing microscope (where the stage moves up and down as the focus knob is moved), then always step the focus in the direction where the stage is moving upward. This will cause you to scan the sample in the direction from top to bottom. If you have a nosepiece focusing microscope, then always step the focus in the direction where the nosepiece is moving upward. This will cause you to scan the sample in the direction from bottom to top.

When adjusting the focus from one sequential slice to the next, move the knob very slowly and cautiously. Also be careful to move the knob only in one direction during the scan.

Once you have started grabbing frames, never reverse the direction of the fine-focus knob (not even a miniscule amount, and not even just once!). Doing so by even a very small amount will ruin the scan due to backlash and hysteresis of the focusing system.

To compensate for erroneous hysteresis effects, do the following before grabbing the first frame. First, focus into the specimen. If you have a stage-scanning microscope, first move the stage down by 20 micrometers or more below the position where you wish to grab the first frame. Then, being careful to move the knob in only one direction (as mentioned earlier, once you are in this stage of operation, never reverse the direction of the knob), slowly and carefully move to the position where you want to grab the first frame. This operation counteracts a hysteresis that is inherent in the microscope’s focusing mechanism.

## Collecting Bias and Flatfield Frames

### Cooled CCD Cameras

Bias and flatfield dataset frames are used by *AutoQuant* to automatically calculate the cooled CCD camera’s dark current distribution, pre-amplifier bias and flatfield non-uniformity. This procedure needs to be repeated every day, or preferably, for every dataset collection. This process compensates for potential misalignments in the optics, which may drift from day to day, and to compensate for dust that may settle in the optical train, which also changes from day to day.

Collect two bias frames and one flatfield frame. The bias frames are taken with the camera shutter closed, or with light to the camera blocked. Select a charge integration time for the first bias frame. The accuracy of the bias frame dataset will increase with increasing charge integration time. As a rule of thumb, use 60 seconds for the first bias frame. The charge integration time for the second bias frame needs to be twice that of the first bias frame, so as a rule of thumb use 120 seconds for the second bias frame.

Collect a flatfield frame. This will be used to calculate the flat-field non-uniformity of your camera. A flat field “sample” can be provided in a number of ways. When using transmitted-light brightfield, the simplest way is to remove the sample from the microscope and grab an image of a blank stage. A rather simple method, with transmitted-light brightfield, is to take the biological sample extremely out of focus, to a point where the image shows a completely flat field. When using fluorescence, the most reliable way is to prepare a “flatfield” slide by placing a drop of fluorescent dye onto a microscope slide and sealing it with a coverslip. In case any debris particles may be contained in this droplet, it is best to grab this image with the droplet well out of focus. Some care must be taken to ensure that no out-of-focus remnants of debris appear in the image, and that no other particles on the slide happen to come into focus. See Table 1: Figure (a) and Figure (b) for an illustration of these conditions. In setting the exposure time, when using transmitted-light brightfield, first try the same exposure time that was used in collecting the optical sections during the axial scan. Adjust the exposure time to ensure the conditions described in Figure C and Figure D. When using fluorescence, since the drop of dye will likely fluoresce at a different intensity than the biological sample, it is especially important to carefully adjust the exposure time to ensure the conditions described in Figure C and Figure D. To emphasize, it is important that the maximum gray value in the image is at about 75 - 85% of the well capacity of the camera, and that there are no saturated pixels in the image.

Carefully record all exposure times and file names for the bias and flatfield images since you will need them later.

## **RS-170 Cameras**

The bias and flatfield frame collection procedure needs to be repeated every day, and for every dataset collection session. This process is used to compensate for potential misalignments in the optics, which will drift from day to day, and to compensate for dust that may settle in the optical train, which will change from day to day. Collect one bias frame and one flat field frame.

The bias frame is taken with light to the camera blocked. This may be accomplished by shifting the dichroic mirror filter set, by turning off the trans-illumination lamp (if transmitted light brightfield is used) or by shuttering the excitation lamp (if fluorescence is used). If you are using frame averaging, average as many frames as possible. As a rule of thumb, average 255 frames.

Collect a flat-field frame. This will be used to calculate the flat-field non-uniformity of your camera. A flat field can be provided in a number of ways. When using transmitted- light brightfield, the simplest and most reliable way is to remove the sample from the microscope and grab an image of the blank stage. Another way, which will take more care is to take the sample extremely out of focus, to the point where, in effect, the camera image shows a flat field (see Table 1). We recommend doing this only if it is not possible to remove the sample without disrupting your experiment setup. Otherwise, one should avoid this method, because it is more prone to errors and misjudgment. For example, Table 1: Figure (a) shows a common mistake, which is quite often made. This occurs when the sample is brought well out of focus, but an obscure out-of-focus remnant of the sample is still present. Avoid this situation. It is extremely important to have a flatfield image that represents a totally flat field as shown in Table 1 (b). When using fluorescence, the most reliable way to prepare a flat-field sample is to place a drop of dye onto a slide and to seal it



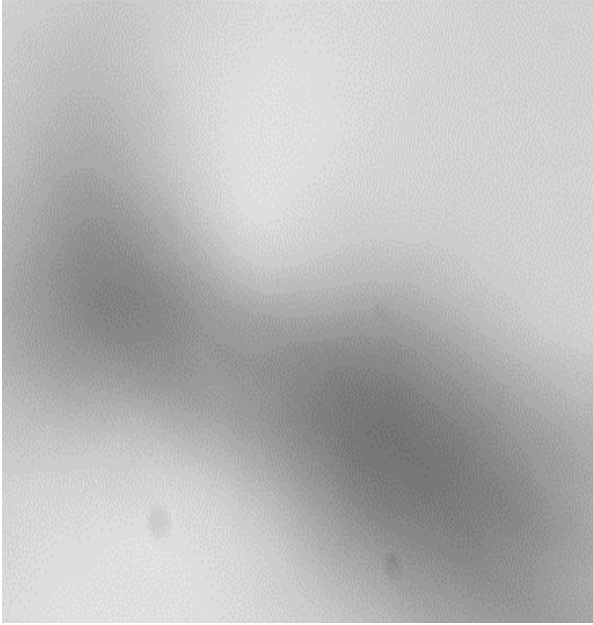
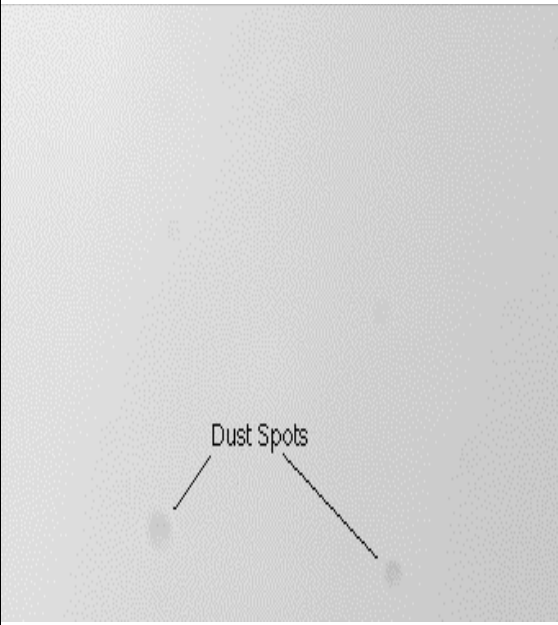
with a coverslip. Although, in principle, it may be possible to take the lens out of focus with the sample in place, for a flat field, we do not recommend doing so with fluorescence. Doing so is prone to problems. With fluorescence, it is much more difficult and prone to errors, to judge that the field is completely flat. Also, it is difficult (and often impossible) to collect an out-of-focus (flat-field) picture that has sufficient signal level, because the signal level decreases as the sample is taken out of focus. This signal-level problem may be compensated for, by increasing your video gain or by increasing your exposure time, doing so is prone to errors and is best avoided.

To summarize: For a flat-field brightfield image, remove the sample. For a fluorescence flat-field image, replace the sample with a slide containing a drop of the fluorescent dye, or try to defocus so that the current slide makes a good flat-field target (Table 1). However, be especially careful that there is no out-of-focus remnant of the sample and make certain that (for fluorescence) your exposure times and gains are set so that you have enough signal level in the flat-field image.

In selecting the camera gain (or gain on the frame grabber, illumination intensity or neutral density filter), refer to the discussion given earlier under **Setting the Exposure, Gain, and Offset**. When using transmitted-light brightfield, first try the same gain that was used for collecting the optical sections during the axial scan. Adjust the gain to ensure the condition described earlier in Figure D. When using fluorescence, since the drop of dye will likely fluoresce at a much different intensity than the biological sample, it is especially important to carefully adjust the gain to ensure the condition described in Figure D. To emphasize, it is important that the maximum gray value in the image is at about 75 - 85% of the well capacity of the camera, and that there are no saturated pixels in the image as shown in Figure C.

Make a careful record of all gains and file names for the bias and flatfield images in your laboratory notebook. Record all exposure times and numbers of frames averaged for all datasets collected, including the flatfield frames, bias frames and the optical sections of the sample.

**Table 1:**

	
Figure (a)	Figure (b)

**Table 1: Figure (a):** An incorrect attempt at a flatfield image by taking the sample well out of focus. This image is not far enough out of focus. An out-of-focus remnant of the sample is still visible.

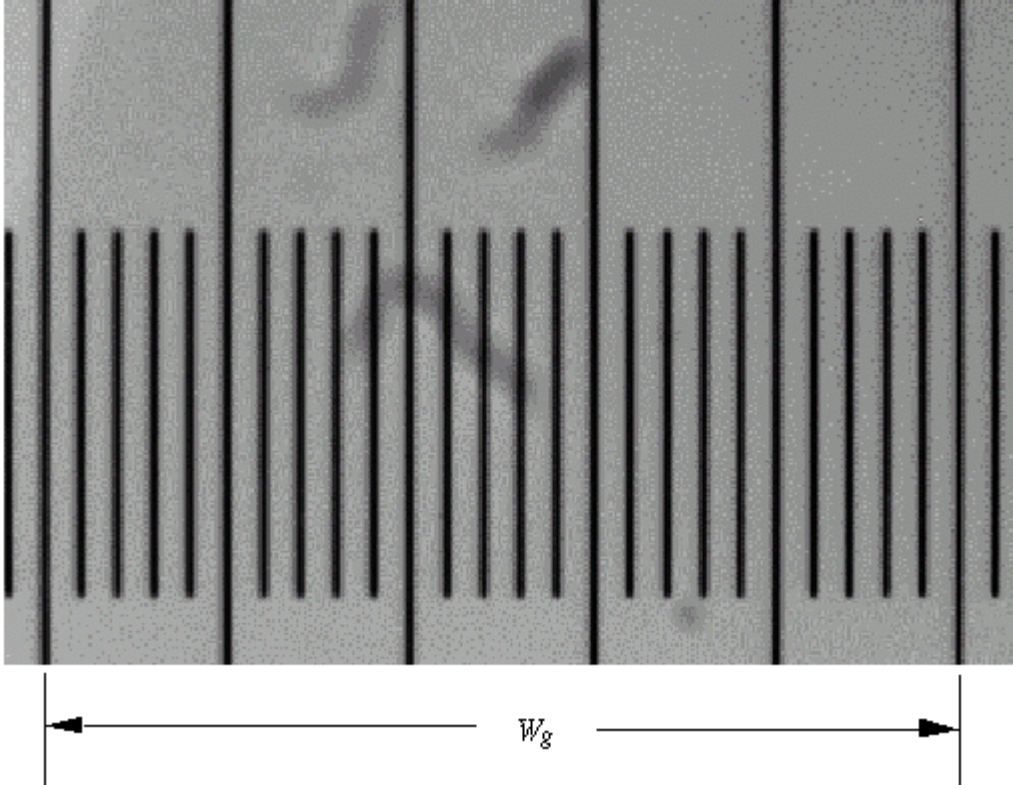
**Table 1: Figure (b)** A correct attempt at a flatfield image by taking the sample well out of focus. Note that no out-of-focus remnant of the sample is apparent. This image will be used to correct the optically sectioned dataset for non-uniform pixel sensitivity and for the dust spots that are highlighted.

## Spatial Calibration

Important parameters that are entered into the *AutoQuant* program in the Data Manager (see page 49 for more information on how to enter these parameters) are the x and y pixel sizes (distance between pixel centers). This may be obtained by using a microscope stage micrometer (e.g. Edmund Scientific product number J30,593 or J30,088). Collect two images of the stage micrometer, using the same lens with which you collected the optical sections. You will need to perform two calibrations - one for the x dimension and one for the y dimension.

Be sure that the graduations of the stage micrometer are properly aligned with the x and y axis. This is done by repetitively adjusting the angle of the camera position on the optical tube, and grabbing an image, until it is clear that the graduation lines are perfectly vertical or horizontal (see Table 2) for the x and y axis calibration, respectively. The stage micrometer will have a specification of micrometers per graduation. Denote this specification as  $\Delta x_g$  micrometers/graduation.

**Table 2:**

	
$N_g = 25 \text{graduations}$	
$N_x = 512 \text{pixels}$	
$\Delta x_p = \frac{10 \times 25 \times 12.7}{11.5 \times 512} = 0.54 \mu\text{m}/\text{pixels}$	
<p>Table 2: Stage micrometer image used for calibrating the pixel size. The distance between the first and the last of the six lines on the upper half of this image was measured to be 11.5cm (<math>W_g</math>) on a video screen. The stage micrometer dimension was known from the manufacturer to be 10<math>\mu\text{m}</math>/graduation. The width of the entire screen (<math>W_s</math>) was measured to be 12.7cm on the video screen. The resulting calculations are illustrated above.</p>	

**AutoQuant** or **5D Viewer** can be used to measure the x-y locations of pixels on a screen using a ROI box, the width of the box will be displayed on the Status bar of the main window. If the number of pixels between graduations along the x and y directions is  $n_x$  and  $n_y$ , respectively, then the number of micrometers per pixel along the x and y directions, denoted  $\Delta x_p$  and  $\Delta y_p$ , respectively, is given by:

$$\Delta x_p(\text{micrometers/pixel}) = \frac{\Delta x_g(\text{micrometers/graduation})}{n_x(\text{pixels/graduation})}$$

$$\Delta y_p(\text{micrometers/pixel}) = \frac{\Delta y_g(\text{micrometers/graduation})}{n_y(\text{pixels/graduation})}$$

If you do not have a stage micrometer available then you can use the following method to approximate the pixel spacing. From the manual for your CCD camera determine the size of the each pixel (e.g., 6 -10 microns is typical). Divide this number by the lens magnification and the camera zoom you are using. This will approximate the pixel spacing in your image. For example, a 40x lens with a 1.5x camera zoom and 6 micron CCD pixels will give you a pixel spacing for the image of 0.1um/pixel. The equation is camera pixel size / (lens magnification \* camera zoom).  
 $6\mu\text{m}/(40x * 1.5x) = 0.1\mu\text{m/pixel}$

## Vibration Control

To maximize the quality of the dataset acquisition, it is recommended that the imaging microscope system be isolated from vibrational disturbances to the extent possible.

The best approach to vibration control is to use a professional anti-vibration table (“air table”), (e.g. see the Ernest Fullam Inc. catalog, Latham, NY, or the Newport Catalog, Irvine, CA). Often, sufficient vibration control can be achieved for video microscopy systems without these air tables. The following section offers some insight into the vibration isolation.

Most vibration-related dataset collection problems arise from *relative motion* between microscope components. This type of relative motion can often be caused by external energy sources.

Mounting the instrument on a maximally rigid platform can minimize relative motion between microscope components. Remember that two types of rigidity are relevant - dynamic and static. Ideally, the platform should be as rigid as possible, dynamically as well as statically, and yet be as light as possible, and respond minimally to effects such as temperature changes. For example, Newport manufactures a special honeycomb sandwich structure that has these desirable properties. In most quiet, climate-controlled laboratories, an aluminum platform with a minimum thick-

ness of 0.5 - 1 inch may suffice. Thicker platforms have higher static rigidity. Try to minimize the length and width of the structure to the minimum necessary amount.

In addition to placing the microscope on a rigid platform, it is desirable to *isolate* the platform from external vibrations. For this, it is useful to be aware of the sources of vibrational energy around the microscope.

Also, vibrations (usually from cooling fans) can be transmitted to the microscope via unavoidable accessories, such as electrical cables. While most of the vibrational disturbances are vertical, it is possible to have disturbances that are along the horizontal direction.

Finally, one must consider shock control in addition to vibration control. Shocks often occur in a laboratory environment due to heavy footsteps nearby, when buildings are being repaired, or when heavy equipment is being moved.

Keep in mind that vibration isolation equipment must be considered not only for the microscope system itself, but also for laboratory vibration *sources* such as computer and electronic systems containing fans and relays. It is important to note that vibrations are best controlled at the source, if at all possible.

### ***Practical and Inexpensive Solutions***

Inexpensive vibration isolator components are available, based on a variety of principles, ranging from pneumatic through those based on the use of viscoelastic polymers, (see Edmund Scientific [Barrington NJ]; catalog no. A35, 264 Sorbothane Vibration Mounts). Such vibration mounts may be simply placed under a heavy pallet made of wood, ceramic, Plexiglas, slate, aluminum, or some other convenient (preferably heavy) material.

Many laboratories have mounted tennis balls on such pallets as vibration isolators. We recommend first deflating the tennis balls by drilling a small hole on one end. Otherwise, we have experienced that inflated tennis balls are too rigid and do not sufficiently dampen building vibrations.

## **Collecting Confocal Datasets**

When collecting confocal datasets that will be used in conjunction with deblurring the following rules-of-thumb apply. We emphasize that these are recommended rules-of-thumb for optimal operation. Owing to the difficult trade-offs among experimental conditions of signal-to-noise, photobleaching, photo-damage, field size, and many other considerations, you may find it necessary to bend these rules. These are *recommended* conditions and not *hard* rules. You should go ahead and bend them, if necessary. Because of the robustness of ***AutoQuant*** your deblurred results may still turn out fine, but this will depend on many factors, including your sample type, and is less predictable than if you follow the rules. To emphasize, keep in mind the following adage *garbage-in = garbage-out*. ***AutoQuant*** is designed to be robust against breakage of these guidelines. Breakage of say one or two of them may cause no problem. However, with each additional guideline that is broken, your chances of achieving satisfactory experimental results

decreases. As a general rule, do not break any of the guidelines for sake of convenience. Break only those guidelines for which your experimental design restrictions absolutely prevent you from following these guidelines. In other words, only break a guideline if you have no choice but to do so.

## Sampling Issues

As a rule of thumb, the in-plane sampling ought to be equal to the *resolution element* (resel) size, or spot size, of the image. According to Webb et al:

$$\Delta d_{xy} = 1\text{resel} = \frac{0.4\lambda}{NA}$$

and

$$\Delta d_z = \frac{1.4\lambda\eta}{NA^2}$$

where  $\Delta d_{xy}$  and  $\Delta d_z$  are the spot sizes in micrometers, for the in plane (XY) and axial (z) dimensions, respectively, and where  $\lambda$  and NA are the wavelength and numerical aperture, respectively. The least critical deviation is to make  $\Delta d_{xy}$  larger than that indicated by the above equation. This is because most of the noticeable smearing in images is along the axial dimension, which is not affected by making  $\Delta d_{xy}$  larger. The next least critical deviation from this rule is to make  $\Delta d_z$  smaller than indicated by the above equation. In fact, there are distinct advantages in doing so. Generally speaking, the finer the sampling is along z, the better is the deblurring along z. The above rule is considered the optimal trade-off between this axial deblurring and the potential photobleaching and other problems (such as extra disk space required) that may be caused by having finer axial sampling. The next least critical deviation is to make  $\Delta d_z$  larger than indicated by the above equation. Doing so lessens the amount of deblurring that is possible along the axial dimension, but there should still be substantial deblurring that is noticed. For example, it is common to process datasets that have  $\Delta d_z$  equal to 2 or 4 times that indicated by the above equation, and substantial deblurring and noise reduction is still usually obtained, albeit generally to a lesser degree. Making  $\Delta d_{xy}$  smaller than as indicated by the above Equation is the most critical deviation from the rule. Generally, it is best to avoid bending this rule, even though it has mixed advantages and disadvantages. Making the in plane sampling finer than the Rayleigh resolution limit will improve the in-plane deblurring. For confocal datasets, however, this desired improvement may come at the cost of degrading the axial resolution, depending upon how well the other rules-of-thumb are followed as explained below. In principle, so long as the confocal pinhole aperture is stopped down to a point that the microscope is considered to be *fully confocal*, then there should not be any problem with making this in-plane sampling finer. In fact, doing so will ordinarily improve the in plane deblurring as explained. On the other hand, it is very often the case (arguably most of the time) that the confocalist *does not* have his/her pinhole aperture stopped down to the fully confocal position (because he/she usually wants to detect more light to improve sensitivity).

Therefore, as a general rule of thumb for optimal results we do not recommend making the in-plane resolution any finer than as indicated in the above equation. If you do so, we recommend that special care be taken to be sure that your confocal microscope aperture is stopped down so that it is in the fully confocal position.

See the next two sections for recommended sampling when the aperture setting is other than fully confocal.

## Confocal Aperture

*AutoQuant* runs on the mathematical assumption that you have a fully confocal microscope. This condition is true so long as the confocal pinhole photodetector aperture is closed down to a size that is less than or equal to the “waist” of the scanning spot (diameter of the Airy disk). In the specimen plane, for a 1.4 NA lens, this spot size is around 0.25 micrometers and the recommended confocal aperture diameter (after projection onto the specimen plane) is then 0.25 micrometers or less. This number, however, does not represent the actual physical diameter of the aperture. For example: The Molecular Dynamics Sarastro<sup>TM</sup> confocal scanning system, for a 60X 1.4 NA lens, a spot size of 0.25 micrometers in the specimen plane is magnified to approximately 50 micrometers in the detector plane. Thus the recommended true aperture size is 50 micrometers for this particular setup.

Please check the specifications of your confocal microscope to determine the aperture settings needed to ensure fully confocal behavior. This condition will differ among confocal manufacturers.

It is well understood that often times this aperture size requirement may not be met. Because of signal-to-noise, photobleaching and other considerations, the confocal aperture is often opened wider. The main point to emphasize is that you should not do so casually, and only if it is necessary. The dataset may still deblur fine, but this will depend upon your sample and other experimental conditions and is not extremely predictable. Remember that the *AutoQuant* system is very robust against noise. It is generally more robust against noise than it is against improper confocal settings, so if you have a choice between fully confocal behavior and noise, for deblurring purposes it is better to sacrifice signal level (noise) for a proper confocal behavior. Remember that *AutoQuant* “thinks” that you have a fully confocal microscope, so it will try to reconstruct a fully confocal PSF. As you open the pinhole aperture, your microscope behavior begins to lean towards widefield behavior and the true PSF begins to have widefield character, such as a flaring, hourglass like shape. This behavior, in a sense, “confuses” the *AutoQuant* software and it will still attempt to reconstruct a fully confocal PSF, so you should avoid this condition, if possible. If this condition cannot be avoided, *AutoQuant* still has the robustness that it may provide a nice deblurred result, but this will be dependent on your sample and other experimental setups and is less predictable than with fully confocal settings. Consider the signal-to-noise requirements explained below.

## Less Than Fully Confocal Conditions

If you find it unavoidable to have the aperture setting opened wider than the fully confocal position, then we recommend likewise broadening your XY sample spacing by the same amount. For instance, if your confocal aperture is broadened to twice the diameter of its fully confocal position, then you should broaden your XY sample spacing (microns per pixel) to twice the above mentioned “optimal” sample spacing (~0.5 micrometers for a 1.2 NA lens or higher). Of course, other experimental conditions may prevent you from doing this, and this may be alright, but as with other rules of thumb, this is the so-called recommended optimal setting.

## Signal-To-Noise Considerations

*AutoQuant* will operate with a wide range of signal-to-noise levels. The Expert Settings have a Noise Smoothing feature to handle different signal-to-noise levels. Widefield datasets generally contain low levels of noise, therefore the default setting is Low. Confocal datasets generally contain medium levels of noise, therefore the default setting is Medium.

For **Signal to Noise Ratio (SNR)** levels of 10 or below, a Noise Smoothing setting of High should be selected. This will cause *AutoQuant* to be extremely robust against the noise. For SNR levels between 10 and 100 the Noise Smoothing setting of Medium should be selected. For SNR levels of 100 or higher, the Noise Smoothing setting of Low should work well. See Table 3 to identify noise levels approximately by visual comparison.

**Table 3:**

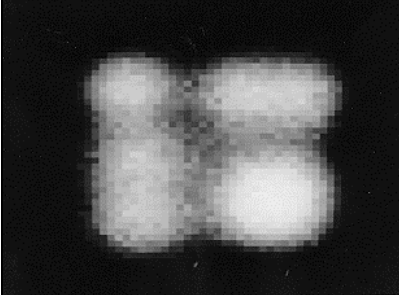
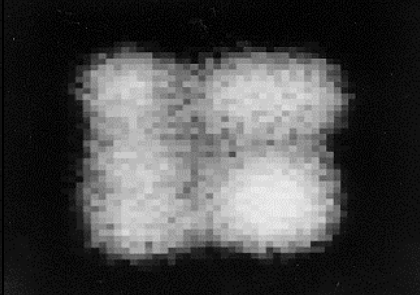
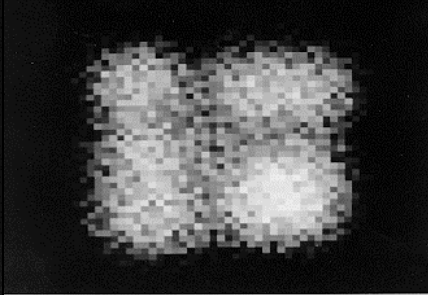
		
SNR 160:1	SNR: 40:1	SNR 10:1

Table 3 (Left to Right). Simulated SNR of 160:1, 40:1 and 10:1 respectively.



## 2-Photon Dataset Collection

### Guide for Obtaining the Correct Collection Parameters

#### ***Sampling Criterion for X, Y***

Sample at 1/2 the Airy disk width with a wavelength that is the excitation wavelength. The excitation optics determine the width of the PSF. This will be approximately  $(0.61\lambda)/(NA \times 2)$ . Where  $\lambda$  is the emissive wavelength and NA is the numerical aperture. Generally the result is 163 nanometers or finer (assuming a 1.4 NA lens). If you can sample finer (say 80 nanometers or 40 nanometers) try this also, as an experiment. It is expected that you may improve the x, y resolution by at least 2 and possibly by 4 (80 nanometers). To improve the “optical” resolution by 4 (to 80 nanometers) you need to increase the spatial sampling by 8 (40 nanometers).

#### ***Sampling Criterion for Z***

Sample at 1/2 of the PSF axial width, which approximately follows the formula:  $(1.4)\lambda\eta/(NA^2)$ . Where  $\eta$  is the refractive index of the lens and NA is the numerical aperture. This is the formula for the axial width of a confocal PSF. For 2-photon microscopy, use the excitation wavelength in this formula. Assuming a 1.4 NA lens, this distance is approximately 800 nanometers. For more substantial deblurring along z, experiment with even finer sampling. For example, 400 or 200 nanometers or finer.

Keep in mind that the above rules are only rules of thumb for ensuring that deconvolution will show substantial deblurring. If, on the other hand, the experimental objectives do not call for an improved x, y resolution but they do call for an improved z resolution, then follow the *Sampling Criterion for Z* and sample as you otherwise normally would do along x, y. If the experimental objectives do not call for an improved z resolution but they do call for an improved x, y resolution, then follow the *Sampling Criterion for X,Y* and sample as you otherwise normally would do along z.

The modality setting of ***AutoQuant***, which is found in the Data Manager (see page 49 for more information on using the Data Manager), both on the Summary tab and the Instrument tab. This is because the optical PSF of the typical 2-photon microscope, theoretically (assuming the detector aperture is wide open), is equivalent to that of a confocal PSF, except that it is determined by the excitation wavelength rather than the emission wavelength.

# Introduction

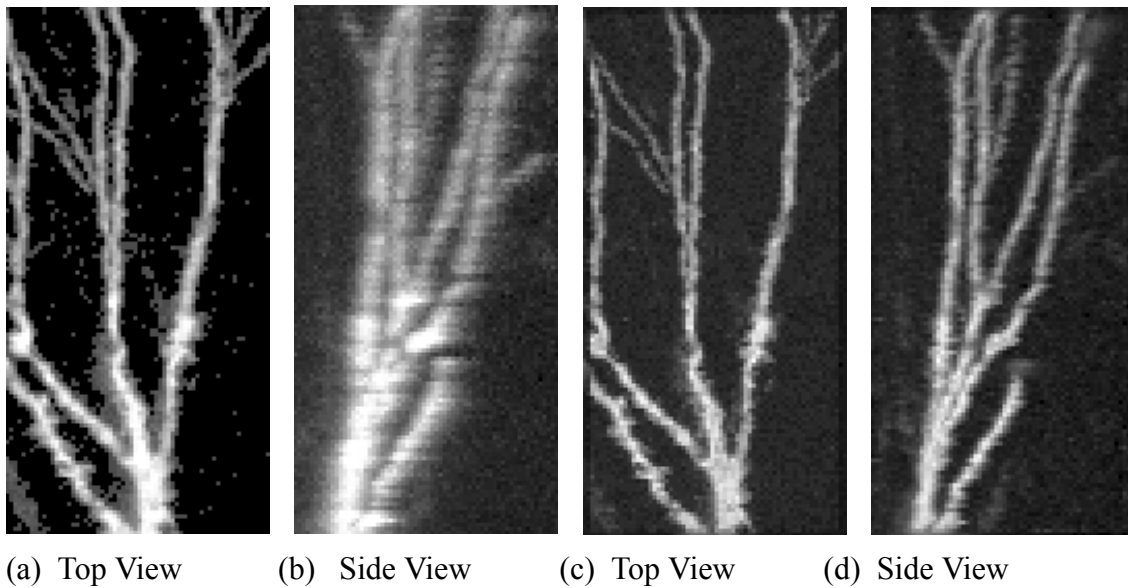
## What is 3D Deconvolution

Also known as “3D image restoration”, deconvolution is a computational technique for removing out-of-focus haze from stacks of optical sections. The out-of-focus haze can be mathematically modeled as a “point spread function (PSF)”. Deconvolution methods can therefore be thought of as methods for inverting the unavoidable and natural blurring effect of the point spread function (PSF).

For example, suppose we focus a microscope having a fine depth of field (DOF) of about 0.4 micrometers, onto a plane in a microscope sample which may be anywhere from a few micrometers (e.g. chromosomes or epithelial cells) to on the order of 50 to 200 micrometers (e.g. neurons or brain slices). The acquired image will contain both sharp, in focus features originating from the plane of focus, as well as hazy, blurry features originating from out-of-focus planes that are above and below the plane-of-focus. After capturing this image electronically, we can refocus the microscope to an adjacent plane that is on the order of a depth of field (DOF) away, and acquire a second image. This process can be repeated until the entire specimen has been scanned. The resulting dataset will be a “3D dataset”, or in other words, a “3D image”. However, it will be a rather poor image, because it will contain all of the out-of-focus haze and blur.

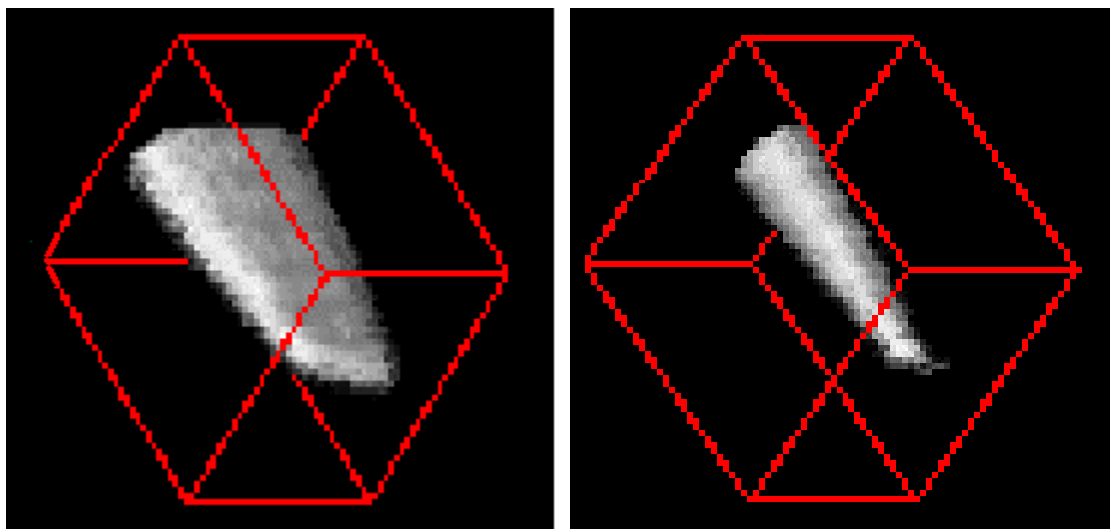
The purpose of deconvolution is to remove the haze and blur and to restore sharpness and clarity to the image, as illustrated for a neuron image below.

The effects of deconvolution are usually most pronounced in the axial direction (z-axis), as illustrated for a confocal image of a rat neuron in the image that follows.

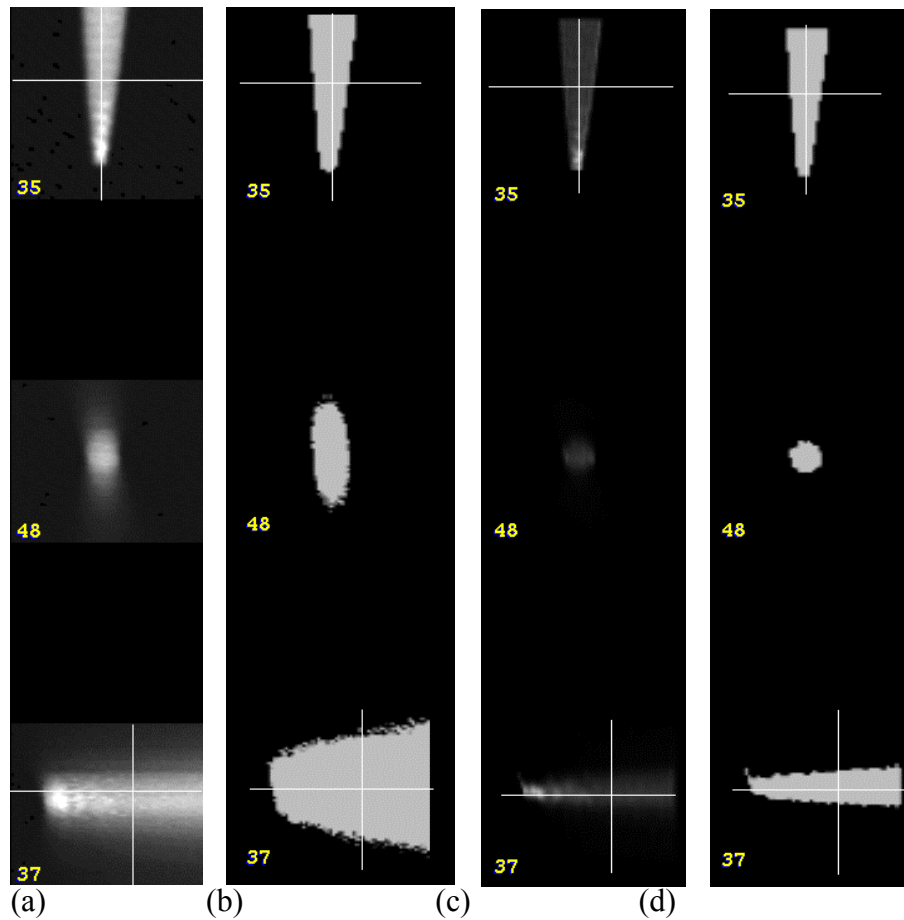


**Figure 2:** Confocal image of a CA3 hippocampal rat neuron. Panels a and b show the projections of the dataset from the top and side respectively. Panels c and d show the same dataset after deconvolution.

The improved axial resolution becomes especially important when accurate morphometry is desired.



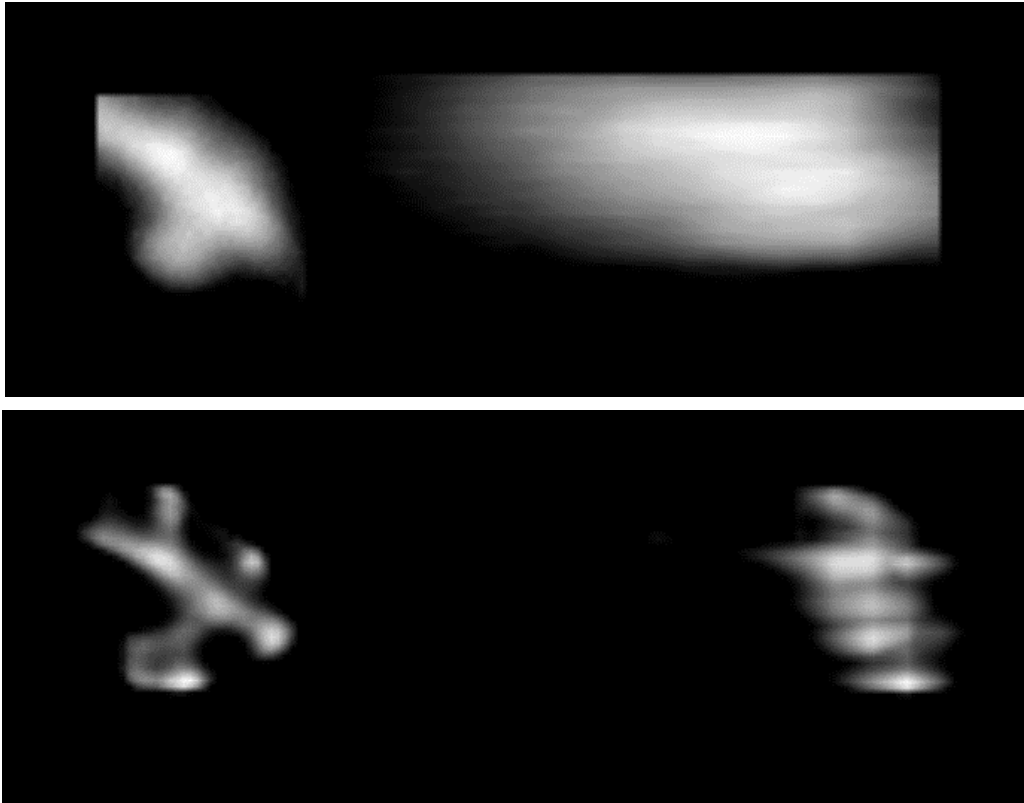
**Figure 3:** On the left is a rendering of the 3D image of a fluorescent dye-filled micropipette imaged by a laser-scanning confocal microscope. On the right is the same object after deblurring. The axis running from left to right corresponds to the microscope axial direction.



**Figure 4: Showing the improved morphometry made possible by deblurring for the image of a known object (fluorescent dye-filled micropipette). (a) Cross-sections through the raw undeblurred dataset (the sectioning positions are indicated by the white lines). (b) Cross-sections through the segmentation of the dataset at the same slices. Panels (c) and (d) show cross-sections through the deblurred dataset. Note the symmetry of the latter result, which agrees well with known reality.**

The improved axial resolution due to deblurring is illustrated in Figures 3 and 4 for the case of a known object (a dye-filled micropipette).

Deconvolution can often reveal fine details that are not directly observable. Figure 5 shows an example of this fact.



**Figure 5:** Example of improved detail revealed by deconvolution. The upper panel shows the top and side projectional views of a portion of a neuron. The lower panel shows the same views after deconvolution. Notice how the spines of the neuron become observable.

## Application of the AutoQuant Deconvolution Package

The *AutoQuant* software package can be used to perform 3D deconvolution in the context of the following forms of microscopy:

1. Transmitted-Light Brightfield Microscopy (TLB);
2. Widefield Epi-Fluorescence Microscopy (WF);
3. Confocal Pin-Hole Laser-Scanned Epi-Fluorescence Microscopy (CLSM);
4. Spinning Disk Scanning Confocal; and
5. Two-Photon Fluorescence.

For the confocal fluorescence microscope, images of adjacent planes are sequentially digitized and optically sectioned in a manner that is generally similar to the widefield case, except that each frame is collected by way of a raster-scanned laser spot and a photomultiplier pinhole detector. Each confocal optical section is, by itself, already relatively deblurred in the sense that the confo-

cal optics rejects most of the out-of-focus light. However, in spite of this improvement over the widefield microscope, the confocal microscope has its own limitations.

- (1) While it indeed rejects *most* of the out-of-focus light, it by no means rejects *all* of it, and thereby retains an obvious haze, even though this haze is much reduced from that of the widefield microscope.
- (2) Another effect of this imperfect out-of-focus light rejection is that the image retains substantial axial smearing. For an image which is not diffraction limited, this smearing appears akin to a linear motion blur in the axial direction. For an image which is diffraction limited, as with empty magnification, this smearing appears as a substantial nonisotropic blur (or nonisotropic spatial resolution), with most of the smearing along the axial direction.
- (3) Owing to this light rejection, far fewer photons are detected, so a rather substantial quantum-photon noise component may be seen in the raw dataset. This noise component routinely causes limitations in protocols where very fine structures need to be seen. For instance, it limits our ability to detect a void in the fluorescence concentration that is often useful to pinpoint impalement sites of electrophysiology probes [Turner, Szarowski et al., 1991], and it limits our ability to determine if a dendritic spine is present. It is difficult to judge whether a void in the image intensity is due to a fine structure that is really there or if it is due to erroneous, random fluctuations caused by the statistical nature of quantum noise.

In the case of confocal fluorescence microscopy, it is the purpose of deblurring algorithms to reduce these three undesirable effects and to thereby improve the utility of the microscope.

Deblurring is best carried out when the dataset is seemingly oversampled. In other words, best results are achieved when the pixel or optical section spacing is finer than what would be used normally, without deblurring. In-plane deblurring is most successful when the sample spacing is finer than 1/2 the Rayleigh width. If in-plane deblurring is desired, we will sample at 0.1  $\mu\text{m}$ , or finer, when the 1/2 Rayleigh width is  $\sim 0.25 \mu\text{m}$ . Axial deblurring is substantial with widefield optics regardless of the  $z$  sampling. The rule of thumb for confocal datasets, without deblurring, is to have the optical sections spaced according to the confocal spot size along  $z$ , which is the resolution element along the  $z$ -direction calculated by the following equation [Webb et al, 1995]:

$$\Delta z = \frac{(1.4\lambda\eta)}{NA^2}$$

where  $\lambda$  is the wavelength in microns ( $\mu\text{m}$ ),  $\eta$  is the refractive index of the immersion medium, and  $NA$  is the numerical aperture of the objective lens. If  $\Delta z$  is 0.5  $\mu\text{m}$ , we will sample at 0.5  $\mu\text{m}$  or finer per slice. The noise suppression mentioned above requires this type of fine sampling.

## What is Blind Deconvolution?

Blind deconvolution is a term that is used to describe methods of deconvolution, which do not require the point spread function (PSF) of the system to be *explicitly known prior* to the deconvolution. **AutoQuant** is based on blind deconvolution. Indeed, a reconstructed estimate of the PSF is produced by **AutoQuant** concurrently with the deconvolved image dataset. There is no need to provide **AutoQuant** with a PSF of any kind.

The important practical implication of blind deconvolution is that it adapts to the real PSF of the microscope system, which can be significantly different from the theoretical PSF (see Figure 6), and from previously measured PSFs due to specimen and instrument variations. The **AutoQuant** blind deconvolution system is able to adapt to PSF changes within a specimen itself. Thus, the deconvolved results are superior to those of methods that utilize theoretical or previously measured PSFs.

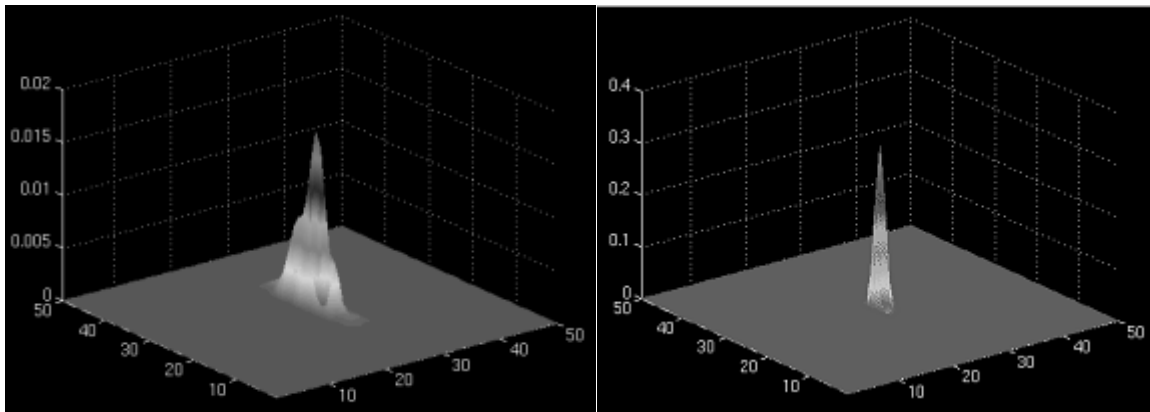


Figure 6: Shown on the left is a theoretical PSF for a confocal microscope. On the right is the PSF as revealed by **AutoQuant**.

Direct, experimental measurement of the PSF is an arcane and difficult procedure that is not straightforward [Hiraoka, Sedat et al., 1990] for routine usage and thereby impedes the wide usage of deblurring algorithms that require it. The PSF measurement method typically involves imaging a sub-resolution fluorescent microsphere. This approach has several limitations. First, noise in the imaging system also shows up in the measured PSF. This can be overcome to some extent, by averaging over several microspheres and/or frames. This is time-consuming and effort-intensive. Second, photobleaching of the microspheres limits the strength of the obtainable signal. Third, the actual PSF changes when the microsphere sample is removed and the biological specimen is inserted. Fourth, the PSF measurement must be done for each sample collection session, and for every optical configuration used. This is because the PSF is a function of the refractive index of the sample or medium, and its nonhomogeneity [Gibson and Lanni, 1991]. Finally, the PSF may vary spatially, especially along the optic axis.

All of the above complications are compounded with confocal microscopy, where light levels are especially low and where the dependency of the PSF to the sample refractive index is especially strong [Visser, Oud et al., 1992].

By using blind deconvolution methods, *AutoQuant* eliminates the need for Point Spread Function (PSF) measurement, improving both the accuracy, as well as convenience of deconvolution. This is a major advantage of using this package instead of those utilizing non-blind deconvolution methods.

Another clear practical advantage of *AutoQuant* is that it inherently reduces noise in the image, especially in severely noisy cases. This is partly because its mathematical foundations have quantum photon noise as a fundamental assumption. Intuitively, this may be explained in that the microscope represents a band-limited system, which means that any good light signal will lie only inside the band limit. Much of the undesirable noise energy lies outside of the band limit, and the algorithm inherently recognizes this and automatically rejects the out-of-band noise, while properly constraining the deblurred image to have only nonnegative values. This advantage is especially important for confocal microscopy, where images often have a large noise component.

## Important Features of AutoQuant

**Resolution Improvement and Noise Reduction:** *AutoQuant* improves the resolution of the microscope in the x, y, and z dimensions. Additionally, datasets from the microscope system may be especially noisy, as is often the case with confocal datasets and with fluorescent widefield datasets using an intensified CCD camera. *AutoQuant* greatly reduces this noise.

**Statistical Optimality:** Because *AutoQuant* is based on Constrained Maximum Likelihood Estimation theory, it is, in this sense, statistically optimal. Intuitively, the algorithm searches for the most likely 3D dye concentration, which caused the collected image dataset.

**Correct Modeling Assumptions and Constraints:** The *AutoQuant* design is based on carefully justified mathematical modeling of the optical system, including Poisson statistical modeling of the noise component in the detector electronics. The Poisson model not only works well with low-photon-count datasets; it also properly constrains the probe concentration and PSF to be nonnegative. This is something that is not done by methods based on Gaussian image models (e.g., Least-squares, Jansson Van Cittert). The model in *AutoQuant* also takes into account the optical characteristic of the dataset collection system (including the sample) and ensures that the estimated probe concentration and PSF agree with the physical system.

**Advantage of an Iterative Method:** Because the method is iterative and nonlinear it is capable of properly constraining the solution (i.e., constraining the probe concentration and PSF solutions) to be nonnegative. This is especially important for both quantitation, and for resolution improvement (especially z resolution). Our algorithm is capable of optimally restoring the missing cone of frequencies that are inherent in the PSF for widefield microscopes. For a detailed description of the problem of *missing cone of frequencies*, the reader is referred to the paper by Streibl (see Bibliography). Linear methods, such as the well-known Nearest Neighbor method and the Inverse Filtering methods [see references] are not capable of implementing such constraints, and are thereby not nearly as suitable for quantitation or resolution improvement. However implementations of both of these simple and fast deconvolution methods are included in *AutoQuant*.



**Dark Current (Bias) and Flatfield Corrections:** The data correction utility is a component in *AutoQuant*. This provides features for automatic data correction, including automatic bias and flatfield correction. This component corrects data from video rate and cooled CCD cameras.

**Processing Many Datasets:** *AutoQuant* allows the sequential processing of many datasets by using its *Batch Processing* feature.

## New in Version X3

This version of AutoQuant contains many improvements, including:

- **New Window 7 User Interface:** An all-new high-resolution *Windows 7* look and feel with intuitive new icons for all available tools.
- **AutoQuant Connect:** *AutoQuant Connect* is a new feature that allows you to automatically import and export your data to partnering software applications for additional analysis or visualization. Using AutoQuant Connect, you can import a "CONNECT Image Set" acquired in partner software applications (such as *Image-Pro Plus* or other software partners). The set includes the acquired image set, the image set meta-data, complete deconvolution settings, instructions, and completion commands. Upon receiving the "CONNECT Image Set", AutoQuant will queue all subsequent sets into the batch processing viewer to launch the next task. Each set is deconvolved using the defined settings with AutoQuant's industry leading algorithms. Once the deconvolution task is complete the image set along with complete meta-data is returned to the location of your choice. It can be returned to the connected software partner application, saved to a custom folder, or displayed in AutoQuant for review, visualization, or analysis.
- **Gibson-Lanni PSF Modeling:** This proven Point Spread Function (PSF) modeling algorithm is now available for Thoretical PSF determination as well as Spherical Aberration (SA) detection and correction. This results in higher-quality PSF modeling.
- **Auto Detect unknown Sample Medium RI and distance from cover-slip:** An automatic calculation is run based on the known accurate parameters to derive both values when the information is not available.
- **Sync Multiple Line Profiles and Measurements for comparison:** Compare original image sets with deconvolved results easily by mapping a line profile to identical pixels on a second resultant image. Display the line distance in microns, compare the angle of the line to the horizontal, auto-scale the intensities, and view the data in a log scale if the values are drastically different. Export the comparison data in \*.csv format for viewing in *Excel*.
- **ROI Deconvolution Preview:** It can be difficult to test the settings of your deconvolution run on larger image sets quickly. This new feature, available for both the 2D and 3D deconvolution algorithms, allows you to preview your results using all the active settings. You draw a small ROI on the image set, and then the optimized acceleration preview begins. Using the current settings in the deconvolution panel, this feature provides an instant look into the results awaiting at the end of the full-stack deconvolution.
- **Batch Viewer Error Reporting:** In the event that the system or application stops working, the Batch Viewer can now report the details of the error.
- **Save Deconvolution and Optical settings files in \*.cfg format:** Save all your active parameters in the 2D and 3D deconvolution dialogs to a file for future use on the same or a different system. You can also save the typical optics parameters that you use most

often. Enter the system parameters once and then simply recall the settings each time you load an image into AutoQuant.

- **Sample RI and Distance from Coverslip added to Data Manager:**

These two parameters in the Data Manager are now visible, so that you can apply them to the core of the deconvolution algorithm. This improves accuracy which leads to better results.

- **Faster Deconvolution Times:** This version of AutoQuant uses optimized math libraries, fewer temp files, and improvements to various algorithms which should result in faster processing times for certain systems.

- **Faster loading of images into the batch processing queue:** The batch queue has been optimized to reduce the loading time for batch processing. The larger the image set, the easier it is to queue up and begin the process.

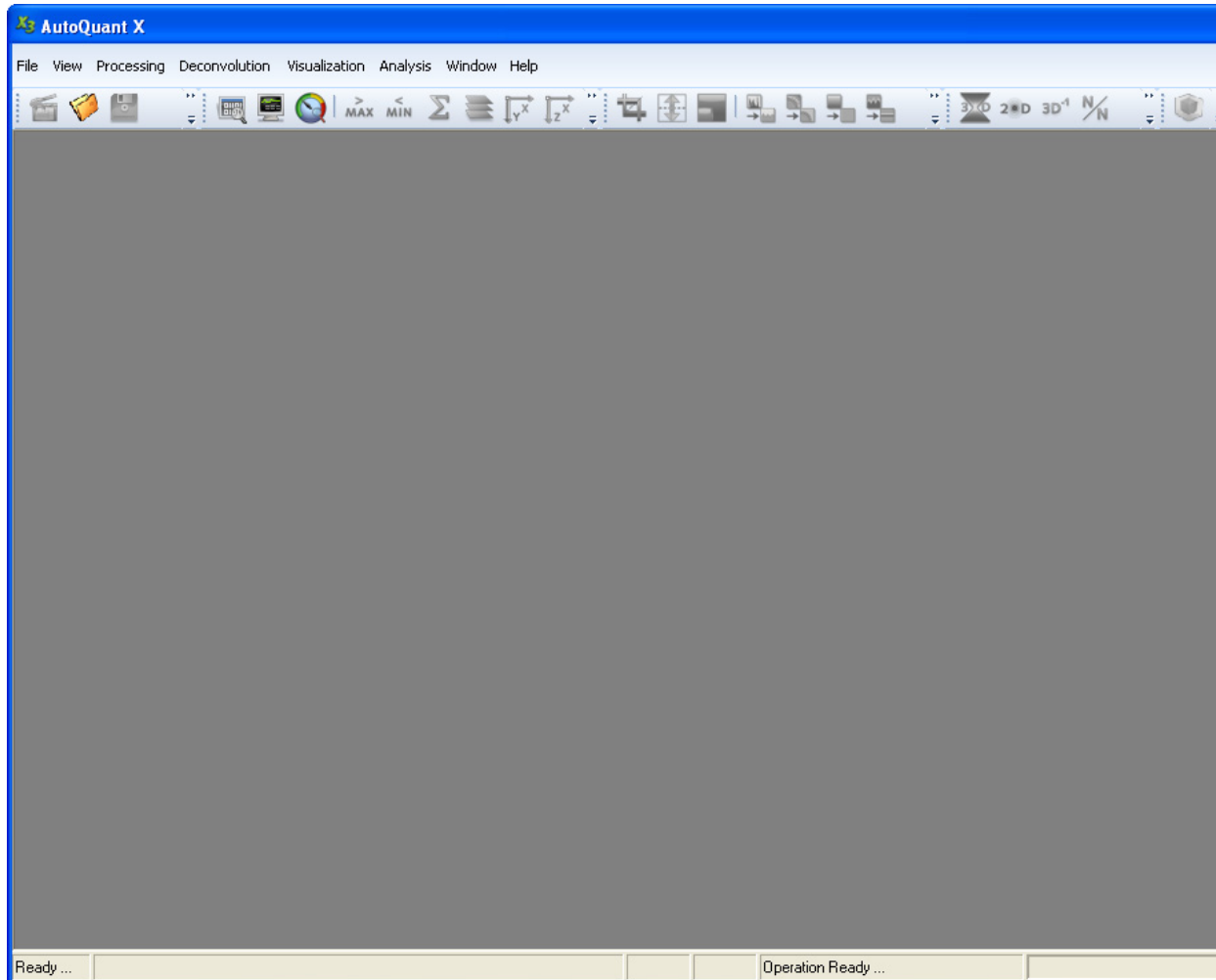
- **New Additions to the Objective, Camera, and Dye lists:** Many new entries have been added to update these lists with the latest hardware and dyes available in the market.

## Basic Principles Underlying the AutoQuant System

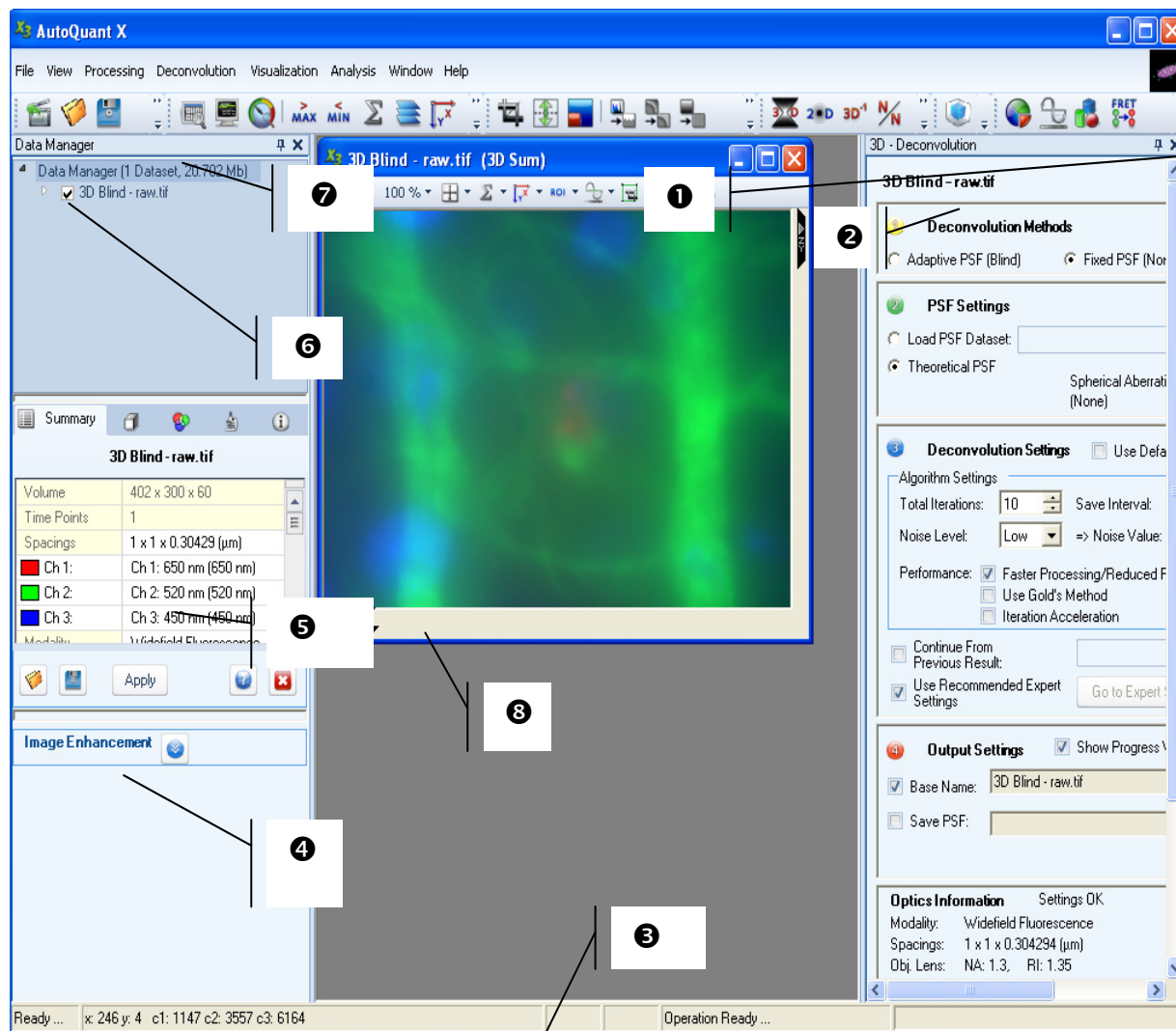
The basic mathematical and conceptual principles underlying the *AutoQuant* system have been widely published in scholarly journals and conference proceedings worldwide. Constrained Maximum Likelihood Estimation may be thought of in two ways. First, consider a 3D fluorescent sample. We may think of the true 3D probe concentration as an unknown random quantity. In an intuitive sense, the image reconstruction algorithm is searching for the 3D fluorescent probe concentration, which is “the most likely one” to have caused the raw optically sectioned image dataset, taking advantage of any available constraints. In a mathematical sense, the algorithm is based on the formulation of a mathematical function, called the likelihood function. This likelihood function depends on both the unknown 3D fluorescent probe concentration and the 3D PSF. The algorithm iteratively solves for the 3D image that maximizes this likelihood function. In other words, the likelihood function is solved for the correct probe concentration and PSF that provide this maximum value solution. This solution then, in effect, becomes our reconstructed 3D probe concentration and PSF. This optimization model is based on an assumption that the noisy camera data follow a Poisson random signal model. This assumption is physically justified; assuming a quantum photon limited photo detector, which is certainly the case in many situations, especially for very noisy low light level situations. Thus, *AutoQuant* has a sound mathematical basis, which makes it statistically optimal. Partly due to this, the algorithm inherently reduces noise in the dataset. Additionally, it incorporates numerous improvements and valuable features.

## User Interface


This section will explain the user interface for the software, explaining what is in each section and how to interact with it. More detailed explanations can also be found in the Online Help that is included with the product. When you first start the application, the user interface will look something like this:




Once a dataset has been opened, it will appear as shown in the following page:




Starting at the top, and moving clockwise, the features indicated are: 1) Docking Controls, 2) Dialog Box, 3) Operation Status, 4) Image Enhancement Dialog, 5) Data Browser, 6) Data Properties, 7) Data Manager and 8) Perspective Tabs. Below, each item will be explained.

1) Docking Controls - There are two icons in the Docking Controls section. The tack  controls whether the dialogs display all the time, or whether they only display when they are clicked on. If the tack icon is vertical, as shown in the previous image, then the dialog remains displayed

all the time. Once the icon is clicked, the tack will appear horizontally , and the dialog will remain open only as long as the mouse is held over it. Once the mouse moves away from the dialog, the dialog will slide closed. It can then be reopened by placing the mouse over the tab for that dialog, located along the side of the application window.

The default for the tack icon is vertical (open at all times).

The other icon in the Docking Controls is the Close icon . Clicking this icon will close the dialog. The Data Browser dialog can be reopened by going to *View->Data Browser*. Depending on which dialog was open on the right side, the dialog can be reopened from its source menu (e.g. the 3D Deconvolution menu would be opened by going to *Deconvolution-3D Decon*).

2) Dialog Box - The Dialog Box appears on the right side of the application window, and is opened either via a menu item or a toolbar icon. If there are multiple dialogs open (e.g. 3D Decon, 5D Viewer, and Statistics), a tab will appear for each of the open dialogs. If they are docked (always displayed, with a vertical tack icon) the tabs will appear at the bottom of the dialog box. If they are not docked (not displayed all the time, with a horizontal tack icon) the tabs will appear along the right side of the application window. Clicking on these tabs will display the tab that has been clicked on.

3) Operation Status - The Operation Status displays the status of all open operations in the application window. Any operation can be right-clicked on and canceled. Right-clicking on a process also gives the option to show details (if it is a process that has sub-processes), as well as to clear all completed processes. This is helpful when the Operation Status becomes cluttered with various processes, many of which are completed.

4) Image Enhancement Dialog - The Image Enhancement dialog contains the controls to adjust the maximum and minimum intensities (by moving the red and orange triangle sliders at the bottom of the graph) and the gamma by either entering a value into the textbox labeled *Gamma* or by adjusting the green triangle slider at the bottom of the graph.

5) Data Browser - The left side of the application window contains the Data Browser, in which is contained the Data Manager, Data Properties Summary, and Image Enhancement Dialog. The Data Browser can be docked or undocked as described in the Docking Controls section above.

6) Data Properties - The Data Properties section contains the image information for the selected dataset. Included in this information are the image dimensions, image spacings, channel information, number of slices, modality information, as well as space to enter notes on the dataset.

7) Data Manager - The Data Manager section displays the opened datasets in a file tree format. With multiple datasets open, the active dataset can be selected by clicking on that dataset in the Data Manager (or by clicking on the dataset display in the application window). Additionally, the display of a dataset can be adjusted by selecting or deselecting channels, slices, and or time points to be displayed. This can be done by clicking on the box next to each channel, time point or z-slice. If there is a green check, that item will be displayed, if there is no green check, it will not be displayed.

Note: If you have multiple datasets for which you cannot import optics data, but the optics information is the same for all of those sets, you can load that information into AQX. Then set the optics for one of those sets, and copy/paste the optics information into the other datasets. You can do this in the data manager by right-clicking on the set that has the optics information, selecting “Copy Optics” and then selecting the remaining sets. Right click-on the sets where you would like the information to go, and select “Paste Optics”.

The toolbar along the top of the application window provides shortcuts to several commonly used functions. These will be shown below as they appear on the toolbar, from left to right.



File Open Icon - Displays the File Open dialog.



Save As Icon - Displays the File Save dialog.



Save Movie Icon - Opens the dialog to save the dataset as an \*AVI file.



Close Dataset Icon - Closes the current dataset.



Save Current View Icon - Saves the current active view..



Data Manager Icon - Opens the Data Manager dialog if it is not already opened.



Display Status Icon - Displays the Status window at the base of the application window if it is not already opened.



Batch Processing Icon - Opens the Batch Processing Dialog if it is not already opened.



Maximum Projection Icon - Displays the active dataset in the Maximum Projection view.



Minimum Projection Icon - Displays the active dataset in the Minimum Projection view.



Sum Projection Icon - Displays the active dataset in the Sum Projection view.



Slice View Icon - Displays the active dataset in the Slice View.



XY Icon - Displays the active dataset in the XY view.



XZ Icon - Displays the active dataset in the XZ view.



YZ Icon - Displays the active dataset in the YZ view.



XYZ Icon - Displays the active dataset in the XYZ view.



Copy Current View Icon - Copies the active view to a new workspace.



Cropping and Reshaping Icon - Opens the Cropping and Reshaping Dialog.



Extend Slices - Opens the Extend Slices Dialog.



Resize Image Icon - Opens the Resize Image Dialog.



Background Equalization Icon - Opens the Background Equalization Dialog.



Background Subtraction Icon - Opens the Background Subtraction Dialog.



Attenuation Correction - Opens the Attenuation Correction Dialog.



Optical Density Correction Icon - Opens the Optical Density Correction Dialog.



Slice to Slice Alignment Icon - Opens the Slice to Slice Alignment Dialog.



Channel to Channel Alignment Icon - Opens the Channel to Channel Alignment Dialog.



Image Algebra Icon - Opens the Image Algebra Dialog.



3D Deconvolution Icon - Opens the 3D Deconvolution Dialog.



2D Deconvolution Icon - Opens the 2D Deconvolution Dialog.



3D Inverse Filter Icon - Opens the 3D Inverse Filter Dialog.



No/Nearest Neighbor Icon - Opens the No/Nearest Neighbor Dialog.



5D Viewer Icon - Opens AutoQuant 5D Viewer and the Properties Form Dialog that controls the AutoQuant 5D Viewer display.



Image Statistics Icon - Opens the Image Statistics Dialog.



Line Profile Icon - Opens the Line Profile Dialog.



Count/Track Icon - Opens the Count/Track Dialog.



FRET Cross Talk Correction Icon - Opens the FRET Cross Talk Correction Dialog.



FRET Efficiency Calculation Icon - Opens the FRET Efficiency Calculation Dialog.



Colocalization Icon - Opens the Colocalization Dialog.



Radiometrics Icon - Opens the Radiometrics Analysis Dialog.





## Using the Interface

This section will go over the basic controls used to navigate and manipulate the User Interface.

### Using the Data Manager

The Data Manager is a simple way to manipulate the display of a dataset. When a dataset is opened, the dataset name will appear in the Data Manager as a file tree. Under the dataset name will be listed the channels, z slices, and or time points associated with that dataset. Deselecting a channel will change the image display of that dataset to not display that channel. A channel can be deselected by clicking the box next to the desired channel so that the check mark in that box disappears.

- (1) Navigate to the Multi-Channel folder in the tutorial data. Open the colorpollenc.tif dataset.
- (2) Notice that when the dataset is displayed in the application workspace, the filename also appears in the Data Manager.
- (3) Click the arrow next to colorpollenc.tif in the Data Manager. The channels will drop down in a file tree structure.
- (4) Uncheck the box  next to Channel 1 and notice that the red channel is no longer displayed in the dataset.
- (5) To remove a dataset from the Data Manager, highlight that dataset, then click on the waste basket icon .

### Using the Data Properties Dialog


The Data Properties dialog contains crucial information about the active dataset. The information is divided into five categories: Summary, Dimensions, Channels, Instrument and Info. The Summary tab displays all of the image information contained and editable in the other three tabs. The Dimensions tab contains the spacing and channel properties of the dataset. The Channels tab contains the controls for changing and naming the channels. The Instrument tab contains the information regarding the microscope used to acquire the dataset. The Info tab contains optional information such as the name of the specimen, when it was acquired and by whom, and notes regarding the dataset. Any of the fields that are white (in any of the tabs) can be edited either by clicking within the desired field and entering text or numbers into the field, or by selecting an item from the dropdown menu within the field.

## Using the Image Enhancement Tool

Depending on the display of the machine running the software, images may appear too dark, or in other cases, too bright. To compensate for these differences it is possible to perform a Gamma correction on a particular view.

(1) From the File menu, select *Open*. Navigate to the SmallHip folder, open the SmallHip.avz dataset and generate an XZ view.

(2) Towards the bottom of the Data Browser, it says *Image Enhancement*. To display the

Image Enhancement dialog, click the expand button , the Image Enhancement dialog will now be displayed.

(3) Enter a value of 1.2 in the Gamma field. Notice that the triangle located in the center of the histogram moves slightly, as well as a change in the image. The Gamma can also be adjusted by clicking on that triangle on the histogram, and dragging it to the left or right.

*Note: A Gamma value of 1.0 restores the image to its original state. Values greater than 1 brighten the image, whereas values below 1 darken the image. Gamma values may be between 0.1 and 10.0.*


(4) You may also adjust the darkness and brightness of the image in any of several ways:  
-Enter a new value in the Intensity text boxes. The Min Intensity text box on the left is for the minimum, and the Max Intensity text box on the right is for the maximum (the Image Enhancement dialog will default to the minimum and maximum values present in the image). The minimum must be less than the maximum.

-Enter a new value in the Percentage boxes between 0 and 100. The Min % text box on the left is for the minimum, and the Max % text box on the right is for the maximum. The minimum must be less than the maximum.

-Left click on the red triangle on the left side of the histogram, and drag it to move the minimum intensity indicator (red line) to the right. Left click on the orange triangle on the right side of the histogram and drag it to the left to move the maximum intensity indicator (orange line) on the histogram. The maximum intensity indicator can only be placed to the right of the minimum intensity indicator, and the minimum intensity indicator can only be placed to the left of the maximum intensity indicator. The Gamma can also be adjusted by left clicking on the green triangle on the bottom of the histogram and dragging it to the left or to the right.

The intensity values are used in a histogram stretching intensity filter. For 8 bit data, this filter sets all the pixel values that are greater than the maximum parameter to 255, and all the pixel values less than the minimum parameter to 0. Pixel values, which fall in between the brightest and darkest parameter values, are linearly scaled between 0 and 255.

Image Enhancement may be done on a single color channel (Red, Green, Blue) or on All Channels by activating the appropriate choice under Channel.

- (5) Close the Image Enhancement box by clicking the up arrow button .

*Note: Only in a Slice View will, the maximum and minimum values for Brightness and Darkness Level be the true maximum and minimum values of the entire dataset.*

In a time-series dataset, the changes can be applied to all time points by clicking the *Apply to all* button and conversely, they can be reset by clicking the *Reset all* button.

## Status Display





The Status Display feature will show the status of all current processes within the AutoQuant/5D Viewer platform. The display is at the bottom of the application window. To display the Status Window, select *Display Status* from the View menu. Within the Status Display, processes can be canceled by right-clicking on that process in the Status Display and selecting *Cancel*. Additionally, the Status Display window can be cleaned up by right-clicking and selecting *Clear Completed*. All completed or canceled processes will be removed from the Status Display. Lastly, the details of a process can be displayed by right-clicking on that process, and selecting *Show Details...* This will open a new tab in the Status Display, showing the details of the processes that are occurring (or have occurred) in that process.

## Using the Dataset Window

Many manipulations can be done on datasets via the tools within the dataset window. There are toolbars that appear across the top of a dataset window.

### Zooming a Dataset

Datasets can be zoomed in three ways:

- Click either the  icon to zoom in or the  icon to zoom out.
- To the right of the Zoom icons, click the image size percentage icon  and a drop down list will appear with pre-determined zoom percentages from which to select.
- To the right of the Zoom Percentage icon is the Aspect Ratio icon .

Click this icon and select *Maintain Aspect* from the drop down menu. Then left click and drag on one of the corners of the dataset window, and the dataset will zoom in and out as the window size increases and decreases.


*Note - Selecting Stretch Aspect can alter the dataset by changing the height and width disproportionately, depending on how the window is resized (if the width is changed more or less than the*

height, and vice-versa). Selecting Maintain Size will not change the size or aspect ratio of the dataset at all; only the window itself will be resized.

## Selecting a Projection

Projections can be selected from two places: the View menu, and the View icon in the dataset window.

(1) In the dataset window, click the projections icon  and the available projections appear in the drop down menu.


(2) Select the Sum projection icon  and click on it.

(3) The dataset window will refresh in Sum projection mode.

For information on how to select Projections using the View menu, see the View Menu chapter.

## Selecting a Single View

As with the Projections, Single Views can be selected both from the View menu and from the toolbar in the dataset window, but additionally, for Single Views, the dataset window contains tabs with which single views can be opened and closed.

(1) In the dataset window, click the Single View icon  and the available single views appear in the drop down menu.

(2) Select the XZ single view and click on it.

(3) The dataset window will refresh in the XZ view.

Alternatively:

(1) With the dataset still in the XZ view, click on the XY Tab on the top of the dataset.


(2) The XY view will open above the XZ view.

For information on how to select Single Views using the View menu, see the View Menu chapter.

(3) Perspective Tabs - These tabs, located along the right side and bottom of the image display window open new perspectives of that dataset within the same image display window. The tab along the right side opens and closes the YZ perspective, and the tabs along the bottom open and close the XY and XZ perspectives. Clicking on one of these tabs will open that tab's perspective. Once opened, clicking on that tab again will close that perspective.

## Creating a Region of Interest

Several functions in AutoQuant and 5D Viewer require a region of interest to be created. This

is done using the Region of Interest icon in the dataset window .

(1) Open a dataset.

- (2) Click the Region of Interest icon, a drop down menu will appear showing the available modes for drawing a region of interest (Square, Circle, Shape, Free hand, as well as options to Delete Selected, Delete All, Select Objects and an option to draw lines).
- (3) Select the Square region of interest.
- (4) Click and drag a region of interest. When the ROI is of the desired shape and size, click again to finalize the ROI. After doing that, a region of interest will appear on the dataset.
- (5) Repeat steps 2-4 several times, creating several regions of interest.
- (6) Click on one of the regions of interest in the dataset.
- (7) Click the Region of Interest icon again and select *Delete Selected*. The region of interest will be deleted.
- (8) Click the Region of Interest icon again, and select *Delete All* and all regions of interest will be deleted.

## Saving and Copying the Current View


There are two ways to save the current view of a dataset. One is to save the current view to a .tiff file, the other is to copy the current view to the clipboard, so that it can be pasted into a document.

- (1) Open any dataset (or, from the tutorial data, use colorpollenc.tif from the MultiChannel folder).



- (2) Click the Save Current View Icon.
- (3) Select the location and name the file *currentviewtest.tiff*, then click *Save*.
- (4) From the File menu, select *Open*, browse to the directory where the *currentviewtest.tiff* file was saved, select the file and then open it.




*Note - Only what is visible in the dataset window when the current view is saved will be shown (and available for viewing) in the .tiff file that is created using the Save Current View feature. If the dataset is zoomed in such a way that the edges are not displayed in the dataset window, those edges will not be included in the .tiff file.*

- (5) Make the original dataset active again by clicking on it.
- 
- (6) Click the Copy Current View Icon.
  - (7) Open a Microsoft Office document such as Word or PowerPoint.
  - (8) In the Edit menu of the Microsoft application, select paste, and the dataset's current view will appear.

## Linking Slice Viewers

Multiple Slice Viewers can be linked together to scroll through slices or time simultaneously, and in synchronization.

- (1) Open any 3D dataset (or, in the tutorial data, use the colorpollenc.tif dataset from the Multi-Channel folder). Once it displays, open the same dataset again.

- (2) In each of the datasets, select Slice Viewer from the Projection menu within the dataset window.
- (3) Click the *Synchronize* icon  in both dataset windows.
- (4) In the slice scroller toolbar along the left side of the dataset window, click the *Play* icon  and notice that both dataset windows scroll through the slices simultaneously.
- (5) Press the *Stop* icon . Notice that both slice viewers stop.

## Tutorials for File Menu Items

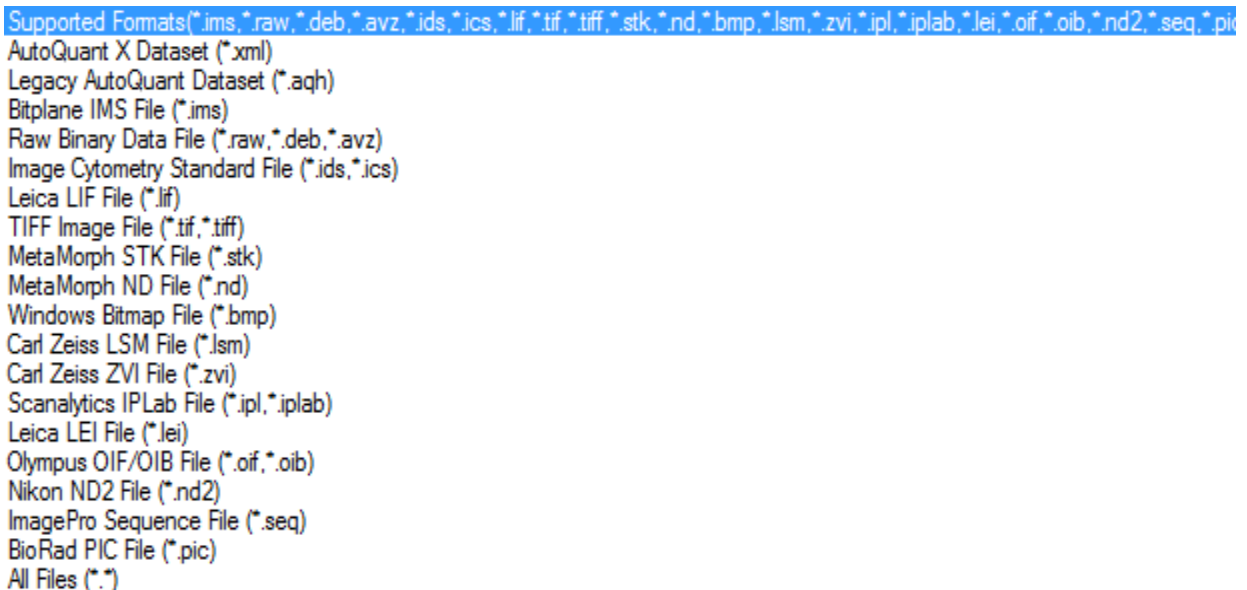
For this tutorial you may use your own data type (Fluorescence [Widefield], Laser Scanning Confocal, Brightfield [Transmitted Light], Spinning-Disk Scanning Confocal, Two-Photon Fluorescence or 2 Dimensional images) or you may follow along with the recommended dataset to use.

### Opening a Dataset

To open a dataset, click on *File* on the menu, and select *Open*. This will open a list of directory of files, from which a chosen file can be loaded into **AutoQuant** and AutoQuant 5D Viewer for processing. For a list of file types that can be opened see “Guidelines for naming your file(s)” below.

*Note: When opening a folder in any of the following tutorials, if the folder appears empty select All Files (\*.\*) in the Files of type field.*

- (1) From the File menu select Open. You will see a list of available file formats.



Supported Formats(\*.ims;\*.raw;\*.deb;\*.avz;\*.ids;\*.ics;\*.lif;\*.tif;\*.tiff;\*.stk;\*.nd;\*.bmp;\*.lsm;\*.zvi;\*.ipl;\*.iplab;\*.lei;\*.oif;\*.oib;\*.nd2;\*.seq;\*.pic;\*.xml)  
AutoQuant X Dataset (\*.xml)  
Legacy AutoQuant Dataset (\*.aqh)  
Bitplane IMS File (\*.ims)  
Raw Binary Data File (\*.raw;\*.deb;\*.avz)  
Image Cytometry Standard File (\*.ids;\*.ics)  
Leica LIF File (\*.lif)  
TIFF Image File (\*.tif;\*.tiff)  
MetaMorph STK File (\*.stk)  
MetaMorph ND File (\*.nd)  
Windows Bitmap File (\*.bmp)  
Carl Zeiss LSM File (\*.lsm)  
Carl Zeiss ZVI File (\*.zvi)  
Scanalytics IPLab File (\*.ipl;\*.iplab)  
Leica LEI File (\*.lei)  
Olympus OIF/OIB File (\*.oif;\*.oib)  
Nikon ND2 File (\*.nd2)  
ImagePro Sequence File (\*.seq)  
BioRad PIC File (\*.pic)  
All Files (\*.\*)

- (2) From the Tutorial Data directory select the Widefield folder. Select the dataset FitcDapi\_crop.tif and click on Open.

*Note: Various types of files including **AutoQuant file**, **AutoQuant 5D Viewer file**, 12 bit data, 16 bit data, Bio-Rad PIC, TIFF File, 8 bit character data, 32 bit float, and STK File can be handled. If your dataset is stored in a series of slice files, the software will recognize the series, and ask if you wish to load the entire series, or simply the file that you selected. If your dataset was stored as 12 bit data on a UNIX machine, and you are running the software on a PC, you must select 12 bit swapped (for UNIX) as the file type. This is also true for 16 bit data, and when transferring 12 bit and 16 bit data from a PC to a UNIX machine.*

- (3) The FitcDapi\_crop.tif dataset will be loaded and displayed in the XY Max Projection view.

*Note: If it is necessary to enter the dimensions of a particular dataset the first time it is loaded, you will be prompted to do so. The next time that dataset is selected, the dimensions will automatically be recalled for you.*

Once a file is opened, subsequent files with the same name (but from a different directory) will be numbered chronologically in the order they are opened. For example, if you have pollen.deb opened from the widefield folder, then you open another file name pollen.deb from a different directory, that file would be named pollen.deb(2), and subsequent pollen.deb files would be named pollen.deb(3), pollen.deb(4) etc.

## Saving a Dataset

This function saves an image or sequence of images as a specified file type.

*If the FitcDapi\_crop.tif file is not still open from the Open tutorial, open it now.*

- (1) Select Save As from the File menu and type BlueGreen as the File name. Click on the Save as type drop-down arrow and select STK FILE (\*.stk).

- (2) Click the Save button. This will save the file in the (\*.stk) format.

*Note: Do not close BlueGreen.stk. Please continue on to the next section.*

## Guidelines for naming your file(s):

*Note: **AutoQuant** and **5D Viewer** follow the same naming conventions as standard Windows applications.*

To move an **AutoQuant** or **5D Viewer** file into and out of a UNIX computer follow the file naming conventions of the UNIX operating system. For example, using an underscore for file names greater than 8 characters, and not using spaces or special characters.

## Save As

The dataset file can be saved as any of the following types:

Raw Binary Data File (\*.raw, \*.deb, \*.avz) - These are raw binary file types that can contain data represented with 8, 12, 16, or 32 bits per pixel.

TIFF (\*.tif) - Tagged Image File Format. This is a common format for image files.



Windows Bitmap File (\*.bmp) - This is a Windows bitmap file, often used to store image data on the Windows platform.

Image-Pro Sequence File (\*.seq) - This is an Image-Pro file format. It is based on the TIFF standard but it includes additional proprietary information used by Media Cybernetics' Image-Pro software.

Image Cytometry Standard File (\*.ids, \*.ics) - Image Cytometry is a standard file format used in some microscopy applications.

Bio-Rad PIC (\*.pic) - This file format is a proprietary file format used by Bio-Rad.

Bitplane IMS File (\*.ims) - This is a proprietary file format used by Bitplane software.

Carl Zeiss LSM File (\*.lsm) - This is a proprietary file format used by Carl Zeiss software.

MetaMorph STK File (\*.stk) - This is a proprietary file format used by Molecular Devices software.

Olympus OIF/OIB File (\*.oif, \*.oib) - This is a proprietary file format used by Olympus software.

Olympus Fluoview File (\*.tif) - This is a file format based on the TIFF standard. It includes proprietary information used by Olympus software.

Scanalytics IP Lab File (\*.ipl, \*.iplab) - This is a proprietary file format used by older Scanalytics IPLab software.

*Note: When you select Save As, if your dataset is in a Slice Viewer Projection and you have changed either the gamma, brightness, or flip settings, a dialog box will appear with the question Save data as shown? "No" will discard the gamma, brightness and flip adjustments. Selecting Yes will keep the gamma, brightness and flip adjustments.*

## **Saving the Application Layout**

The layout of the application can be saved to avoid having to customize toolbar locations in the future. As all of the toolbar options are moveable, this is a helpful tool. Once all of the toolbar items are in their desired locations, select *Save Layout* from the File menu.




## Editing the User Options

Since multiple users may use the same workstation with AutoQuant/5D Viewer on it, there is the ability to have multiple user options programmed for each user. The following features can be customized to each user:

- Temporary File Directory: This is where the temporary files created by the program will be stored.
- Default Output Directory: This is where the program will default to when a dataset is saved.
- Startup Directory: When first opening a file, the program will default to this directory.
- Number of Recent Files Shown: This determines how many files will appear in the recent files list on the File menu.
- NoCPU Intensive Operations: This determines how many CPU intensive operations (such as deconvolution) can be performed at once. Generally, however many CPU's the computer has is the maximum number of operations that should be allowed.
- Transfer to Imaris: Determines what bit depth should be used to transfer files to Imaris. Your options are:
  - UNSIGNED\_INT16\_WHEN\_POSSIBLE: If the maximum intensity of the dataset is equal to or less than 65536 (16-bit limit) the dataset will be transferred as 16-bit integer data. If the maximum intensity of the dataset is larger than 65535, it will be transferred as 32-bit floating point data.
  - FLOAT\_32: Forces data to be transferred to Imaris as 32-bit floating point data in all cases.

If Imaris is not installed, changing these options will have no effect on the behavior of AQX.

To change any of these settings:

- 1) From the file menu, select *User Options*.
- 2) Click on *Temporary File Directory*. This icon  will appear, click on that icon. A Windows Browser will appear.
- 3) Scroll to the desired location, highlight that folder, and click *OK*.
- 4) Repeat this procedure for any settings that need to be changed.
- 5) When all changes have been made, click on the OK icon . Or, to discard the changes, click the Cancel icon . This will cancel all changes, and close the User Options dialog.

## Exiting AutoQuant/5D Viewer

This function closes the software and all files that are open. You do not need to save the images generated in this tutorial section. From the File menu, select *Exit*. If there are any open, unsaved datasets, you will be prompted to save them.

## Recent Files List

This feature displays five of the most recently loaded and closed datasets. This list is found in the File menu underneath Exit. To open one of these most recent files, select the file from the list by clicking on it.

Please close all views before going on to the next tutorial.

## AutoQuant Connect

*AutoQuant Connect* is a new feature that allows you to automatically import and export your data to partnering software applications for additional analysis or visualization. Using AutoQuant Connect, you can import a "CONNECT Image Set" acquired in partner software applications (such as Image-Pro Plus or other software partners). The set includes the acquired image set, the image set meta-data, complete deconvolution settings, instructions, and completion commands. Upon receiving the "CONNECT Image Set", AutoQuant will queue all subsequent sets into the batch processing viewer to launch the next task. Each set is deconvolved using the defined settings with AutoQuant's industry leading algorithms. Once the deconvolution task is complete the image set along with complete meta-data is returned to the location of your choice. It can be returned to the connected software partner application, saved to a custom folder, or displayed in AutoQuant for review, visualization, or analysis.

## Export to Image-Pro

If you have Image-Pro installed on your machine, you can export files to it for analysis. To launch Image-Pro, select the Image-Pro icon on your toolbar. The Image-Pro icon and menu item are only available if you have Image-Pro installed on the same machine as AQX.

When you export a file to Image-Pro, it will be saved as a 16-bit \*.seq (sequence) file and opened in Image-Pro. Multi-channel files are opened as several single-channel files. The Image-Pro Color Composite tool is opened so that the files can be re-merged.

## **Export to Imaris**

If you have Imaris installed on the same machine as AQX, you will have an option on the File menu called Export to Imaris, as well as an icon on the toolbar. If you do not have Imaris installed on your machine, you will not have these options.

If you have these options, load a dataset into the AutoQuant X workspace. Click the Imaris icon, or select Export to Imaris from the File menu. The dataset is saved as an ICS/ICD pair and opened in Imaris. If the maximum intensity of the dataset is equal to or less than 65536 (16-bit limit) the dataset will be transferred as 16-bit integer data. If the maximum intensity of the dataset is larger than 65535, it will be transferred as 32-bit floating point data. You can force data to be saved as 32-bit floating point by going to the User Options and changing Export to Imaris to FLOAT\_32.

## **Export to MetaMorph**

If you have MetaMorph installed on the same machine as AQX, you will have an option on the File menu called Export to MetaMorph, as well as an icon on the toolbar. If you do not have MetaMorph installed on your machine, you will not have these options. Active files are sent to MetaMorph through a COM-based inter-application exchange.

## Tutorials for the View Menu Items

For this tutorial you may use your own data type (Fluorescence [Widefield], Laser Scanning Confocal, Brightfield [Transmitted Light], Spinning-Disk Scanning Confocal, Two-Photon Fluorescence) or you may follow along with the recommended dataset to use. All 3D datasets can be viewed in a Single view. All 3D datasets can be Zoomed, Enhanced, Inverted, and its Aspect Ratio changed. Choosing different views will display the image from different perspectives.

### Displaying the Data Browser


The Data Browser option, also known as the Data Manager, displays the Data Browser along the left side of the application window. For more information on the Data Browser, see the User Interface chapter.

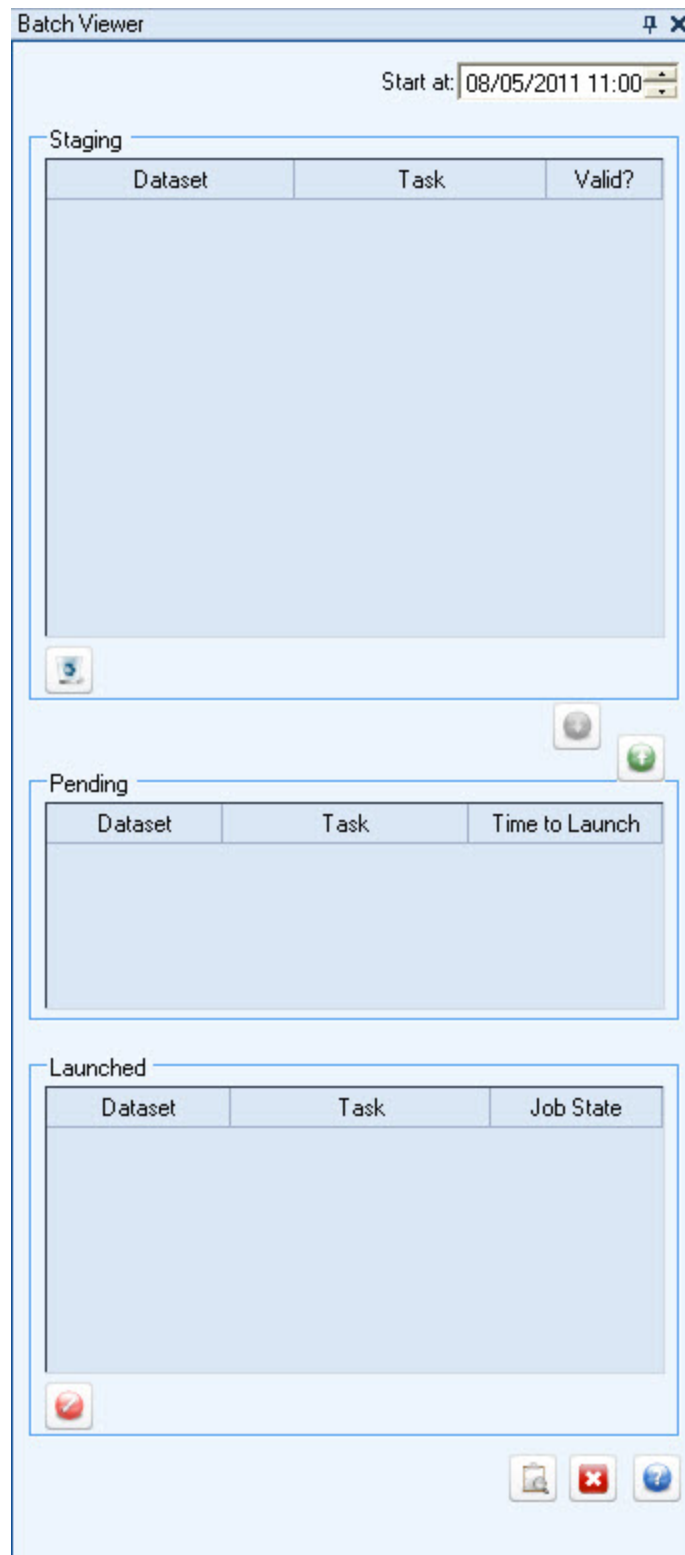
### Displaying the Status Dialog

To display the status of all operations, click on View->Display Status. This will display the Status Dialog at the base of the application window.

### Displaying the Task Batch

To display the Task Batch (the list of jobs in the Batch Processing queue), click on View->View Task Batch. This will open the Batch Processing Dialog along the right side of the application window.

- (1) Navigate to the Widefield folder under the Tutorial Data directory and open the dataset FitcDapi\_decon.tif. This dataset has had the Standard Settings and the Expert Settings set up in the previous tutorial. The information is contained in its header file, and the header file is automatically read when the dataset is loaded.
- (2) From the Deconvolution menu, select 3D Deconvolution. The 3D Deconvolution box will appear. Extensive research and development has made entering or even changing the settings not necessary for most uses. As such, leave the settings at their default setting.
- (3) Click the Batch icon  in the 3D Deconvolution Dialog. The 3D Deconvolution dialog will close.
- (4) Click the Batch icon in the toolbar. This will open the Batch Viewer Dialog, showing the task just added to the batch. If any settings were incorrect, there will be a red "X" in the Valid? column of the Staging Area. Clicking that icon will bring you to the dialog for the operation listed (in this case, the 3D Deconvolution dialog), so that the setting can be fixed.



- (5) Select a different operation (3D Inverse Filter) from the drop down menu in the Operation frame of the Batch Viewer dialog.
- (6) In the top right part of the Batch Viewer is the *Start at* section. This is where you can schedule a batch by entering a date and time (in the future). Set the time for 2 minutes ahead of the current time.
- (7) Click the arrow to the left of where it says FitcDapi, highlighting the entire row. At the bottom of the Staging Area, click the down arrow to move the process into the pending area.
- (8) In roughly 2 minutes, the process will move into the Launched area, where the 3D Inverse Filter will begin processing.

## Displaying a Dataset

Sometimes, a dataset gets closed out of the workspace, but it is still left in the Data Manager. To display this dataset again, highlight the dataset in the Data Manager by clicking on its name, then click View->Display Dataset. The dataset will now be displayed in the workspace.

## Using the Max, Min and Sum Projections

There are various projections in which datasets can be displayed. These are Maximum Projection, Minimum Projection and Sum Projection. Each of these projections can be selected from the View menu. Selecting the desired projection will display the dataset in that projection, without opening a new dataset window.

## Using the Slice Viewer

To display and view a 3D image one slice at a time, use the *Slice Viewer*. This function generates slices through the image “optically” and then displays the “face” of each slice in the Slice Viewer. The thickness of each slice corresponds to the Z spacing or step size. Viewing in the Slice Viewer is a good way of examining the degree of optical sectioning in the dataset.

It is advisable to examine datasets in a Slice Viewer because it allows for viewing of subtle and not so subtle changes from slice to slice.


Make the Pollen.deb: XY Max Projection dataset active by clicking within its view.

- (1) From the View menu, select Slice Viewer to generate an optical slices projection of the dataset from the XY perspective.

*Note: When a projection operation is selected, the resultant view formed depends upon the currently active view. For example, in the previous step the XY Max projection was the active view. Thus, when the Slice Viewer menu item was selected, optical slices were created from an XY per-*

*spective.*

(2) To step through the optical slices, move the slider on the left side of the dataset window. This can be accomplished by clicking on the button and dragging it up and down.

(3) The Slice Viewer also has the ability to cycle through all the slices automatically. To do this, click the *Play* icon  on the left side of the dataset window.


(4) Clicking the *Stop* button  will halt the Slice Viewer.

*Note: Do not close Pollen.deb: XY Max Projection. Please continue on to the next section.*



## Synchronizing Slice Viewers

To compare multiple Slice Views simultaneously, use the *Slice Synchronizer* feature. This feature links up multiple Slice Viewers to scroll through the slices simultaneously.

- (1) Open any 3D dataset (or, in the tutorial data, use the colorpollenc.tif dataset from the Mult-Channel folder. Once it displays, open the same dataset again.
- (2) In each of the datasets, select Slice Viewer from the Projection menu within the dataset window.
- (3) Click the *Synchronize* icon  in both dataset windows.
- (4) In the slice scroller toolbar along the left side of the dataset window, click the *Play* icon and notice that both dataset windows scroll through the slices simultaneously.
- (5) Press the *Stop* icon. Notice that both slice viewers stop.


If there are datasets that have different numbers of slices, (e.g., one dataset has 50 slices, one has 30 slices and one has 90 slices), all datasets will scroll through all of their slices, the smaller datasets will pause at the end while the larger datasets finish scrolling. Once the largest dataset has completed scrolling, all datasets will start scrolling again at slice 1.

## Scrolling through Slices

The large scrollbar on the left side of the Slice Synchronizer dialog box allows you to move through the slices in a manner similar to that of a single optical slice viewer (i.e., the arrows in the scroller move one slice at a time, and multiple slices can be jumped at a time by moving the slider about).

If the *Play* button  is clicked, the views automatically scroll through their slices.

## Adding and removing Datasets from Slice Synchronization

Datasets can be added and removed from Slice Synchronization by clicking the Slice Synchronizer icon  in that dataset window. If a dataset is synchronized, a blue outline will appear around the Slice Synchronizer icon. If it is not synchronized, there will be no outline around the icon.

## Selecting a Single View

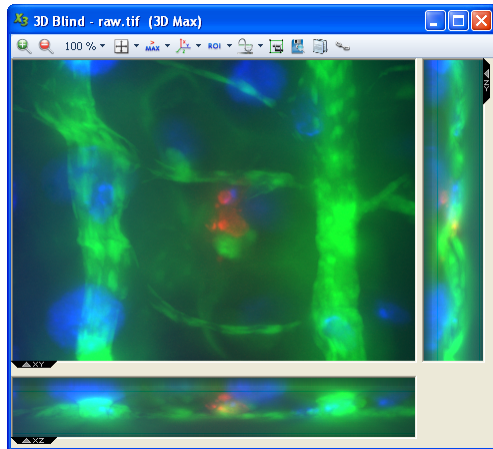
There are several views in which the dataset is presented. The dataset can be viewed from three orthogonal perspectives. The choices available are: XY, XZ, and ZY where X represents the image width, Y the image height, and Z is the depth or may be thought of as the optical axis in microscopy applications. When a choice is made, the single view that is generated will be for the currently active projection. For example, if a Sum Projection is the active view, selecting ZY will generate a ZY Sum Projection of the view.

- (1) Navigate to the TLB folder and open the TLBview.tif dataset. Select XZ from the Single View option under the View menu.

*Note: Transmitted Light Brightfield datasets will open in the Min Projection view.*

- XY - This is the view of the “face” of the dataset or the front view of each slice.
- XZ - This is an “edge-like” view of the dataset or of one of its slices. It can be thought of as a horizontal slice through the XY view.
- ZY - This is a “side” view of the dataset or each slice. It can be thought of as a vertical slice through the XY view.

## Triple View



The Triple View creates three orthogonal views of the dataset. On the top left of the Triple View display is the XY view, on the top right is the ZY view, and on the bottom left is the XZ view of the dataset. The TLBview.tif:XY Min Projection from the previous section should be active.

- (1) From the View menu, select Triple View. The XY, YZ, and XZ views of the TLBview.tif dataset are displayed in the Triple View box.
- (2) The Triple View can also be used on datasets displayed in a Slice Viewer projection. From the View menu select Slice Viewer. The Triple View will be displayed in a Slice Viewer.
- (3) The XY, YZ, and XZ Optical Slice Viewers are displayed in the Triple View box. The red lines indicate where each slice is in relative position to the others in the dataset.

*Note: Do not close TLBview.tif:XYMin Projection. Please continue on to the next section.*

## **Saving the View**

To save the view of the active dataset as a .tiff file, use the *Save View* feature. This will save only the image within the dataset, without the borders and toolbars. Alternatively, click the Save Current View icon in the dataset window.

## **Copying the View**

To copy the image within the active dataset (for pasting into a document, for instance), use the *Copy Current View* feature. The current view of the active dataset is now in the clipboard, and can be pasted into a document or image. Alternatively, click the Copy Current View icon in the dataset window.

# Tutorials for Processing Menu Items

## Cropping an Image

Once an ROI has been created (see Creating a Region of Interest on page 58) the dataset can be cropped in three different ways: Crop the Stack, Slice by Slice or by entering crop dimensions. Cropping the image creates a new dataset based on whatever pixels fall inside that selected region. The following tutorials take you through the steps of each different cropping method.



Cropping and Reshaping

**3D Blind - raw.tif**

Crop Selected ROI   Crop Dimensions   Reshape Dataset

Enter the new dimensions for the image:

Top Left Front Corner		Bottom Right Back Corner	
X	1	X:	402
Y	1	Y:	300
Z	1	Z:	60

New Dimensions

Width	Height	Depth
402	300	60

Time Point Options

☒ Crop Time Series   ☐ Crop Time Point

✓ ✗ ?

Cropping and Reshaping

**3D Blind - raw.tif**

Crop Selected ROI   Crop Dimensions   Reshape Dataset

☐ Swap Z/T

Enter the new dimensions size for the Dataset

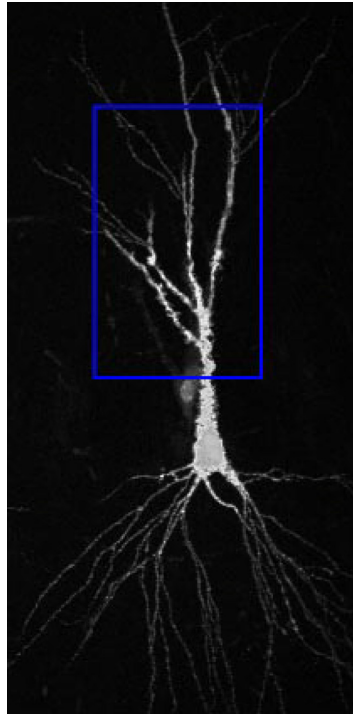
Time :	1	Dimensions product : 180
Depth :	60	
Channel :	3	

Select new dimensions order for the Dataset

Channel\_Time\_Slice


✓ ✗ ?

- (1) Go to File - Open, select the Confocal folder from the Tutorial data directory. Select the Neuron.deb dataset and open it.



- (2) From the Processing menu select Cropping and Reshaping.
- (3) If an ROI had been drawn, its top left corner position and dimensions would be displayed in the *Crop Selected ROI* tab. For this tutorial, you will enter an ROI manually to crop a section like the one displayed above.
- (4) Click on the *Crop Dimensions* tab. In the Top Left Front Corner frame, enter the following parameters; X:51, Y:72, and Z:15. In the Bottom Right Back Corner frame, enter the following parameters; X:150, Y:272, and Z:37.

*Note: In a time-series dataset, by default, all time points will be cropped. To crop only certain time points, deselect the time points that you do not want cropped in the Data Manager section on the left side of the application.*

- (5) Click the OK icon . The image will be cropped and displayed in the viewing window.
- (6) From the File menu, select Save As.
- (7) When prompted for a file name to save the cropped image type Neuron\_crop1 in the File name field and click the Save button. You may close the Neuron\_crop1 view.

*Note: Do not close Neuron.deb. Please continue on to the next section.*

## Extending Slices

In order to protect meaningful data in the outermost slices of a 3D dataset, the image can be extended (have additional optical slices added to it) along its Z axis, using the *Extend Slices* feature.

This function allows you to generate additional image slices based on linear extrapolation, and appends them to the top and bottom of an image. It is not the same as adding blank slices to the top and bottom of a stack as mentioned in (1) below.

An extended dataset will have false slices attached to its top and bottom. These false slices serve the purpose of a buffer to keep the guardband region from overlying meaningful features, thereby keeping the meaningful data from being obscured in the deconvolution process.

This tutorial continues with the Neuron.deb dataset from the Crop tutorial above.


(1) From the Processing menu select Extend Slices. The Extend Slices Dialog will appear. You have the option of extending the data by using false slices or by using the Zero-fill option which when selected, adds blank slices to the Top and Bottom of the dataset devoid of any information.

The Extend Depth indicates how many slices you want to add to the top and bottom of your sample.

In the Extend Depth box enter 4 for the Top and Bottom field values.

The new Depth value will be displayed in the New Dimension frame.

Leave the Zero Pad box unchecked.

(2) Click the OK icon . The extended dataset will be displayed. Click on the XZ view icon, and notice the difference between the real data slices and the false slices just generated.


(3) Close all views before going on to the next tutorial.

## Resizing an Image

In order to reduce data storage requirements, reduce pixelation (blockiness) in small datasets, and improve rendering, a dataset can be resized using the *Resize Image* feature. This function allows you to change the overall size of an image. There are two methods for doing so, the Linear and the Ideal method. The Ideal method is recommended, however the Linear method is somewhat faster than the Ideal method.

Resizing a dataset resamples the data to produce a new stack of optical sections with dimensions different than those of the original stack, unlike changing the size of the image on the screen.

*Note: Resizing an image and Sizing (zooming) an image are different functions.*

- (1) Navigate to the Widefield folder and open FitcDapi\_crop.tif.
- (2) From the Processing menu, select Resize Image. The Resize Image dialog box appears. This dialog box contains three ways in which an image can be resized: Dimensions, Voxel Size or Resize Factor. There is also the option of choosing Linear or Ideal as the method employed in the Resize function. As one parameter is changed the others are automatically updated to preserve the true image size.
- (3) In the Resize Methods section, select Linear.
- (4) In the text boxes, enter and verify the following:  
  
For the Dimensions (New) X, enter a value of 250.  
For the Dimensions (New) Y, enter a value of 250.  
For the Dimensions (New) Z, enter a value of 100.  
  
Voxel Size:  
  
Verify the Voxel Size (New) X is 0.108.  
Verify the Voxel Size (New) Y is 0.144.  
Verify the Voxel Size (New) Z is 0.196.  
  
Resize Factor:  
  
Verify the X Resize Factor (New) is 2.77777.  
Verify the Y Resize Factor (New) is 2.083333.  
Verify the Z Resize Factor (New) is 2.040816.
- (5) Click the *OK* icon . When completed an XY Max projection of the newly resized dataset will be created.

Conversely, if a dataset is too large to view easily, the volume can be resized to a smaller three-dimensional volume.

Please close all views before continuing on to the next section.

## Performing CCD Correction

The CCD Correction function performs the Flat-Field and Bias Field corrections and the corrections for the camera and lamp flicker, whose principles are described in the Guidelines for Collecting 3D Image Data section of this manual. The Data Correction function is for widefield datasets (fluorescence or brightfield) only. Do not use it for confocal datasets.



It is recommended that you do a CCD Correction on your dataset before processing a Widefield dataset (fluorescence or brightfield).

## Slow Scan Cooled CCD

The CCD Correction feature can correct for Slow Scan Cooled CCD cameras, when charge integration time is selectable.

(1) Navigate to the TLB folder in the Tutorial Data directory. Set the Files of type to All Files and select star.1. Next, open star.bs0, star.bs1 and star.lt0. These will be the Flat Field, Bias Field 1 and Bias Field 2 images.

*Note - With your own datasets, If you have not collected these fields then their selection boxes should be unchecked. The recommendation is to collect and load all three Field image stacks.*

(2) From the Processing menu, select CCD Correction and click on Cooled CCD. The Cooled CCD Data Correction box will appear.

Cooled CCD Correction

**3D Blind - raw.tif**

Files

☐ Flat Field

☐ Bias Field 1

☐ Bias Field 2

Exposure Times

Average: 0.00 sec

Flat Field: 0.00 sec

Bias Field 1: 0.00 sec

Bias Field 2: 0.00 sec

Bad Pixel Treatment:

None


✓ ✗ ?

- (3) To load the Flat Field file (\*.lt0), check the Flat Field box and select star.lt0 from the drop down list.
- (4) To Load the Bias Field 1 file, check the Bias Field 1 box and select star.bs0 from the drop down list. Click Open.
- (6) To Load the Bias Field 2 file, check the Bias Field 2 box and select star.bs1 from the drop down list.
- (8) For the tutorial data, enter the following parameters:  
Average: 0.03  
Flat Field: 0.03  
Bias Field 1: 60  
Bias Field 2: 120

*Note - When using your own datasets, these numbers should have been recorded and archived when the image stacks were collected. The Average exposure time is the exposure time of the CCD camera that was set while the 3D dataset was collected. To learn how to obtain these numbers refer to the operator's manual of your camera, check the documentation of the software used to collect the image stacks, or contact the camera and data collection software manufacturer.*

- (9) The Bad Pixels Treatment setting should have the default value of None.

*Note - Bad pixels occur when the dark current of a pixel is so severe that it causes a bright spot in the picture. Most image stacks do not require any treatment of bad pixels. Therefore the recommended default selection is None. If the selection of None causes the corrected pictures to become black, this is an indication that the dataset requires the treatment of the bad pixel(s). In this case, the recommended Bad Pixel Treatment setting is Median Filter. This is the most accurate means of treating bad pixels. It replaces the bad pixel's intensity with the median value of its nearest neighboring pixels. The Wiener Regularization setting is available because it executes much faster than the Median Filter setting, but it is not as accurate as the Median Filter setting.*

- (10) Click the OK icon  to begin the data correction of the Slow Scan Cooled CCD dataset. Data correction will take approximately 1-2 minutes depending upon your machine.
- (11) Once the image has been corrected, the result will open in a new dataset window.


Close all views before continuing on to the next section.

## High Speed CCD

The High Speed CCD selection under CCD Correction is for datasets that were collected from a standard Video camera, an intensified CCD camera, a standard digital camera or any other camera besides the Slow Scan Cooled CCD camera. There are two tabs: the Files tab and the Parameters tab.

- (1) Navigate to the Data Correction folder under the Tutorial Data directory. Set the Files of type to All Files. Open the Strfish.tif dataset. Create the ZY view of the dataset.
- (2) From the Processing menu select CCD Correction, and click on High Speed CCD. The High Speed CCD Data Correction Dialog will appear.
- (3) Next, open the StrfishFfv.tif file from the same directory. Check the Flat Field box, then select StrfishFfv.tif from the drop down list.
- (4) Open the StrfishBfv.tif file from the same directory. Check the Bias Field box, then select StrfishBfv.tif from the drop down list.

*Note - There is only one Bias Field for High Speed CCD Data Correction, because Dark Current does not effect the image in this situation.*

- (6) Enter the value for the frame(s) per optical slice. Leave the default setting of 1 for this example. For Bad Pixel Treatment select Median Filter. Click the OK icon  and **AutoQuant** will correct the High Speed CCD dataset.

*Note - To slow down the viewing time per each frame increase the frame(s) per optical slice value.*

## Correcting Attenuation

If your dataset suffers attenuation due to any of the following issues:

- Excitation light absorption
- Fluorescent emission light absorption
- Photobleaching
- Changes in the optical point spread function as a function of depth, which in turn is fundamentally due to refractive index mismatches between the sample embedding medium, coverslip, and lens immersion medium.

The *Attenuation Correction* feature can correct the attenuation as a function of depth into the sample. To determine if a dataset requires Attenuation Correction, look for darkening within an image as you go from the top to the bottom of the view.

- (1) From the File menu, select Open. Navigate to the Confocal folder and select RedNuc.0.
- (2) From the View menu, select Projections, then select Sum projection. From the View menu select Single View and click on XZ to generate an XZ View of the Sum Projection. Notice how the dataset appears darker as you go from the top to the bottom of the view.
- (3) From the Processing menu select Attenuation Correction.  
The Attenuation Correction will be performed automatically on the dataset, and the result will be displayed in a new dataset window.

- (4) Compare the raw XZ Sum Projection dataset to the newly corrected XZ Sum Projection dataset. The bright to dark effect has been diminished in appearance.


Attenuation Correction is most often applied to Confocal datasets where there is frequently a decrease in image intensity due to attenuation, light scatter, and spherical aberration as the depth into the sample is increased. Apply Attenuation Correction to the raw dataset before processing.

- (5) Close all views before going on to the next section.

## Equalizing the Background

*Note: This section applies to users of **5D Viewer** only.*

In order to equalize a dataset that has an uneven background, use the *Background Equalization* feature. This function allows you to remove unnecessary background from an image.

- (1) From the File menu select Open. Navigate to the SmallHip folder and select SmallHip.tif.
- (2) From the Processing menu select Background Equalization. The Background Equalization box will appear. Adjust the Background by typing in the Background Equalization box or by scrolling the bar until the Maximal Equalization value equals 45%.
- (3) Check the Remove Negative Intensities box to enable it. When this box is checked any background points that have negative intensity values will be removed. These negative pixel intensity values can come from the autoscaling function in the software or from artifacts in the dataset itself. Click the *OK* icon . The background adjusted dataset will be displayed.

## Subtracting the Background

Sometimes, image analysis is performed more accurately with the background removed from the dataset, especially if the dataset has background noise. For instance, the FRET or Ratiometrics features provide more accurate results with the background removed. To accomplish this, use the *Subtract Background* feature. Removing the background is different than equalizing the background, in that this feature actually removes the background, rather than evening it out.

There are five different ways to subtract the background from a dataset:

- ROI - This involves drawing an ROI around an area that is known to be the background. The algorithm will remove all data with the same or lower intensity as the intensity within the ROI.
- Histogram Peak - Selecting this algorithm will remove the most commonly occurring intensity level, using the assumption that the majority of a dataset is background and contains no relevant data.
- Minimum Value - Selecting this algorithm will remove the lowest intensity value from the image.

- Constant Value- This involves entering an intensity value, below which, all intensities will be removed.
- Background Image - This involves taking an image of a blank specimen (which will create a dataset that is entirely background) and selecting this dataset from the drop down menu (the dataset needs to already be opened for this to happen). The algorithm will then analyze the intensity in that image, and subtract that intensity from the desired dataset.

- (1) Click on File from the main menu and select *Open*. Navigate to the Ratiometrics folder and open *HighCal.001*.
- (2) Select *Background Subtraction* from the Processing menu. The Background Subtraction dialog box will open.
- (3) There are five different methods of background removal: Region of Interest (ROI), Histogram Peak, Minimum Value, Constant Value, and Background Image.

-For the ROI option, create a region of interest (see page 58 for instructions on creating a ROI) in the image that is empty (background) by clicking the mouse, dragging it to create a box, and then releasing. Make sure that nothing is inside this box.


-The Histogram Peak option requires no input, it looks at a histogram of the intensities, then removes the highest occurring intensity (this is typically the background, in an image in which the object takes up less than half of the image).

-The Minimum Value option also requires no input, and it removes the minimum intensity from the image.

-The Constant method allows an intensity to be entered into the text box, and anything at or below that intensity level will be removed from the image.

-The Background Image option requires an image to be acquired with no object in it (a blank image). Select this file from the drop down menu. If the background image file has not been opened, open it using the *Open* button on the main toolbar.

Click on the radio button for the desired method of background removal.

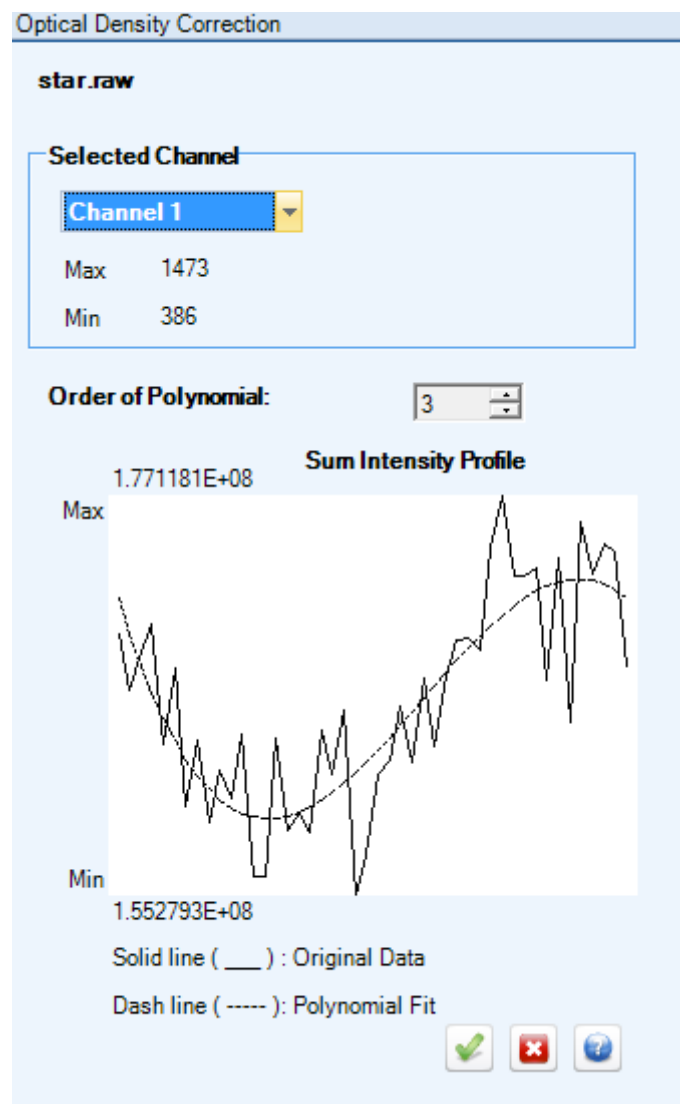
- (4) Click the OK icon . A new dataset window will open with the background removed.

## Correcting Optical Density

To correct fluctuations in the image intensity values across the depth of an image, use the *Correct Optical Density* feature. This only works on images with depth > 1. The image intensity value often fluctuates erroneously because of random flicker from the camera shutter or the lamp instabilities. Flicker occurs with nearly all widefield microscope systems. It is due to several causes. Most Cooled CCD cameras have a randomness in their shutter's speed. This shutter speed fluctuation causes variations on the order of several milliseconds (typically) from one exposure to the next. The effect can be seen best in a side-view projection of a dataset.


To determine if your dataset needs Optical Density Correction look for abrupt fluctuations in image intensity summations from one depth to the next.

- (1) From the File menu select Open. Navigate to the TLB folder and click on star.1. In the Files of type field select *All Files (\*.\*)*, if the folder appears empty.
- (2) In order to appreciate fluctuations in image intensity, an XZ projection needs to be generated. From the View Menu, click on Projection, then select Sum Projection. Then, from the View menu, click on Single View, and select XZ View. Notice the horizontal lines in the dataset.
- (3) From the Processing menu select Optical Density Correction. The Optical Density Correction box will appear.



The Order of the Polynomial is set at 3. This is the default value that can be used to correct for optical density. The order of Polynomial can be changed by clicking the arrows in the box either up or down, raising or lowering the Order of Polynomial, respectively.

*Note: Increasing the order of the Polynomial will increase the correction in the image intensity to a certain point, after which possible aberrations will be observed depending on the dataset.*

(5) Change the Order of Polynomial to 4. Notice how the dotted line more closely follows the intensity profile of the image. Click the OK icon  .

(6) Place the star.1.tif\_OpticalCorrection XZ Sum Projection view below the star.1 XZ Sum Projection view, and compare the differences between the views.

Notice how the Optical Density Correction has reduced the amount of horizontal lines in the corrected dataset.

*Note: The intensity profile displayed is for a Grayscale dataset. For a multi-channel dataset the intensity profile displayed is Channel 1, which is generally the red channel. This is the default. The intensity profile for each channel can be viewed separately by making the appropriate selection under Selected Channel. For each channel an order of the polynomial can be chosen which best fits the image intensity profile (each channel can use a different Order of Polynomial).*

Other types of cameras often have a similar problem. Ordinary Video Rate CCD cameras, intensified CCD cameras, digital cameras and other types of cameras will show this flicker effect. There are several factors that contribute to this effect, and it is unknown to what degree each factor contributes. Some camera manufacturers may claim that their cameras do not have these imperfections. Even so, the effect is still likely to arise and is due at least in part to other factors in the optical train that the camera manufacturer cannot control. Such factors include: arc-lamp flicker, instabilities in the power supply to the lamp, instabilities in the power supply to the camera, fluctuation in the 110 Volt A.C. power that supplies the camera electronics, microscope and other electronics in the microscope system. Other remote possibilities are adverse laboratory conditions, temperature changes, room lights being turned off and on, and movement in the room causing shadows over the microscope.

Although the source of these flicker effects is difficult to pinpoint exactly, it is certain that they occur and that they occur regularly.

(7) Close all views before continuing on to the next section.

## Aligning an Image

*Note - This feature (including Manual, Automatic and Channel to Channel Alignment) is only available as an added plug-in. Contact your dealer for purchase information.*

For several reasons, a 3D dataset can be misaligned, and in several different ways. There can be shift in between slices, a live cell image can move during acquisition, or rotate or even become warped, and a multi-channel dataset can have misalignment from channel to channel. The *Image Alignment* feature can correct for these problems, producing an aligned image for easier viewing and more accurate analysis.

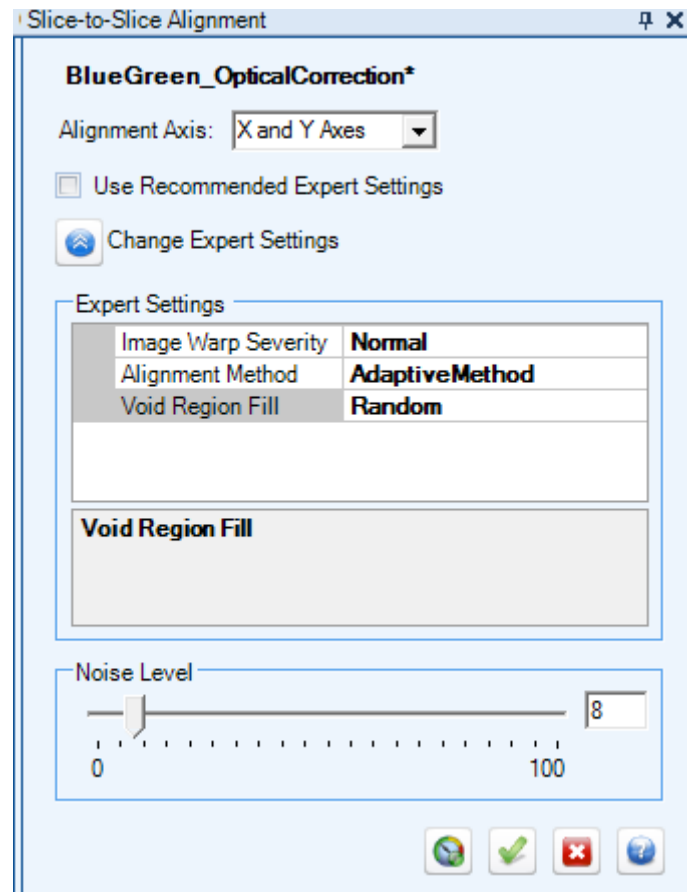
## Slice to Slice

### ***Automatic Alignment***

This feature corrects for X & Y translation and warp (nonlinear distortion) between frames. Misalignment and warp can be caused by stage motion, movement of the sample during live imaging and other causes.

- (1) Open the Unaligned.avz file from the Ocular folder in the Tutorial Data directory.
- (2) From the Processing menu select Image Alignment, then select Slice-to-Slice. The Image Alignment box will appear. Click the Advanced Settings button, and then the dialog will appear as shown here:

:





## **Alignment Axis**

This features allows the alignment to be restricted to the X axis, the Y axis, or both axes. The default setting is Both X and Y Axes.

### **X**

This setting will only align an image horizontally.

### **Y**

This setting will only align an image vertically.

*Note-Selecting both X and Y will align the image on both its horizontal and vertical axes.*

- (3) Leave the default setting of Both X and Y axes enabled.

## **Advanced Settings**

Unchecking the *Use Recommended Expert Settings* checkbox and then clicking the *Change Expert Settings* button will display additional Slice-to-Slice Image Alignment settings. These settings are already set based on the image information, however, if you feel that these are not as accurate as you would like them, they can be fine tuned.

### **Image Warp Severity**

This feature corrects for warp due to motion between frames. The more serious the warp, the longer the processing time. There are five available settings: None, Tiny, Normal, Severe and Very Severe.

- (4) Verify that the Image Warp setting is None (this is the default).

### **Alignment Method**

This is the frame to which all other frames are aligned.

- Adaptive Method: This function finds the base slice which has the highest similarity in features and structures to all other slices.
- Compare to Nearest Slice: This function compares each slice to its nearest slice in order to align the dataset.
- Always Compare to Base: This function compares every slice in the dataset to the base slice (slice 1).

- (5) Select Adaptive Method.

### **Void Region Fill**

This function will put a frame like cover over the areas that no longer contain data. After the slices are aligned, gaps and spaces are created at the edges. This function chooses how those areas are filled in.

*Note: The Frame may have straight, scalloped or irregular edges to it.*


- Black: This will put a black frame on the edge of the image where the aligned data has left a void. This is recommended for Fluorescence and Darkfield datasets.

- White: This will put a white frame on the edge of the image where the aligned data has left a void. This is recommended for Transmitted Light Brightfield datasets.
- Random: This will place a frame on the edge of the dataset made up of random pixel values where the aligned data has left a void. This is recommended for datasets that will be deconvolved or for images that have obvious structures at the edges of the frame.

(6) Select Random from the drop down menu.

### Noise Level

This function allows you to set the noise threshold for which the Image Alignment feature will compensate.

(7) Click the OK icon . The application will launch the alignment process and will display the results upon completion. This result will need to be saved.

(8) Create Slice Viewers of the original and the aligned images, then click on the Slice Synchronizer icon. Scroll through the datasets. Notice the motion between the slices in the Unaligned.avz dataset, as compared to the Aligned.avz dataset.

To add the alignment to the Batch, click the *Batch* icon .

Close all views before going on to the next section.

## Channel to Channel

The Channel to Channel alignment feature allows the user to align multi-channel datasets that have shift between the channels due to a filter cube or some other image acquisition problem. To open the Channel to Channel Alignment feature, go to the Processing menu option, select Image Alignment, then Channel-to-Channel. The Channel-to-Channel Alignment dialog will appear.

### Alignment Axis

This section allows the user to select the axis, along which the image needs to be aligned: along the X axis, along the Y axis, or along both axes. The default setting is to align to both axes.

### Base Slice

This section is where the Base Slice (the slice to which all others will be aligned) is chosen. The default setting is to let the application select the base slice. Deselecting *Set Automatically* will enable the slice scroll bar, with which the user will then need to scroll to the desired base slice. Alternatively, the user can enter the slice number into the text box to the right of the scroll bar.

## **Advanced Settings**

Deselecting the *Use Recommended Expert Settings* and then clicking the *Change Expert Settings* button will open the Advanced Settings section. It is recommended that the default settings be used. Clicking this button again with the Advanced Settings dialog showing will close the Advanced Settings section.

### **Image Warp Severity**

This feature corrects for warp due to motion between frames. The more serious the warp, the longer the processing time. There are five available settings: None, Tiny, Normal, Severe and Very Severe.

### **Void Region Fill**

This function will put a frame like cover over the areas that no longer contain data. After the slices are aligned, gaps and spaces are created at the edges. This function chooses how those areas are filled in.

*Note: The Frame may have straight, scalloped or irregular edges to it.*

### **Black**

This will put a black frame on the edge of the image where the aligned data has left a void. This is recommended for Fluorescence and Darkfield datasets.

### **Random**

This will place a frame on the edge of the dataset made up of random pixel values where the aligned data has left a void. This is recommended for datasets that will be deconvolved or for images that have obvious structures at the edges of the frame.

### **White**


This will put a white frame on the edge of the image where the aligned data has left a void. This is recommended for Transmitted Light Brightfield datasets.

## **Noise Background**


### **Set Level**

This option sets the noise level for the dataset. Use the slider to select what percentage of the dataset is noise.

## **OK**

Clicking the *OK* button  will initiate the Channel to Channel Alignment feature, and will create a new, aligned image, that will need to be saved.

## **Cancel**

Clicking the *Cancel* button  will cancel and close the Channel to Channel alignment dialog. No aligned image will be created or saved.

## Inverting Image Data

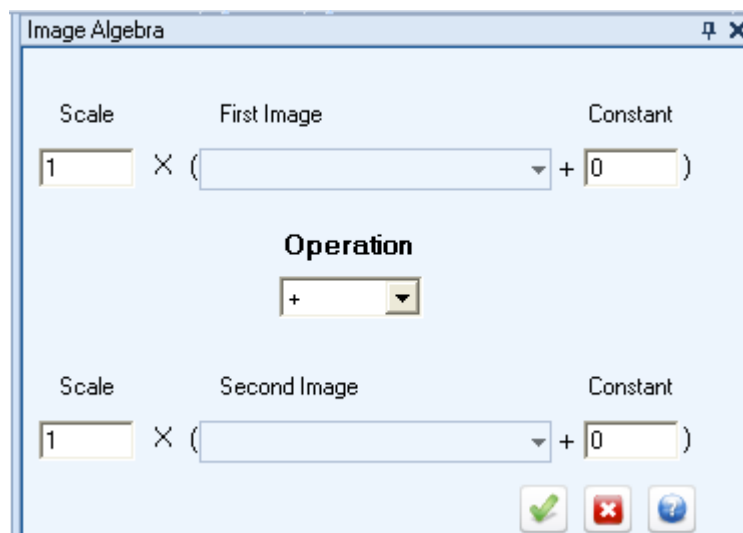
In order to create a new dataset with an inverted intensity scale, use the *Invert Image Data* feature.

- (1) From the File menu, select Open. Navigate to the Small Hip folder, and open SmallHip.tif.
- (2) From the Processing menu select Invert Image Data. The SmallHip.tif dataset will be processed automatically and displayed in a new dataset window upon completion of the inversion.

## Image Algebra


*Note: This section applies to users of AutoQuant 5D Viewer only.*

In order to perform a mathematical operation on two volumes, use the *Image Algebra* feature. The volumes must be the same dimensions. The available Image Operations are: Addition, Subtraction, Multiplication, Division, AND, OR, and XOR. If Subtraction is selected, the Second Image will be subtracted from the First Image. If Division is selected the First Image will be divided by the Second Image. The two original files will remain intact, and a new data file will be created with the processed results. To use the AND, OR and XOR one of the datasets must be optically sliced (in a Slice Viewer projection), in the XY view and segmented.



- (1) Go to File-Open. From the Widefield directory open the PollenPlus5.deb dataset.
- (2) From the Processing menu select Image Algebra. When the Image Algebra dialog appears, from the First Image drop down menu, select PollenPlus5.deb, and select Pollen.deb from the Second Image drop down menu. Leave the Scale for both the First and Second Image at 1. This is the default value. The Constant Value field will be inactive.

*Note: To add a Constant Value to an Image, select None for the Second Image and enter a number in the Constant Value field. The Constant Value is for intensity offset. Every pixel's intensity will be either decreased (by entering a negative number) or increased (by entering a positive number) by the Constant Value number.*

- (3) Select Subtraction from the Operation drop-down menu, and click the OK icon .

**5D Viewer** will subtract the Pollen.deb dataset from the PollenPlus5.deb dataset, creating a new file.

- (4) The resultant dataset, PollenPlus5\_AlgebraResult, may now be saved as PollenPlus4.deb.
- (4) Close all views before going on to the next section.

## Tutorials for Deconvolution Menu Items

### Performing 3D Deconvolution

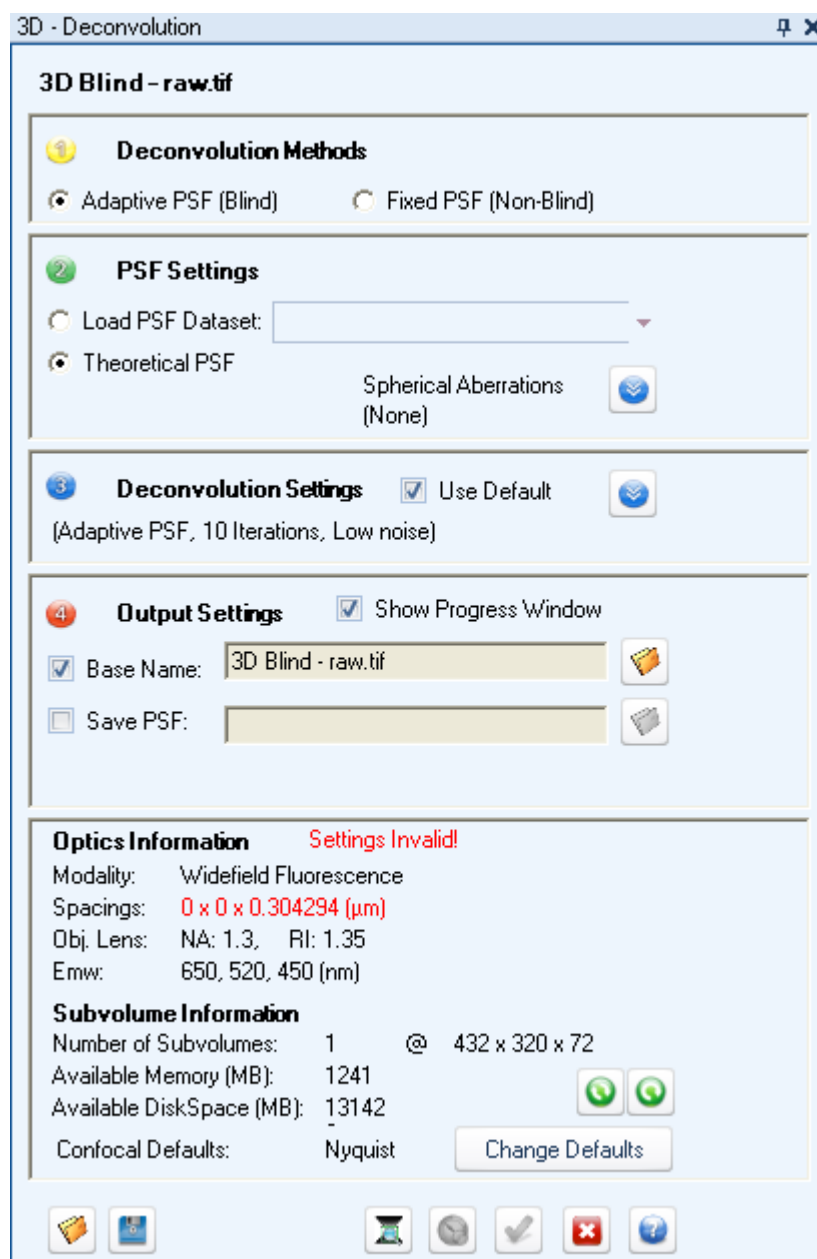
*Note: This section applies to users of **AutoQuant**.*

Blind Deconvolution is considered to be the most accurate algorithm available with **AutoQuant**. Blind Deconvolution does not require the calibration and measurement of the Point Spread Function (PSF). Blind Deconvolution can be used with Fluorescence [Widefield], Laser Scanning Confocal, Brightfield [Transmitted Light], Spinning-Disk Scanning Confocal, or Two-Photon Fluorescence. In addition, structured illumination images from the Optigrid Structured-Light system and the Quorum Angstrom system have been successfully deconvolved using the confocal modality.

3D Blind Deconvolution is an iterative and constrained algorithm. It is iterative in the sense that it repeats the same computational operations many times while converging to the enhanced image solution. It is constrained in the sense that it only accepts deconvolved images that have the correct mathematical properties of nonnegativities, (that is, it does not allow the tracer concentration to have negative values) and smoothness (suppresses snowy-like noise due to low-light levels). 3D Blind Deconvolution is a method of deconvolution that adapts itself to the real PSF of the microscope system (which can be significantly different from the theoretical PSF and from the previously measured PSFs) due to specimen and instrument variations. The **AutoQuant** Blind Deconvolution system is able to adapt to PSF changes within a specimen itself. Thus, the deconvolved results are superior to those methods which utilize theoretical or previously measured PSFs.

For this example you may use your own data type (Fluorescence [Widefield], Laser Scanning Confocal, Brightfield [Transmitted Light], Spinning-Disk Scanning Confocal, or Two-Photon Fluorescence images) or you may follow along with the recommended dataset to use.

- (1) Navigate to the Widefield folder under the Tutorial Data directory and open the dataset `FiteDapi_crop.tif`. This dataset has had the Standard Settings and the Expert Settings set up in the previous tutorial. The information is contained in its header file, and the header file is automatically read when the dataset is loaded.
- (2) From the Deconvolution menu, select 3D Deconvolution. The 3D Deconvolution box will appear. Extensive research and development has made entering or even changing the settings not necessary for most uses. As such, leave the settings at their default setting.



(3) Click the OK icon . The 3D Blind Deconvolution will start. The results will be displayed in the main viewing window.

Close all views before continuing on to the next section.

## Performing 2D Deconvolution

2D Blind Deconvolution allows you to apply either the Adaptive PSF ( Blind) Deconvolution or Fixed PSF (Non-Blind) Deconvolution to a single two-dimensional image (the image can be Fluorescence (widefield), Laser Scanning Confocal, Brightfield (Transmitted Light), Spinning-Disk Scanning Confocal, and Two-Photon Fluorescence) without requiring previous knowledge of the microscope parameters or the image parameters. The 2D Deconvolution algorithm is also capable of improving the resolution of an image for the restoration of features at a sub-pixel resolution level. You may process multi-frame (time series) image sets, individual color channels or intensity images. The 2D algorithm is able to suppress noise while retaining quantitative accuracy (total number of photons) in the image, thus allowing you the ability to make valid quantitative measurements.

- (1) Navigate to the Demo\_Data folder, and open 2D Data.tif.
- (2) From the Deconvolution menu, select 2D Deconvolution. The 2D Deconvolution Dialog will appear.




2D - Deconvolution

### 3D Blind - raw.tif(1)



**1 Deconvolution Methods**  
☐ Adaptive PSF (Blind) ☒ Fixed PSF (Non-Blind)

**2 PSF Settings**  
☐ Load PSF Dataset:   
☒ Theoretical PSF  
☐ Derived PSF








**3 Deconvolution Settings** ☐ Use Default   


**Algorithm Settings**  
Total Iterations:  Save Interval:   
Noise Level:  => Noise Value:   
Performance: ☒ Faster Processing/Reduced Resolution  
☐ Use Gold's Method  
☒ Iteration Acceleration

  
☐ Continue From Previous Result:   
☒ Use Recommended Expert Settings

**4 Output Settings** ☒ Show Progress Window  
☒ Base Name:    
☐ Save PSF:  

**Optics Information** Settings OK  
Modality: Widefield Fluorescence  
Spacings: 1 x 1 x 0.304294 (µm)  
Obj. Lens: NA: 1.3, RI: 1.35  
Emw: 650, 520, 450 (nm)  
**Subvolume Information**  
Number of Subvolumes: 60 @ 240 x 180 x 1  
Available Memory (MB): 1279  
Available DiskSpace (MB): 13126  
Confocal Defaults: Nyquist

- (3) Make sure that Adaptive PSF (Blind) is selected in frame 1 of the dialog.
- (4) Select Theoretical PSF in frame 2 of the dialog.
- (5) In frame 3, set the Total Iterations to 10, and check the Faster Processing/Reduced Resolution box. Ensure that the Use Recommended Expert Settings box is checked.
- (6) In frame 4, leave the Base Name as 2D Data.tif.
- (7) Click the OK button . The 2D Deconvolution will start. The results will be displayed in the main viewing window.

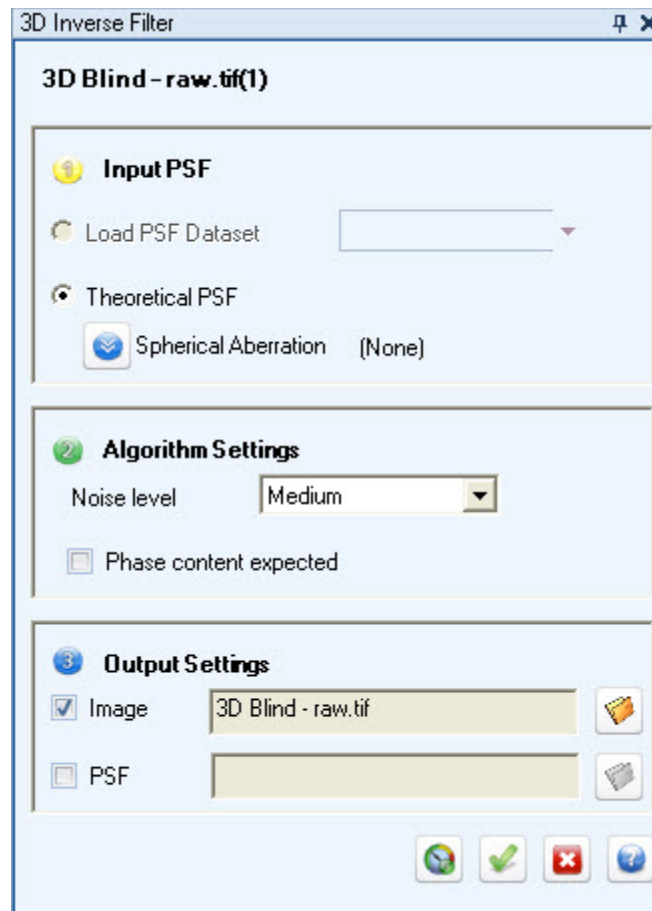
## Using the Inverse Filter


*Use with Widefield Fluorescence and Transmitted Light Brightfield datasets only. Inverse Filter does not work with Confocal datasets.*

The Inverse Filter is a one step non-iterative deconvolution method based on inverse-filtering theory. It utilizes optimal linear filtering.

Inverse Filter is one of the simple deconvolution methods offered in ***AutoQuant***. It is very useful for obtaining quick results, but is not as accurate as Blind Deconvolution. Inverse Filtering is typically more robust than the Nearest Neighbor or No Neighbor deconvolution methods. The execution speed of the Inverse Filter is between that of Nearest Neighbor and Blind Deconvolution. The Inverse Filter should be used in cases where speed is important. A typical processing time is under 2 minutes with a 256x256x32 dataset (Pentium III, 450 Mhz).

- (1) Navigate to the Widefield folder and open Pollen.deb.
- (2) From the Deconvolution menu select 3D Inverse Filter. The Inverse Filter Dialog will appear.



- (3) Select Medium from the Noise Level drop down menu.
- (6) The Phase Content Expected should be used if the specimen is a Transmitted Light Bright-field dataset, and exhibits significant phase characteristics, e.g. areas of the specimen appear brighter than the background. For this dataset leave the Phase Content Expected box unchecked.
- (7) Click the OK icon . **AutoQuant** will now execute the Inverse Filter deconvolution. The status bar will indicate the progression of the Inverse Filter.
- (8) When the Inverse Filter method has finished, a new XY-Max Projection view will appear. This is the result of the Inverse Filter deconvolution. Do not close the Inverse Filter result inv\_Pollen.raw dataset. You will compare it to the results of the Nearest Neighbor in the Nearest Neighbor's section on page 91.

## Using the Nearest Neighbor/No Neighbor Algorithms

*Use with Widefield Fluorescence and Transmitted Light Brightfield datasets only. Confocal datasets do not work with the Nearest Neighbor/No Neighbor algorithms.*

The Nearest Neighbor algorithm is the fastest algorithm available. It works by deconvolving one image slice at a time. As a trade-off in order to achieve this speed, it is less accurate than either the Blind Deconvolution or the Inverse Filter. It should be used in cases where speed is most important. Typical processing times are less than 1 second for a 256x256 single image slice and less than 1 minute for a 256x256x32 3D dataset (Pentium III, 450 Mhz).

**AutoQuant** contains two methods for running the Nearest Neighbor deconvolution on a dataset. One method is called Processing Stack and will run the specified deconvolution on the entire image stack. The other method is called Processing Current Slice and is run while viewing one slice of an XY-Slice Viewer of the image stack.

### To Process the Current Slice

- (1) Click on the Pollen.deb dataset to make it the active view. From the View Menu select Slice Viewer. The slice currently displayed will be the slice processed.
- (2) From the Deconvolution menu select No/Nearest Neighbors. The Nearest/No Neighbor Slice Operation dialog box will appear. The Haze Removal Factor, the Z Kernel Width, and the No Neighbor selections will have values preset for you. The default value for the Haze Removal factor is 0.97, and for the Z Kernel Width it is 3. The No Neighbor algorithm and Phase Content Expected are both unchecked by default.


*Note* - The Phase Content Expected should only be checked if the specimen is a Transmitted Light Brightfield dataset, and exhibits significant phase characteristics, e.g. areas of the specimen appear brighter than the background. For this dataset leave the Phase Content Expected box unchecked.

- (3) Click the *Apply to Current Slice* button. The Nearest Neighbor deconvolution will be applied to the current slice and will be displayed in the current Slice Viewer. To view the original view of the slice click the *Press and Hold to View Original Image* button. Release the button to return to the processed view.

To use the No Neighbor algorithm instead of the Nearest Neighbor algorithm, check the No Neighbor box on the Nearest/No Neighbor Slice Operation box.

- (4) Adjust the results by trying different values for the Haze Removal Factor and the Z Kernel Width, also try the No Neighbor deconvolution by clicking in the flag box to activate and deactivate it.

(5) Set the Haze Removal Factor to 0.98, the Z Kernel Width to 2, and leave the No Neighbor check box unchecked. These values produce satisfactory results for the Nearest Neighbor or No Neighbor algorithm.

(6) Click the Close button  to exit from the Nearest/No Neighbor Slice Operation box. The algorithm chosen (Nearest Neighbor), and the values selected for the Haze Removal Factor and the Z Kernel Width are now saved for this specific dataset.

*Note: Do not close the Pollen.deb dataset. Please continue on to the section.*

## To Process a Stack

(1) Click on the Pollen.deb: XY MAX Projection to make it the active view. From the Deconvolution menu select No/Nearest Neighbor.

(2) The Nearest/No Neighbor Parameters box appears. The settings for the Haze Removal Factor (0.98), the Z Kernel Width (3), and the Nearest Neighbor algorithm were determined above in the Processing Current Slice section.

(3) Click Start Processing Stack. The deconvolution will be performed on each slice of the entire stack. A status bar will indicate the progression. A new dataset window will appear with the results.

(4) Compare the results of the Nearest or No Neighbor deconvolution to the original dataset and note how the deconvolution removes haze and sharpens features. Examine the Maximum projections and the Slice Viewers of the original Pollen.deb dataset, the Inverse Filter, and the Nearest Neighbor results in both the XY and XZ views.

Close all datasets before continuing on to the next section.

## Performing DIC Restoration

*Use with Differential Interference Contrast datasets only.*

The DIC Restoration feature converts a DIC image into an image that represents the optical thickness of the specimen.

### Restoration Method

This section allows you to select between an Iterative and an Inverse Filter process to restore your dataset. The Iterative process will yield better results, whereas the Inverse Filter will yield satisfactory results in less time.

## Explaining the DIC Image Settings

### ***Slice Selection***

In this section, select the slice(s) to deconvolve. Choose from All Slices, Current Slice, or Slice Range. If Slice Range is chosen, you must enter a range into the From: and To: text boxes.

### ***Image Characteristics***

In this section, input the data about your dataset to attain the best restoration results possible.

#### **Shear Angle**

This is the angle at which the light in your acquisition equipment meets your object. Select the proper angle for your equipment, and the shadow box to the right of the drop down menu will mimic the lighting of your dataset.


#### **Image Noise**

Select the image noise level that best matches your dataset, Low, Medium or High. Channel to Deconvolve

Select the channel to restore. If you are restoring a grayscale image, the only available option is Intensity; if you are restoring a multi-channel image, select the channel to restore.

## ***Advanced Settings***

Clicking the Advanced button will open the Expert Settings dialog. Only change these settings if you are experienced with the DIC Restoration feature. Otherwise, use the default settings.

- (1) Navigate to the DIC folder and open test.tif.
- (2) From the Deconvolution menu, select DIC Restoration.
- (3) In the Restoration Method section, ensure that Iterative is selected.
- (4) In the DIC Image Settings, select All Slices (the current image is only 1 slice).
- (5) Still in the DIC Image Settings section, select 135 from the Shear Angle drop down menu. Once the angle is selected, a sample will appear to the right of the drop down menu; the light pattern in this image should match that in the dataset.
- (6) Select Low for Image Noise.
- (7) Click Advanced to view the advanced features. Verify that Total Iterations is set to 15. Verify that the Regularization Iterations and Regularization Contribution are both set to 10. Make sure all of the Processing Options are checked.
- (8) Click the OK icon . The results will display in a new dataset window.

## Expert Settings

*Note: This section applies to users of **AutoQuant** only.*

- (1) Navigate to the Widefield folder and open FitcDapi\_crop.tif.
- (2) Go to the 3D Deconvolution dialog.
- (3) Uncheck “Use Default” to expand the deconvolution settings.
- (4) Uncheck “Use Recommended Expert Settings”.
- (5) Click the “Go to Expert Settings” button. The Expert Settings dialog will appear.

Deconvolution Expert Settings

**BlueGreen\_OpticalCorrection\***

Subvolume

☒ XY Montage      Sub-volume overlap (Pixels): 14

☐ Z Montage      XY-Guardband (Pixels): 14

☒ Dynamic Subvolumes      Z-Guardband (Pixels): 6

Pre-processing

☒ Intensity Correction

☒ Minimum Intensity Removal

Object First Guess: Filtered Original

Spherical Aberration Detection

☒ Accelerated

☐ Automatic

Super Resolution




☐ Activate Sub-pixel Processing      XY factor 1

Fixed PSF Deconvolution

Gold's Method Parameters

Gaussian Width (FWHM in pixels) 1

Smoothing Interval (Iterations) 3



### **Subvolume section**

- (4) This section of Expert Settings affects the handling of the object data.

The default values are as follows:

XY Montage is selected (enabled).  
Z Montage is deselected (disabled).  
Dynamic Subvolumes is selected (enabled).

Leave these fields at their default values.

- (5) Verify that the following fields are set to their default values:

The Subvolume overlap (Pixels) is 10.  
The XY Guardband (Pixels) is 10.  
The Z-Guardband (Pixels) is 6.

### **Pre-processing section**

- (6) Verify that the Intensity Correction box is checked.
- (7) Verify that the Minimum Image Intensity Removal box is checked.
- (8) Verify that the Accelerated SA Detection box is checked.
- (9) Verify that the Pre-Condition Imported PSF box is checked.
- (10) Verify that the Object First Guess is set to Original Data.

### **PSF Section**

*Note: Depending on whether Load PSF Dataset is checked or Theoretical PSF is checked in the main dialog, only one of the following tabs will appear. If Load PSF Dataset is checked, the Fixed PSF Deconvolution tab will be shown, and if Theoretical PSF is checked, then the Adaptive PSF Deconvolution tab will be shown.*

- (11) In the Adaptive PSF Deconvolution tab:
- (12) Verify that the Axial Stretch Factor is 1.
- (13) Verify that the PSF Waist is 1.
- (14) Verify that the PSF First Guess is set to Theoretical Estimate.
- (15) Verify that the Disable PSF Constraints box is unchecked.
- (16) Click on the Fixed PSF Deconvolution tab.

- (17) Verify that the Gaussian Width is set to 1.
- (18) Verify that the Smoothing Interval is set to 3.
- (19) Click OK to close the Expert Settings box. This sets the Expert Settings for the FitcDapi\_crop.tif dataset.
- (20) Close all views before going on to the next tutorial.

## **Expert Settings Explanations**

### **Z Montage**

The Z Montage setting allows the deconvolution application to break the dataset into sections along the optical axis and to deconvolve these subsections separately. The valid settings are: On (checked) or Off (unchecked).

*Guidelines:* The default for this option is “Off”. It should only be turned “On” for image stacks with a large “Depth” setting (e.g. >100 slices). This option reduces the amount of RAM required by the deconvolution process. It may also be useful in rare cases where the sample thickness is so large that the PSF changes dramatically along Z. In such cases, Z Montage allows the blind deconvolution to find different PSF solutions for different depths.

### **XY Montage**

The XY Montage option allows the deconvolution application to break the dataset into sub-volumes along the XY dimensions.

*Guidelines:* The default for this setting is “On”. This option reduces the amount of RAM required by the deconvolution application. It should be turned “Off” only if the deconvolution application is producing rigid, box-like artifacts in your dataset.

### **Z Montage**

The Z Montage option allows the deconvolution application to break the dataset into sub-volumes along the Z dimensions. The default for this option is unchecked.

### **Dynamic Subvolumes**

The Dynamic Subvolumes selection allows the deconvolution application to subdivide the dataset into the largest size the processing computer’s RAM can handle. The advantage of larger subdivisions of data being processed is the increase in processing speed and therefore a decrease in the amount of time it takes to deblur a dataset.

### **Subvolume overlap (Pixels)**

The Subvolume overlap setting determines the number of pixels that the montaged subvolumes will overlap. The possible values are integers from 0 to  $N/2$ , where  $N$  is the width or height of the XY field in pixels, whichever is smaller.

*Guidelines:* An overlap of 10 or 25 pixels usually works best. If the result of the deconvolution contains artifacts having rigid lines, edges or an obvious grid structure start with a value of 10. If doing so reduces the problem, but does not eliminate it, then increase this number again. Overlapping regions are deconvolved twice, so making this number too large (e.g. 100) will increase the deconvolution time.

### ***XY Guardband (Pixels)***

The Guardband size defines the width of a border surrounding each subvolume. This border is the region where the subvolumes are processed to seam them together. This guardband prevents artifacts at the seams. The possible values are integers from 0 to  $N/2$ , where  $N$  is the width or height of the XY field in pixels whichever is smaller.

*Guidelines:* Generally, the larger the guardband, the fewer the artifacts and the better the image quality. However, deconvolution time increases with guardband size, so a default value is set which minimizes deconvolution time and eliminates artifacts in most cases. The default value is 10. This number may be increased if seaming artifacts appear. If so, first increase the number to 15, then 20, and then 25 until the seaming artifact is gone.

### ***Z-Guardband (Pixels)***

The Z-Guardband specifies the number of slices that will be added at the top and bottom of the subvolume. This Guardband prevents artifacts at the seams of these subvolumes. The possible values are integers from 0 to  $N/2$ , where  $N$  is the depth of the XZ or YZ field.

*Guidelines:* The Z-Guardband should never be larger than the subvolume overlap region. A value of 6 is adequate for most image stacks.

### ***Intensity Correction***

Selecting this option will correct for differences in intensities between slices. The default setting for this option is unchecked for Confocal images and checked for Widefield and TLB images. It works much like the Optical Density Correction feature in the Processing menu.

### ***Minimum Intensity Removal***

The Minimum Intensity Removal feature will subtract the minimum intensities from the image. Thus, with Minimum Intensity Removal selected, an image that has intensities ranging from 11-245 will be adjusted to have intensities of 0-234. The default for Minimum Intensity Removal is Yes.

### ***Accelerated SA Detection***

Selecting Accelerated (this is selected by default) will accelerate the process of detecting the spherical aberration of the dataset. Accuracy is slightly better with acceleration not selected, but the trade-off of speed with the acceleration is greater than the accuracy compromise.

### ***Pre-Condition Imported PSF***

This option will pre-condition an imported PSF.

### ***Object First Guess***

The Object First Guess has four options: Flat Sheet, Original Data, Object Input and Filtered Original. The First Guess selects the initial estimate of the object that is used to initiate a blind deconvolution process.

- Flat Sheet: Select this option to use a constant array as the object guess.
- Original Data: Select this option to use a smoothed version of the original image as the object first guess.
- Filtered Original: Select this option to use an Inverse Filter as the object first guess.

- **Object Input:** Select this option to use the previous result from an already performed deconvolution as the object first guess.

### **Super Resolution**

#### **Activate Sub-pixel Processing**

This option will perform deconvolution on a sub-pixel grid to achieve super-resolved results. It will also, however, result in longer processing times.

#### **XY Factor**

If the Activate Sub-pixel Processing feature has been checked, the XY Factor feature will become enabled. This feature allows you to select the number by which the pixels will be divided. The options are 1 (which is for when the Activate Sub-pixel Processing feature is not activated), 2 and 3, with three being the highest resolution, and longest processing time.

### **PSF Section**

In the PSF section there are two tabs: the Adaptive PSF (Blind) and the Non-Blind.

#### **The Adaptive PSF (Blind) tab**

##### **PSF First Guess**

This is the initial PSF estimate. It contains the following PSF Guess methods:

- **Theoretical Estimate:** Selecting Theoretical Estimate causes the deconvolution to generate, as its first PSF Guess, a model of the theoretically-expected spread of light based on the optics settings that have been applied to the dataset. The default Axial Stretch Factor is 1 for widefield data and 3 for confocal data. It is recommended that the **Disable PSF Constraints** box be *unchecked*. Doing this maintains limitations placed on the Point Spread Factor (PSF) to prevent the deconvolution from considering highly unlikely or impossible solutions. The PSF Waist default setting is 1 for both widefield and confocal data. This is the size of the narrowest part of the PSF, usually measured in Airy Disc diameters.

- **Autocorrelation:** This is used by the 2D Deconvolution to generate a PSF by examining the image itself, rather than the optics used to obtain the data. It is not presently applicable to 3D Deconvolution.

- **Flat Sheet:** This option uses as the first guess, a volume filled with a constant value equal to the average of the entire image stack. For very noisy data, this is an appropriate starting point for the deconvolution.

- **PSF Input:** Selecting "PSF Input" allows the user to provide a dataset containing the first guess PSF. This is used anytime a measured PSF is supplied to the non-blind deconvolution. It is also the appropriate selection when using a PSF saved from a previous deconvolution as a starting point for a new deconvolution.

#### **Axial Stretch Factor**

The axial stretch factor is used to stretch a theoretical first guess PSF in the axial direction. Axial stretch factor can be used to correct for geometric distortion caused by severe Spherical Aberration (SA) in cases where SA detection does not work. Axial stretch factor can also be used to reduce noise in undersampled confocal data. Undersampled data is no longer quantitative as there is no way to know the true size of features in the image. Over-deconvolution with a stretched PSF can still be useful in the case, to reduce noise and reveal the presence (though not the true size) of features in the image. This section allows you to set how much axial stretch to allow from the theoretical first guess PSF. The default for Widefield datasets is 1, and for Confocal datasets the default is 3.

### **PSF Waist**

The PSF Waist is the size of the narrowest part of the PSF, usually measured in Airy Disc diameters. The default setting for both Widefield and Confocal is 1.

### **Disable PSF Restraints**

This removes the limitations placed on the Point Spread Function (PSF).

### **The Fixed PSF tab**

This tab contains the parameters to use with Gold's method.

### **Smoothing Interval**

The Smoothing Interval defines the iteration interval at which to apply noise smoothing when using Gold's method. Regular applications of noise smoothing can help to control a noise over-enhancement side effect associated with Gold's method.

### **Gaussian Width (FWHM in pixels)**

This field is based on the Noise Level, found directly below. If the Noise Level is set to *Other*, then this field becomes editable, and you must enter in the Gaussian Width. Otherwise, if you select *Low*, *Medium*, or *High*, then the Gaussian Width automatically defaults to a set number. This feature will be used to smooth the resultant image.

### **Noise Level**

Select the noise level that best represents your dataset. If the Low, Medium, High options are not sufficient, then select *Other* and then enter in your own Gaussian Width in the Gaussian Width section above.

## 2D Blind Deconvolution (Interactive)

This option gives you more control over the 2D Blind deconvolution process:



To use this feature, follow the steps below:

1. Press the Process button.
2. A preview window will appear. The current view in the image display window will be processed. This means that if the current view in the image display is a max projection, then the 2D max projection will be deconvolved. If the view is a slice, the 2D slice will be deconvolved. As the 2D deconvolution proceeds, the preview window will be updated with the results.
3. After the 2D deconvolution is finished, the user can use the Preview Control Iteration scroll bar to move through and examine the results of the 2D process at each iteration.
4. You should set the scroll bar at the iteration that contains the best results. Then press the Apply All button. At this point, the 2D deconvolution will be performed on the entire dataset (not just the current view) using the selected number of iterations.

### ***Notes on Spherical Aberration Detection:***

**Spherical Aberration (SA):** Spherical Aberration occurs when peripheral rays have a different focal length than rays near the center of the lens. Spherical aberration is the dominant aberration in microscopy. Spherical aberration causes asymmetry in the PSF which distorts the data. Detecting spherical aberration before deconvolution will improve the accuracy of the theoretical PSF calculation. The theoretical PSF can be used in both fixed deconvolution and as a first guess to blind deconvolution. Improving the accuracy of the theoretical PSF calculation will in turn improve deconvolution results.

This control is available on both the 3D Deconvolution dialog and the Inverse Filter dialog. The control is hidden by default and can be made visible by pressing the Spherical Aberration drop-down button in the **PSF Settings** group of the 3D Deconvolution dialog or the **Input PSF** group of the 3D Inverse Filter dialog. This control specifies the spherical aberration correction from -15 to +15 (no correction is 0). The values used for SA calculations are “wavelengths”. You may obtain an estimated value using the **Auto Detect** method, the **Manual** method, or the **Calculate SA** method. This will fill in the SA value with the estimate, which then can be used in conjunction with the deconvolution or inverse filter. If you have already performed an SA estimate calculation for the conditions under which the image was collected, you may enter that previously-calculated SA value directly.

**Auto-Detect SA:** This option extracts the SA correction factor from the image data using a patented blind optimization technique and displays the results in the Spherical Aberration control so that you can see if the wavelength values are reasonable. It is recommended that Optical Density Correction be performed before SA detection.

**Manually Detect SA:** When the **Auto Detect/Manual** option is checked, a previous SA estimate calculation for the conditions under which the image was collected should be entered manually using the **Wavelengths of Phase Error** edit box or scroll bar.


**Calculate SA:** This option calculates the spherical aberration based on Obj. Lens immersion RI, Specimen Embedding RI and Distance from Coverslip. It displays the results in the SA control to that you can see if the wavelength values are reasonable.



## Tutorials for Analysis Menu Options

### Counting and Tracking

The Counting and Tracking feature can be used to count objects within a dataset, track those objects if it is a time series dataset, and give statistics on the objects, and if applicable, their movements and changes over time.

- (1) From the File menu, select *Open*. Navigate to the Counting and Tracking folder, click on *Nitella1.tif*. Check the *Use Sequence Detection* box, and then select *Time* from the drop down menu that appears.
- (2) From the Analysis menu, select *Counting and Tracking*.
- (3) Adjust the threshold of the image in the *Counting and Tracking Preview Window* by moving the minimum (red slider) and maximum (orange slider) sliders in the histogram in the Counting and Tracking dialog.
  - The goal is to adjust the thresholding such that the objects to be counted appear as separate white areas.
- (4) Once the dataset is thresholded properly, click the *Count All T* button to count all of the objects in all time points. The section at the bottom of the Counting and Tracking dialog will contain all of the statistical information about each object found, and beneath that grid will be displayed the total number of objects found.
- (5) Click the *Export Objects* button. This will open a new window, with just the segmented objects displayed. This image can be saved and used for publication or presentation purposes.
- (6) Click the *Track* button. The program will track the counted objects through all of the time-points. The statistics from these tracks will be displayed on the *Tracking Grid* and *Tracking Summary Statistics* in the statistics frame.
- (7) Click the *Save*  icon, which will save all of the statistics into an .xls file format so it can be opened and manipulated in a spreadsheet.

### Track Selected Objects

This option allows you to track a small number of selected objects. To select an object, click on it in the Counting and Tracking preview window or highlight it in the Object Stats tab of the grid control in the Counting and Tracking PlugIn Dialog.

You have the option to specify the maximum displacement of the object between time frames. If the object moves farther than the maximum the track will end. You may also specify a maximum deformation (number between 0 and 100 that is the percentage energy difference of the feature vector) of the object between time frames. If the object deforms too much the track will end

## Preprocessing

The Preprocessing menu contains tools for refining a dataset before performing the Counting and Tracking functions on it. Clicking the down arrow next to *Preprocessing* will display the Preprocessing tools.

- Background Equalization - Move the slider to adjust the amount of background equalization to perform on the dataset.
- Noise Smoothing - Move the slider to adjust the amount of noise smoothing to perform on the dataset.
- Wavelet Preprocessing - Move the slider to adjust the amount of wavelet preprocessing to perform on the dataset.

## Histogram


The Histogram is used to segment the dataset for counting. Move the Max (red) and Min (orange) sliders to adjust the intensities displayed. The objective is to adjust the thresholds such that the objects to be counted are filled in with white and disconnected with other objects.

### ***Count All T/Count Current T***

These buttons will activate the Counting function, with *Count All T* counting all of the timepoints, while *Count Active T* will only count the currently displayed timepoint.

## Statistics Frame

At the bottom of the *Counting and Tracking* dialog is the Statistics frame. This frame contains the statistics about all of the objects and tracks found in a dataset once the operations are performed. The statistics can be saved as an .xls file by clicking the *Save* icon. Objects can also be deleted by

highlighting the objects to be removed, then clicking on the trash can icon . Clicking Auto-Quant 5D Viewer icon will open the objects in AutoQuant 5D Viewer.


## Export Objects/Export Segmented Data

These buttons will open a new dataset window, the *Export Objects* button will display the objects, whereas the *Export Segmented Data* button will display the segmented data. These can be saved and used for publication or presentation purposes.

## Statistics

This function provides you with specific statistical information about a dataset. The Statistical Information displayed will be the following: Min. Intensity, Max. Intensity, Total Number of Voxels, Sum of Voxel Intensities, Average Intensity, Standard Deviation, Variance, ROI (Region Of

Interest) Area, ROI Perimeter, and a Histogram of ROI of Current View. You can obtain statistical information about the dataset in any view, except for Montage view and Movie view.

- (1) Open the TobaMC.tif dataset from the Multi-Channel folder. From the View menu, select *Slice Viewer*.
- (2) Under the Analysis menu, select Statistics or click on the Statistics icon. A Statistics dialog box will appear.
- (3) Check the *All Slices* box, which will perform statistics on all slices.
- (4) Left-click on the *Plots Over Slices of Current Time Point*. Notice the dashed red line in the graph.
- (5) In the Slice Viewer, scroll through the slices. Notice that the red dashed line moves to display the values for that slice, and that the values in the *Statistics of XY View* also update automatically to correspond to the new slice. Click on the Histogram button.
- (6) In the dataset window, left-click the ROI icon , then select *Square*. Left-click and drag out a rectangle, then click to complete the rectangle. Right-click in the image to turn the cursor into a *Select Object* cursor. Left-click on the ROI, and notice that the histogram and statistics update automatically.

### **Three-Dimensional Statistics**

*Note - this section applies only to users of 5D Viewer.*

Three-dimensional statistics may be calculated when an image is in a Slice Viewer projection. To calculate three-dimensional statistics, check on the *All Slices* box. **5D Viewer** will then perform the calculations on the selected volume. Selecting Single Slice will cause **5D Viewer** to analyze the displayed slice (two-dimensional) only.

Please close all views before going on to the next section.

### **Line Profile**

This function allows you to obtain line profile information from the dataset by drawing lines on any view.

- (1) Open a view you would like to draw lines on (you can use the TobaMC.tif from the multi-channel folder for an example).
- (2) From the Analysis menu select Line Profile. A Line Profile dialog box will appear.
- (3) Click on the Line Mode, and select *Square*.

(4) Left click the mouse within the view and move the cursor, left clicking again will complete the current line segment. Observe the Line Profile within the Line Profile dialog will show the data properties for the line being drawn. The distance of the line segment is displayed beneath the Line Profile.

*Note: For multi-channel datasets, the Line Profile will display colored lines (red, green, and blue) each of which represents its respective channel color. For example, the red line is the relative intensity values with respect to the Max. and Min. intensity of the image's Red Channel. The orientation of the intensities is relative to the image, not the start and endpoint of the line. When in the Slice Viewer, the Intensity Profile only displays intensities values for a single slice for the line being drawn. At the bottom of the Line Profile box the Angle displays the angle in degrees between the last line drawn and the current line. This works only in the Piecewise Line mode.*

*Note: The distances displayed will be in microns per pixel.*

## Using the FRET Module

*Note: This feature is only available as an added plug-in. Contact your dealer for purchase information.*

One method of studying the interaction between proteins is by analyzing FRET, or Fluorescence Resonance Energy Transfer. When doing so, there are some issues to overcome in order to obtain the most accurate results possible. **Media Cybernetics'** FRET feature allows for the elimination of Cross Talk, so FRET can be accurately calculated.

## Correcting FRET Cross Talk

### Algorithms

In the Algorithm section, there are three algorithms from which to choose. There is Elangovan and Periasamy, Gordon and Herman, and Media Cybernetics Inc., Inc. which is our own proprietary algorithm. For more information on the differences between the algorithms, go to [www.aqi.com](http://www.aqi.com) and download the FRET Application Note from the Support section. Select the desired algorithm by clicking on its radio button.

### **FRET Images and Calibration for Cross Talks**

This section makes associations between the images and excitation/emission/specimen set. To optimize the use of the FRET feature, the proper datasets must first be obtained. Below is a list of headings, and the images they correspond to that will allow for optimal use of the FRET feature (when naming your datasets, make sure you know which one corresponds to which excitation/emission/specimen set):

#### **Headings and Corresponding Datasets**

DDf - Donor excitation and Donor emission applied to the FRET specimen

DAf - Donor excitation and Acceptor emission applied to the FRET specimen

AAf - Acceptor excitation and Acceptor emission applied to the FRET specimen

DDd - Donor excitation and Donor emission applied to the Donor specimen

DAd - Donor excitation and Acceptor emission applied to the Donor specimen

\*AAd - Acceptor excitation and Acceptor emission applied to the Donor specimen

\*DDa - Donor excitation and Donor emission applied to the Acceptor specimen

DAa - Donor excitation and Acceptor emission applied to the Acceptor specimen

AAa - Acceptor excitation and Acceptor emission applied to the Acceptor specimen

*\*Not necessary, but can be used with the Gordon and Herman and Media Cybernetics Inc. Inc. algorithms.*

If it is not possible to obtain the calibration images (DDd, DAd, AAd, DDa, DAa, and AAa) the Cross Talk Coefficients tab can be used to estimate the Cross Talk. See the *Use Cross Talk Coefficients Tutorial* for more information on this.

### **Specified Cross Talk**

#### **Donor Only, Acceptor Only, Donor and Acceptor**

This section allows you to select which datasets to correct for cross talk. Typically, if both donor and acceptor calibration images are available, it is best to select *Donor and Acceptor*. However, if only one or the other set of calibration images are available, for instance, only the donor calibration images, then select that set to correct, in this instance, select *Donor Only*.

#### **Calibration Images**

Checking this button allows you to load in calibration images with which to perform FRET. By default, this button is checked.

#### **Cross Talk Coefficients**

Checking this button allows you to enter cross talk values in lieu of using calibration images. For each image (donor and acceptor) there are three values to be filled in. These should be between 0.0 and 1.0. Ideally, settings should be set between 0.0 and 0.5. Settings of 0.5 and 1.0 are acceptable, but not ideal. The only way to verify is to visually inspect the resultant images. For this reason, unless you know what these coefficients are from previous experiments, it is ideal to have calibration images.

## **PreProcessing**

### **Use Automatic Alignment**

Checking this box will align each set of images using the Channel to Channel Alignment feature, using its default settings. For more information on the Channel to Channel Alignment, see page “crossreference”.

### **Background Subtraction**

This section allows the user to choose from three options regarding the Subtract Background algorithm: *None*, *Minimum\_Value*, and *Histogram\_Peak*. Selecting *None* will perform no background subtraction on the datasets. Selecting *Minimum\_Value* will subtract the background by eliminating the minimum value present in the dataset. Selecting *Histogram\_Peak* will subtract the background based on the most commonly occurring intensity level. There are other options which can be performed on images individually available from the Processing menu, under the Data Correction option. For more information on these, see page “crossreference”. The Subtract Background feature should be used to clean up images that have background noise. Doing so will allow for more accurate results with the FRET algorithms.

### **FRET Conversion Factor G**

This is the relationship between the loss of donor signal due to FRET with the Donor filter set (DD) and the increase in acceptable signal due to FRET with the FRET filter (DA) set. The default is 1.

*Note-This is only available when using the Elangovan and Periasamy or Gordon and Herman algorithm.*

### **Intensity Range Factor**

This divides the intensities into whatever number of sets is selected (e.g., if the number 10 is chosen, then the intensities will be processed in 10 evenly distributed sets). The recommended range is between 5 and 20. This number can either be typed into the text box, or scrolled to using the up and down arrows next to the text box.

*Note-This is only available when using the Elangovan and Periasamy algorithm.*

### **Result Name Prefix**

This text box allows the user to enter a prefix which will be used in the file name of each resultant image created by the experiment.


### **Calculate**

Clicking this button initiates the selected Correct FRET Cross Talk algorithm.


### **Reset**

Clicking this button will clear all file associations made in the *FRET Images* and *Use Calibration Images* sections. There is no undo for this.

### **Close**

This button  closes the FRET dialog box. Any files created using the FRET dialog will remain open, only the dialog box closes.

### **Help**

This button  will launch the Online Help file, opened to the FRET section.

## Calculate FRET Efficiency

Once the Cross Talk has been corrected, the analysis of FRET can begin.

### ***Data for FRET Efficiency Estimation***

This section allows you to select the images to run statistical analysis on. Do this by clicking on the dropdown arrow, then selecting the correct image (based on the heading above the dropdown menu). In other words, for the first (top) dropdown menu, select the image that is the corrected FRET image (DAf). If you have just run the Correct FRET Cross Talk algorithm, the proper images will already be assigned. All three images must be assigned in order to run statistical analysis on them.

### ***Dye Pair's Foster Distance $R_0$***

This section is where you can enter the dye pair used in the images. This will allow for a calculation of the actual Forster's distance when calculating the FRET efficiency. Select the proper dye pair from the dropdown list by clicking on the dropdown arrow, or if the distance is known, yet not displayed in the dropdown menu, it can be entered into the textbox manually by checking the *Use New  $R_0$*  then entering the  $R_0$  into the text box underneath the checkbox.

### ***Fixed Threshold***

This section has two sliders, which allows you to set the upper and lower percentages of the intensities to be analyzed. Set the intensities by either clicking on the pointer and dragging it to the desired location, or by clicking on the desired location to incrementally move the pointer to that spot, or by entering the desired intensity in the text box.

### ***Current Rectangle ROI***

The Image ROI section displays the coordinates of the selected region of interest. Select the region of interest by clicking and dragging on the desired image to create a rectangle around the desired region. *Top Left* refers to the top left corner of the rectangle, whereas *Bottom Right* refers to the bottom right corner of the rectangle. For each, the X, Y and Z refers to the axial coordinates of the respective corners.

### ***Calculate FRET Efficiency in ROI***

This button calculates the efficiencies for a selected region of interest for the active image set.

## **Save Statistics**

This button will save the data into a .csv file, which can then be copied and pasted into an Excel spreadsheet.

## **Delete**

This button removes an already processed region of interest, removes it from the table and erases all of the efficiencies calculated for it. Highlight the desired ROI(s) to be removed, then click *Delete*.

## **Close**

This button will close the FRET Analysis dialogue.

## **Help**

This button will open the Online Help file, opened to the section regarding FRET.

## **Correct FRET Cross Talk Tutorial**

- (1) Open a Windows Explorer and navigate to the FRET folder in the tutorial images. Drag and drop Image-AAa, Image-AAf, Image-DAa, Image-DAd, Image-DAf, Image-DDd, and Image-DDf into the AutoQuant X2 workspace.
- (2) From the Analysis menu, select FRET, then select FRET Cross Talk Correction.
- (3) Select the desired algorithm from the Algorithms section.
- (4) Associate each excitation/emission/specimen with an opened file by clicking on the arrows in the FRET Images section, selecting the file with the Donor excitation and Donor emission applied to the FRET specimen for the DDf heading, the file with the Donor excitation and Acceptor emission applied to the FRET specimen for the DAf heading, and the file with the Acceptor excitation and Acceptor emission applied to the FRET specimen with the AAf heading. *Once a set of files has been assigned and processed, the file associations will be saved in the header file, and only one file association will need to be made in the FRET Image section, the rest will automatically fill in upon clicking anywhere in the dialog.*
- (5) Select *Remove Both* in the *Specified Cross Talk* box.
- (6) In the Calibration for Cross Talks section, associate the files with the appropriate headings as outlined in the preceding list. Create a rectangular region of interest in each one of the calibration sets. Create the region of interest around the area in which FRET is suspected of occurring by clicking and dragging on the image, creating a rectangle around the desired area. *If no calibration images are available, proceed to the Use Cross Talk Coefficients section.*
- (7) Check the *Use Automatic Alignment* box
- (8) From the *Subtract Background* dropdown menu, select *Minimum\_Value*.



- (9) Enter `tutorialset1` for the Result Name Prefix.
- (10) Click the *Calculate* button. FRET will start and three new images will be generated, all with the prefix *tutorialset1*.

## Use Cross Talk Coefficients Tutorial

With the Gordon and Herman as well as the Media Cybernetics Inc., Inc. algorithms, there is an option to enter the Cross Talk Coefficients rather than loading calibration images. It is still highly recommended that calibration images be used, however, the Use Cross Talk Coefficients tab exists for when these images are not available.

- (1) Select *Gordon and Herman* from the Algorithms section.
- (2) Click on the *Cross Talk Coefficients* radio button on the top right side of the Calibration for Cross Talks section.
- (3) Enter the estimated Cross Talk Coefficients for each of the 6 entries on the tab.
- (4) Enter the FRET Conversion Factor  $G$  into the text box. If this value is not known, the default setting of 1 can be used, or an estimation between 0 and 1 can be entered (if using the Gordon and Herman algorithm, a value higher than 1 can be entered).  
 *$G$  is defined as the ratio of acceptor that is excited due to FRET versus the donor that is quenched due to FRET.*
- (5) Click *Calculate*. The application will create 3 new files with the results.
- (6) Changes can be made to the Cross Talk Coefficients settings. Once the changes have been made, to create new files, click the *Calculate* button.

## Calculate FRET Efficiency Tutorial

Once the algorithm has run on the images, and the new files are opened, analyses can be run on the images.

- (1) At the bottom of the dialogs section, click on the Calculate FRET Efficiency tab.
- (2) In the Data for FRET Efficiency Estimation section, if the images are not already loaded in, assign the appropriate images to the Corrected FRET, Corrected Donor and Corrected Acceptor images by clicking the dropdown arrows and selecting the correct image from the list.
- (3) Select a region of interest to process by clicking the ROI icon in the dataset window, selecting the rectangular ROI, then clicking and dragging on one of the images around the area to be analyzed, creating a rectangle around it, then clicking the mouse button again.

(4) Select the dye pair used in the images in the Dye Pair's Foster Distance  $R_0$  section by clicking on the dropdown arrow and selecting the appropriate pair from the list.

*Note - If the dye pair that was used in the images is not listed, but you know the Forster's distance, you can enter that distance in the text box as well. Check the Use New  $R_0$  box, then enter the known  $R_0$  into the text box below the check box.*

(5) Select a percentage range of intensities to analyze by adjusting the Fixed Threshold sliders to the desired percentages. *The minimum must be lower than the maximum.*

(6) Click *Calculate Efficiency in ROI*. The efficiencies will be calculated and displayed in the table in the dialog box. The statistics can be saved as a .csv file by clicking *Save Statistics...* and then specifying a file name and directory to which they will be saved.

(7) Close all images before moving on to the next section.

## Ratiometrics

*Note: This feature is only available as an added plug-in. Contact your dealer for purchase information.*

In order to display and analyze the effects that changes in the sample environment have on the subject, it is necessary to use the *Ratiometrics* feature on the two datasets, each dataset being taken in a different sample environment. The most commonly studied changes studied are changes in calcium concentration and changes in pH.

## Images

In this section, the images to be analyzed are assigned to either the numerator or the denominator. The feature will perform an image division, and as such, the image assigned to the numerator will be divided by the image assigned to the denominator.

## PreProcessing

### ***Use Automatic Alignment***

Checking this box will align the channels using the Channel to Channel Alignment feature, using its default settings. For more information on the Channel to Channel Alignment, see page 85.

### ***Maximum/Minimum Thresholding***

This section has two sliders, which allow you to set the upper and lower percentages of the intensities to be analyzed. Set the intensities by either clicking on the pointer and dragging it to the desired location, or by clicking on the desired location to incrementally move the pointer to that spot, or by entering the desired intensity in the text box.

## Use Grynkiewicz Equation for Ion Concentration

Select this option if you are measuring the effects of changes in calcium concentrations in your sample. To get the best results out of this equation, a calibration set of images should be taken. This set should consist of four images: two each at two separate wavelengths; one with no calcium, and the other saturated with calcium. Once a calibration has been performed, the values can be entered into the textboxes manually in later analyses.

### **Calibration Parameters**

#### **Rmax**

This is the ratio of the two wavelengths for the calcium saturated specimen.

#### **Rmin**

This is the ratio of the two wavelengths for the calcium free specimen.

#### **B(SI2/Sh2)**

This is the ratio between the calcium free and calcium saturated specimens.

#### **Calibrate**

If the above mentioned set of calibration images are open, click on the *Calibrate* button to assign these files to the proper setting.


#### **Numerator Wavelength**

Assign the images associated with the numerator wavelength to this section. The calcium free sample needs to be assigned to the Low Ion image, and the saturated calcium sample needs to be assigned to the High Ion image.


#### **Denominator Wavelength**

Assign the images associated with the denominator wavelength to this section. The calcium free sample needs to be assigned to the Low Ion image, and the saturated calcium sample needs to be assigned to the High Ion image.


#### **Browse**

These buttons  , located to the right of each image textbox, will open a Windows Explorer browser from which you can navigate to, and click and drag files into the AutoQuant/5D Viewer workspace.

#### **OK**

This button  will calibrate the images, to fill in the necessary parameters in order to use the Ratiometrics feature.

#### **Cancel**

Clicking this button  will cancel the calibration process and close the Calibration Images dialog.

#### **Help**

Clicking this button  will open the Online Help file, opened to the topic relating to the Calibration dialog.

**K<sub>d</sub>**

This is the affinity of a fluorescent dye to calcium. Different dyes have a different affinity. Below is a table listing the most popular dyes used in measuring intracellular concentration:

**Table 1: K<sub>d</sub>**

Dye	K <sub>d</sub> for Ca <sup>++</sup>
Fura-2	224 nM
Quin-2	115 nM
Indo-1	250 nM
Ca-Green	190 nM

**Viscosity Factor**

This is the Viscosity factor, which is the effect that the viscosity will have on the spectra of the dyes. Viscosity is defined as the measure of the resistance to flow that a fluid exerts. If the viscosity factor is not known, it is recommended that 0.7-0.85 be entered into this field.

**PostProcessing**

This section allows you to clean up noisy images during processing. This will result in a much more accurate analyses of the datasets.

**Remove Spot Noise**

This option lets the user remove “spot noise” caused by a few bright pixels that may be generated during the ratio process.

**Gaussian Smoothing**

This feature removes noise by using a Gaussian filter, thus creating a more accurate image.

**Calculate**

Clicking this button will run the Ratiometrics process, and create a new image.

**Statistics**

Clicking this button will open a statistics window, displaying a histogram and statistics of the intensities. For more information on the Statistics window, see page “crossreference”.

**Close**

This button  will close the Ratiometrics dialog.

**Help**

This button  will open the Online Help, opened to the Ratiometrics topic.

**Ratiometrics Tutorial**

(1) Navigate to the Ratiometrics folder, then open *HighCa1.001*, *HighCa2.001*, *LowCa1.001*, and *LowCa2.001*. For each file, it will ask which pattern denotes the slice number; select the pattern in which the pound sign (#) falls at the end of the filename.

(2) From the Analysis menu, select Ratiometrics.

- (3) Click on the *Use Grynkiewicz Equation for Ion Concentration* box.
- (4) Click on the *Calibrate* button.
- (5) For the Numerator Wavelength section, assign the *LowCa\_Num.1* and *HighCa\_Num.1* images to the *Low Ion* and *High Ion* images, respectively. For the Denominator Wavelength section, assign the *LowCa\_Den.1* and *HighCa\_Den.1* images to the *Low Ion* and *High Ion* images, respectively. Click the *OK* button. This will populate the Calibration Parameters frame in the Ratiometrics dialog.
- (5) Go to File, then Open, and navigate to the Ratiometrics folder, then the Calibration folder, and open *BILT1\_Num.01.tif* and *BILT2\_Den.01.tif*. For each file, it will ask which pattern denotes the slice number; select the pattern in which the pound sign (#) falls at the end of the filename.
- (6) Assign *BILT1\_Num.01.tif* to the Numerator and assign *BILT2\_Den.01.tif* to the Denominator in the Images frame of the Ratiometrics dialog.
- (7) Verify that the *Remove Spot Noise* feature has been checked. Click *Calculate*. A new image will appear. This is the result of the ratio.
- (8) Click the *Analysis* button in the Ratiometrics dialog to display the statistical information on the new image.
- (9) Close all images before moving on to the next section.

## Colocalization

*Note: This feature is only available as an added plug-in. Contact your dealer for purchase information.*

Often times in multi-channel datasets, two separate dyes will occupy the same area. In order to display the areas in which two dyes become colocalized, use the *Colocalization* feature. This function displays and analyzes the overlay of two separate dyes in a multi-channel dataset. To use Colocalization, a multi-channel dataset must first be opened.

## Image 1/Image 2

In the *Image 1* and *Image 2* sections, the channels to be analyzed are selected and assigned a color in which to be displayed. All available datasets will appear on the dropdown menus, so make sure that the channels selected are the same dimensions. If 2 channels with different dimensions are loaded into these dropdown menus, an error message will appear stating *The dimensions of this image are different from the other image. Do you want to change to this new image for colocaliza-*

*tion analysis*? Selecting *Yes* will load the new image, and clear the other image, at which point a second channel will have to be selected. Selecting *No* will not load the new channel, and the channel loaded for the other image will remain loaded.



Next to each image dropdown menu is a dropdown menu with a color selector. The channel selected in this dropdown menu will determine how the channel is displayed in the Coloc Viewer. The options are Red, Green and Blue. Once a color is selected for one channel, that color will not be available on the dropdown menu for the other channel.

## Colocalization Preview

The Colocalization dialog controls the Colocalization Preview, which appears in the AutoQuant X2 workspace.

### Tools

This section contains tools to manipulate the Colocalization Preview.

-  The zoom controls will zoom the image in and out. The button on the left decreases the image, and the button on the right increases its size. Holding either of these buttons down will continue to either increase or decrease the image's size, depending on which button is being pressed. The dropdown menu displays the image size, relative to its original size as a percentage. Alternatively, it is possible to either select a predetermined percentage from the dropdown menu, or manually enter a percentage into the textbox.
-  The Region of Interest icon allow the user to draw a region of interest in the colocalization image in the dialog.

The image in the Colocalization Preview displays the two selected channels (in the colors selected) and shows the colocalized areas in white.

On the left side of the Colocalization Preview, are the controls to display and move the Coloc Viewer through the slices, if it is being displayed in slice mode.

Checking the *All Slices* checkbox in the Operations section of the Colocalization dialog will display all slices, and the slice scroll bar will disappear. To bring the scroll bar back, uncheck the *All Slices* checkbox.

The scroll bar can be used to move through the slices sequentially by clicking on the pointer and dragging it from side to side.

## 2D Histogram

The 2D Histogram displays the intensity distribution for the two channels. The brighter an area is, the higher the occurrence of that intensity. The left side of the graph represents *Image 1* and the bottom represents *Image 2*. Within the histogram are two lines forming an angle, with boxes at the ends of them, which act as sliders. These lines create an intensity mask. Only the intensities that fall between the two lines will be displayed. The top line controls the displayed intensities for *Image 2* in the Coloc Viewer. The bottom line controls the displayed intensities for *Image 1* in the Coloc Viewer. To move the sliders, place the cursor over the slider (box) and the cursor will turn into a hand. Left click on the slider, then move either up and down, or left to right.

## **Color Map**

The Color Map selection changes the color map of the 2D Histogram. The options available are Hot and Cool.

## **Intensity Range**

The Intensity Range frame displays the intensity ranges of each image. This frame is not editable.

## **Operations**

The operations section contains tools to toggle between slice view and Sum projection, as well as to add a Region of Interest to the Colocalization Statistics.

### **All Slices**

Checking this box will display and calculate colocalization on the Sum projection of the dataset. Unchecking this box will display and calculate colocalization on individual slices as they are displayed. When in slice mode, the slice can be changed via the scroll bar in the Colocalization Preview.

### **Add ROI**

Once an ROI has been created in the Colocalization Preview (see page 53 for more information on creating a region of interest), clicking the *Add ROI* button will add that ROI to the Colocalization Statistics section, with the statistics displayed. Until the *Add ROI* button is clicked, while these statistics are displayed in the *Active ROI* row, they are not saved, and will be lost if any changes are made to the Colocalization Preview, be it a change to the 2D Histogram, changing the displayed slice, or changing the ROI.

## **Colocalization Statistics**

This frame displays the statistics for the Region of Interest in the Colocalization Preview.

Along the top of the ROI frame are the statistics that are gathered on the ROIs.

The first ROI listed is always the Active ROI. If no ROI has been drawn, then the Active ROI will be the entire Coloc Viewer.

## **Delete**

The *Delete* button will delete the selected ROI(s). Multiple ROIs can be selected by either clicking on the first and last ROIs to be deleted while holding the shift button, or clicking and dragging from the first to the last ROI to be deleted.


## **Save Statistics**

Clicking the *Save Statistics* button will save the statistics as a .csv file. A *Save As* browser will appear, in which the user can navigate to the desired directory and name the file. Clicking *Save* will complete the save.


## Generate Coloc Image

Clicking this button will generate the image showing the colocalized areas based on the intensity ranges set up in the 2D Histogram. This new image can be saved as an independent file.


## Close

Clicking this button  will close the Colocalization dialog box.

## Help

Clicking this button  will bring up the Online Help menu, opened to the section on Colocalization.

### **Colocalization Tutorial**

- (1) From the File menu option, select open and navigate to the Multi-Channel folder. Open *colorpollenc.tif*.
- (2) From the Analysis menu at the top of the screen, select Colocalization. The Colocalization dialog will open, as well as the Colocalization Preview window in the workspace.
- (3) For *Image 1*, select *colorpollenc.tif* then select *Channel 1* to the right of that, with *Red* selected as the color next to it, and for *Image 2*, select *colorpollenc.tif* then select *Channel 2* to the right of that, with *Green* selected as the color next to it.
- (4) In the 2D Histogram in the Colocalization Dialog, change the maximum and minimum intensities displayed in the Colocalization Preview by either clicking and dragging the boxes at the bottom and left side of the axes, or by clicking and dragging the line-ends in the histogram area. Notice how the colocalized areas change in the Coloc Viewer.
- (5) Now move the Slice scroll bar back and forth. Notice how the Coloc Viewer changes as the slices change. Now click the *All* checkbox. All slices will now be displayed, as well as the colocalization in all of the slices.
- (6) In the Coloc Viewer, click on the ROI icon  and select *Square*. Inside the image, click then drag out a rectangle and click again to complete it, creating an ROI. Click *Add*. The statistics for the selected ROI will be loaded into the Colocalization Statistics section. Create another ROI and click *Add* to load it into the statistics window.
- (7) Click *Save Statistics....* Name the file *statistics1.txt* and click *Save*.
- (8) Highlight the first ROI by clicking on it in the Colocalization Statistics section, then click the *Delete* button.



- (9) Click *Generate Coloc Image*. A new image will open, displaying the colocalized areas. Open the Slice Viewer from the new image. Scroll through the slices, noticing how the colocalized areas change.
- (10) Close all images before moving on to the next section.

# 5D Viewer Documentation

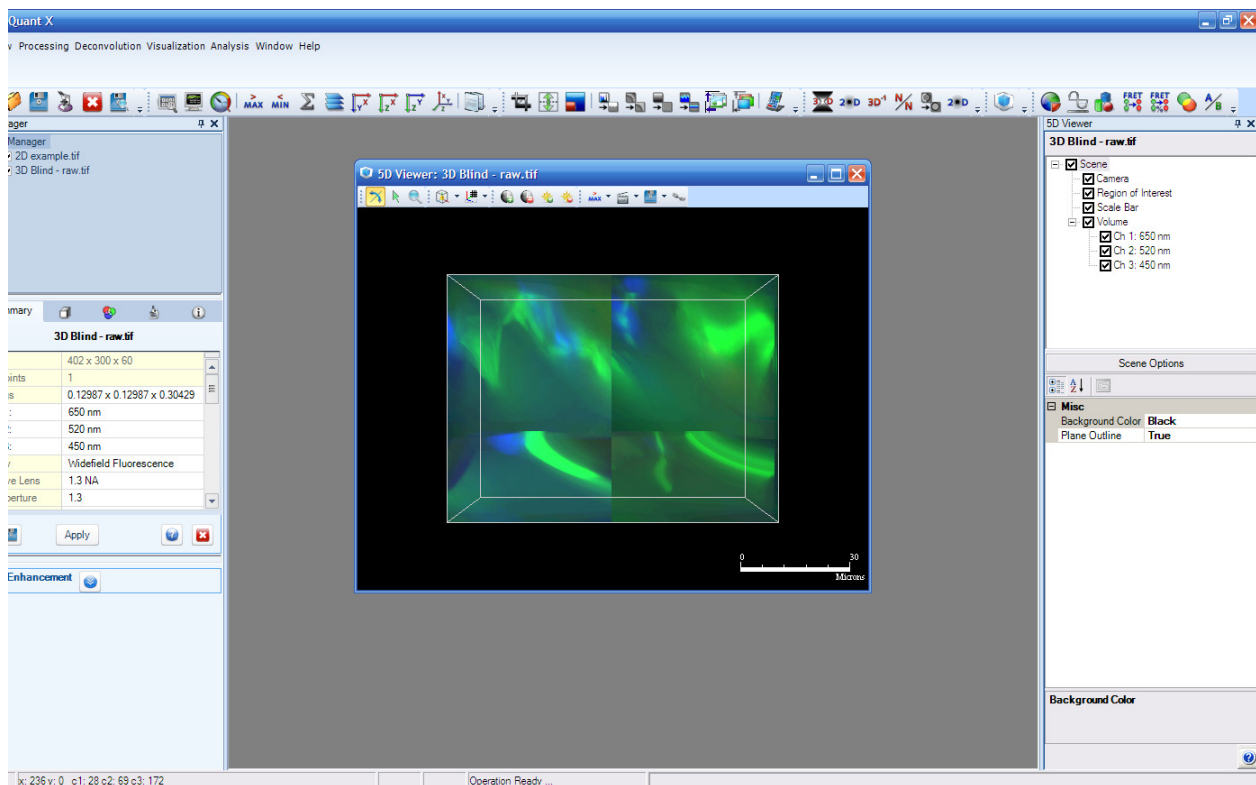
## Introduction

The AutoQuant 5D Viewer 5D Viewer is a rendering tool that allows you to explore your three dimensional, multi-channel and multi-time point data. You can rotate, zoom, color, section, threshold, isosurface and make movies of your data. AutoQuant 5D Viewer can be run as its own application or in combination with the AutoQuant X2 deconvolution software.

## Requirements

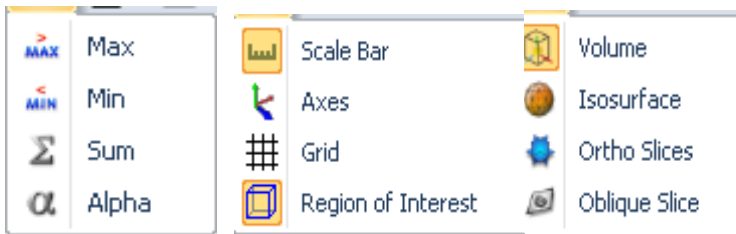
To use AutoQuant 5D Viewer, you will need an *nVidia GeForce 5600* or *nVidia Quadro FX 2000* video card or better with 128MB of texture memory or more. ATI video cards are not supported.

## Interface Overview



The view can be changed using the mouse, keyboard, and tool buttons or by changing parameters in AutoQuant 5D Viewer dialog.

The top portion of AutoQuant 5D Viewer dialog is an object tree containing a list of all objects displayed in the scene or view. Only a few default objects are present when AutoQuant 5D Viewer is started. New objects can be added by using the object tool menus:





Objects can be turned on or off by checking or unchecking their associated box within the object tree.

Volumes, isosurface, ortho slice, and oblique slices are treated as different views of the same object. If one is added to the scene for the first time all other views are turned off in the object tree. They can be manually turned on to display several views at the same time by checking multiple objects in the object tree.

The scale bar, axes, grid and region of interest are treated as decoration objects. They can be toggled on or off from the tool menus or from the object tree.


## Toolbar

AutoQuant 5D Viewer contains a toolbar for some of the more common actions performed on the dataset.

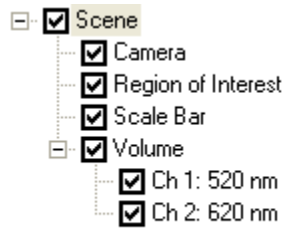
- Rotate  - When the *Rotate* button is depressed, use the left mouse button to click and drag on the object to rotate it along any axis.
- Select  - To select an oblique slice or orthogonal slice, click this icon, then click on the desired slice/plane.

- Projections - To select a projection mode, click on the icon to see the drop-down menu: Maximum Projection, Sum Projection, and Alpha Blend Projection.

- Contrast - Click one of the icons to adjust the brightness and contrast in the Viewer

- Zoom  - When the *Zoom* button is depressed, hold down the left mouse button and move the mouse from left to right and back. Dragging the mouse to the right will zoom out, dragging the mouse to the left will zoom in. Use the right mouse button to click and drag on the object to rotate it along any axis.

If the dataset is more than one channel view objects will have child objects, one for each channel:



Each channel can be turned on or off by using its associated check box.

## Tutorial:

- (1) Load Tutorial\_Data/Widefield/BlueGreen.stk
- (2) Launch AutoQuant 5D Viewer by going to Visualization|5D Viewer
- (3) Display an isosurface as the current view
- (4) In the object tree turn the volume rendering on.
- (5) Turn the volume's child object named "Ch 2: 520 nm" off
- (6) Turn the isosurface child object named "Ch 3: 520 nm" off

This will display the blue channel as a volume rendering and the green channel as an isosurface:



[Insert picture here]

## Time series

When a multi time point dataset is loaded into AutoQuant 5D Viewer the time series scrollbar is shown at the bottom of the display window:



The scroll bar can be used to move between time points one at a time using the scrollbar or the

play forward  and play back  buttons. The time series can be continuously played using forward and back play buttons.

There will be a rendering delay the first time a time point is visited. The volume must be loaded from disk, preprocessed and textures generated. This information is cached for each time point

visited. The next time the time point is displayed it is instantaneously rendered.

## **Object Parameters**

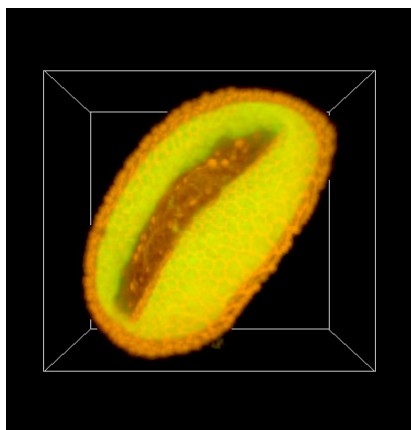
When an object or child object is select in the object tree, the options for that object are displayed in the bottom of AutoQuant 5D Viewer dialog.

If an object containing child objects is selected and its parameters are changed these changes are also applied the child objects.

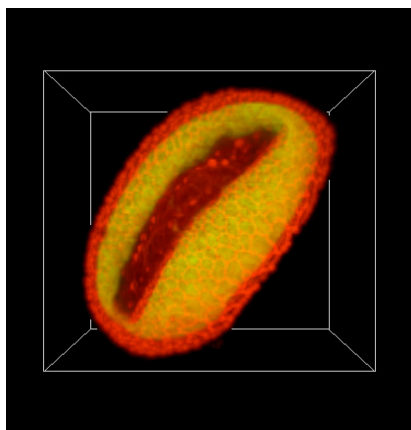
## Tutorial:

- (1) Load Tutorial\_Data/Multi-Channel/Colorpollenc.tif
- (2) Launch AutoQuant 5D Viewer
- (3) Highlight “Volume” in the object tree
- (4) Change the volume’s gamma to 2 and press enter.


This should brighten both channels:




- (5) Highlight “Ch 2: 520 nm” under “Volume”
  - (6) Change the first channel’s gamma back to 1 and press enter.
- This should keep the red channel bright but the green channel dim:




## Navigation

The navigational mode toolbar  allows you to change between rotation, selection and zoom modes.

Rotation is the default mode for AutoQuant 5D Viewer.  The image can be rotated by clicking the left mouse button, dragging the mouse, and releasing the left mouse button.

While in rotation mode right click and drag to zoom in and out. Zoom is set as the current

navigation mode by selecting the zoom  button. Left click and drag while in zoom mode will zoom the image.



The  button enables selection mode. Zoom and rotation are disabled. In selection mode the region of interest, ortho slices, and oblique slices can be moved.

## Keyboard Hotkeys

- Home – Brings the image back to default rotation and zoom.
- Up arrow – rotates the scene up (counter clockwise around the x axis)
- Down arrow – rotates the scene down (clockwise around the x axis)
- Right arrow – rotates the scene right (counter clockwise around the y axis)
- Left arrow – rotates the scene left (clockwise around the y axis)
- PgUp – increases the minimum threshold by 5%.
- PgDn – decrease the minimum threshold by 5%.

## Image Display Controls

The image display controls affect all view objects (volumes, isosurfaces, ortho slices and oblique slices). When these tool buttons are pressed they change the associated parameters of each object. The exact value can be viewed and changed in the parameter list of each object.





The increase/decrease threshold tool buttons   can be used to change the minimum threshold of volume renderings, isosurfaces, ortho slices and oblique slices. This reduces the amount of background so that bright features are easier to see.

If a volume is displayed as a minimum projection the threshold buttons affect the maximum threshold instead.

The increase/decrease brightness tool buttons  can be used to change the brightness of the volume rendering, ortho slices and oblique slices. Use these buttons to brighten up dim data.

## Coloring

Channel coloring is done through the *Summary* and *Coloring* tabs in the **Data Manager**. When the mouse hovers over a channel line on the summary tab a drop down combo box becomes visible:

Volume	402 x 300 x 60
Time Points	1
Spacings	0.13 x 0.13 x 0.304 (µm)
 Ch 1:	 650 nm
 Ch 2:	520 nm
 Ch 3:	450 nm
Modality	Widefield Fluorescence
Objective Lens	1.3 NA
Imm. Medium	1.518

Selecting this combo drop down displays the color picker window. Here you can choose from a standard palette of colors or a wavelength. Different colors can be assigned to each channel of a multi channel dataset. For single channel images a third color tab containing look-up tables is available. This is available for single channel datasets only.

Changing the channel color will affect the coloring of volume renderings, isosurfaces, orthogonal slices and oblique slices.

## Volume Rendering

By default a volume rendering is displayed when AutoQuant 5D Viewer is first opened and a “volume” object is added to the object tree.

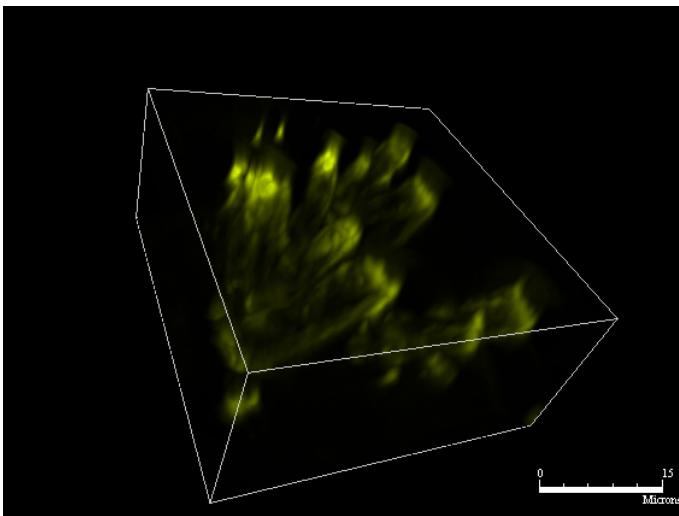
Some preprocessing is done to the data prior to displaying it the first time as a volume. This includes rescaling of intensity ranges to fill the full dynamic range of the monitor, resizing it in X and Y to speed up rendering, and converting it to 8bit so the graphics card can display the volume. Though some contrast and XY resolution are lost this allows the image to be rotated in real time.

- Bright – Change the overall brightness of the volume rendering.
- Gamma – Changes how intensities are mapped to screen colors from a linear mapping to a curve. A gamma value of one will create a one to one intensity to color display of

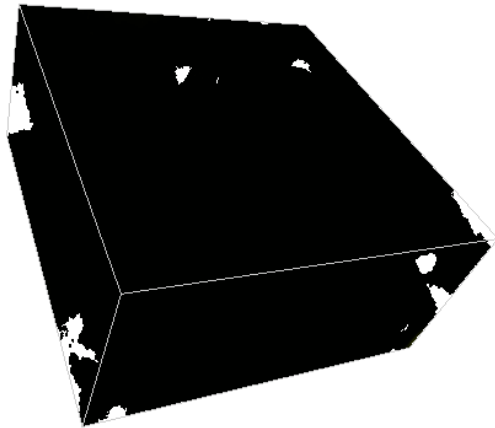


the image. A gamma larger than one will emphasize dim features. A gamma smaller than one will suppress dim features.

- **Min/Max Threshold** – Used to include or exclude certain intensity ranges. By increasing the minimum threshold background is removed. By decreasing the maximum threshold bright features are removed.
- **Number of slices** – Changes how accurate the volume rendering is. Most volumes can be over one hundred slices thick in any direction. Instead of blending all slices together, a subset of slices are blended together. The more slices the better the rendering but the slower the performance. If the number of slices control parameter is set to “Manual” or “All” changing this value will have no affect on the image.
- **Number of slices Control** – Allows you to manually set the number of slices to render the volume with, use all the slices or have the software do it automatically (recommended).
- **Projection** - Affects how volume slices are combined or blended.
- **Maximum projection** – The brightest portions of the volume are used to render the volume. This is the default rendering style for volumes. It is the recommended projection for fluorescent images (bright features on a dark background).

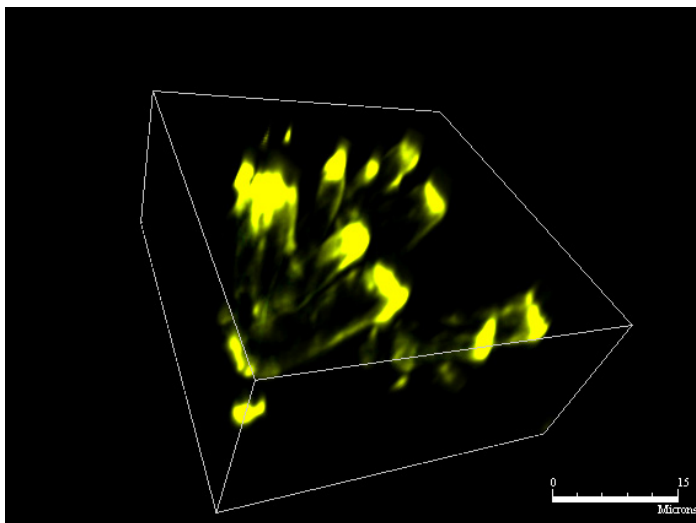


- **Minimum projection** – The parts of the image with the lowest intensities are displayed. This is the opposite of the maximum projection. This is recommended for viewing brightfield data (dark features on a bright background).

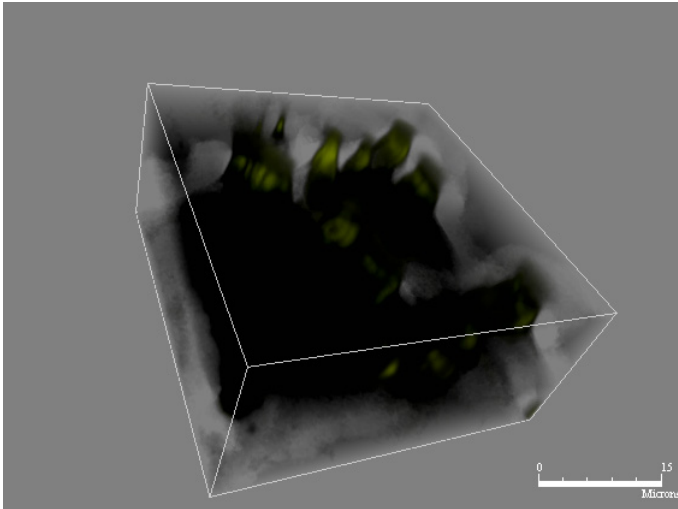


When a minimum projection is selected the background is automatically set to white. It can be set back to black or another color by selecting Scene in the object tree and changing its Background color parameter.

- Sum projection – Each slice is added together to give a composite rendering of the entire volume. This makes it easier to see trends in the data but many times background is summed and makes it hard to see features.



- Alpha (blend) projection – Each slice in the image is blended together from back to front. Details far away from the point of view impact the final rendering the least. Details close to the point of view influence the final rendering the most. This gives a true to life rendering of the image



When an alpha projection is selected the background is automatically set to grey. It can be set back to black or another color by selecting Scene in the object tree and changing its Background color parameter.

The projections can be changed from the projection tool menu as well:



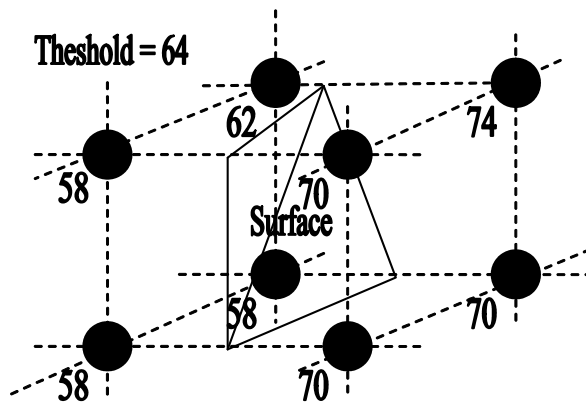
The projection tool menu and the volume object's projection parameter are linked.

**Storage Hint** – Render the volume as a 3D or 2D texture. 3D is the default and recommended mode.

**Volume size** – Change the subsampled volume size. If the original dataset's width or height are larger than this value then it is resized such that the larger of the width and height are equal to this value.

## Isosurface

An isosurface is a surface that represents a constant value within the dataset. The constant value, or threshold, is automatically selected for each channel by a background removal algorithm. Each pair of voxels are examined incrementally throughout the volume. If the voxel values transition from below the threshold to above the threshold one or more triangles are created between the voxels.




The location of the surface is weighted by the voxel values. If the voxel with an intensity of 62 was increased to 64 the top back corner of the surface would be moved to coincide with the location of the voxel.

In the previous example if the selected threshold was decreased from 64 to 60 the entire surface would move to the left. If the threshold was increased to 68 or more the surface would move to the right. As more voxel pairs are visited the isosurface is gradually built. If the threshold is changed the geometry of the isosurface is affected. This in turn will affect volumetric measurements such as surface area and 3D volume.

- Bin factor – Allows the isosurface to be generated from every 2<sup>nd</sup>, 3<sup>rd</sup>, 4<sup>th</sup>, etc. voxel. This limits the complexity of the isosurface. For smaller bin factor values more triangles are generated and the surface appears more detailed. Bin factors of one or two are not recommended for large datasets because there may not be enough memory to hold all the triangles generated. A large number of triangles also can slow down the rendering time and make rotation and interaction with the viewer cumbersome.
- Clip – If true and the oblique slice is present the oblique slice will cut the isosurface. This is useful in showing a cut away view of the isosurface.
- Display sides – Renders an isosurface on the sides of the volume.
- Max threshold – Picks the upper value at which the isosurface will be generated. For fluorescent data this usually remains at the maximum intensity of the dataset and no isosurface is generated using this threshold. For brightfield data decrease this value to remove background and dim feature.
- Min Threshold – Picks the lower value at which the isosurface will be generated. For fluorescent data increasing this will remove background and dim features.

- Use Smoothing – Applies a 3x3 two dimensional Gaussian smoothing filter to the data before generating an isosurface. This removes noise and yields smoother surfaces. This is the recommended default.
- Wire Frame – When true shows the triangles which make up the isosurface.

## Ortho slices


If an ortho slice is added to the scene an “Ortho Group” is added to the object tree with three child ortho slices, one along each of the major axes. Each ortho slice can be slid along its respective X, Y or Z axes. They can be moved by entering selection mode () and then clicking and dragging the slice.

Slice Number – Allows you to manually set the location of the slice along its axis.

The ortho slice objects can be animated by selecting Set Start Frame from the movie tool menu, moving the slice and then selecting Set End Frame from the movie tool menu. When the movie is played the slice is animated between the two locations.

## Oblique slices

The oblique slice object displays a slice on an arbitrary axis.

To move the oblique slice enter selection mode () toolbutton), select the slice by clicking on it and then click somewhere in the middle of the slice and drag.

To rotate the oblique slice enter selection mode, select the slice and then click and drag on the outer edge or crosshairs.

Click outside the oblique slice to hide the oblique slice rotation and drag controls.


If an isosurface or volume rendering is displayed at the same time the oblique slice will by default clip, or cut, the isosurface. Clipping can be turned on or off for each channel of the volume or isosurface. When clipping is turned on for one channel and off for another you can “peel away” a channel by moving the oblique slice.

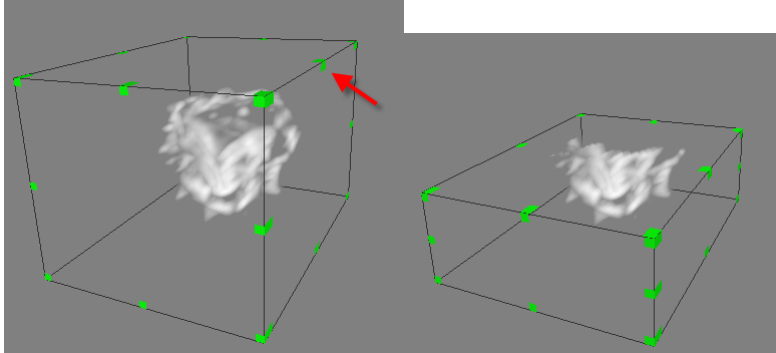
Distance from Origin – The location of the oblique slice along its axis. Increase or decrease this value to manually set the location of the oblique slice

Normal X, Y, and Z – Controls the orientation of the axis on which the oblique slice sits.

The oblique slice can be animated by selecting Set Start Frame from the movie tool menu, moving the slice and then selecting Set End Frame from the movie tool menu. When the movie is played the slice is animated between the two locations.

## Region of interest

By default the entire extents of the object are displayed. The object can be subvolumed by entering selection mode (  toolbutton) and then clicking and dragging any of the green control boxes at the edges of the volume:



The exact locations of the sides can be set within the parameters section of the Region of Interest object.

The region of interest can not be animated.

## Scalebar

Scalebar for the image appears at the lower right-hand side of the image window. The scale bar can be turned on or off from the object tree or decoration menu toolbar.

The scale bar responds to spacing changes in the data manager. If no spacing information is available in the data manger then pixel values are displayed.

## Axis

Displays the X-axis as red, the Y-axis as green and the Z-axis as blue.

## Grid

The grid is displayed on the back sides of the image's bounding box. The grid is linked to the scale bar's markings and will automatically resize when zooming.

- Show labels – Turns on or off the “X”, “Y” and “Z” text.
- Show Spacing – Turns on or off the numbers along each axis.

## Camera

Controls how the scene is being viewed.

- Complexity – When rotating the number of slices blended for the volume and the number of isosurface triangles are decreased for faster rendering. A complexity of one represents full complexity which gives best image quality but slow performance. A value close to 0 will minimize the complexity, show poor image quality but give best performance.
- Scale – Allows the zoom factor for the entire scene to be manually set.
- Orientation X, Y and Z – Describe the scene’s orientation vector. Change this to manually orient the scene.
- Angle – Amount of rotation along the scene’s orientation vector. Set in degrees.
- Perspective – Toggles between perspective and orthogonal viewing modes.
- Perspective Angle – Changes the amount of perspective. This parameter has no effect if perspective is turned off.

## Scene

Holder for generic scene parameters.

- Background color – Changes the background color. This is automatically set to grey when an alpha projection is selected, white when a min projection is selected and black when maximum or sum projection are selected.
- Plane Outline – Turn on or off the bounding lines of the region of interest and ortho slices.



## Movie Making



Quick movies that rotate around a major access can be made by selecting the movie tool. Then select Quick Movies, select an access and then select an amount of rotation. The movie is automatically played. A movie navigation bar is displayed at the bottom of the screen and a new movie scene object is added to the object tree.

More elaborate movies can be made by selecting a zoom and rotation, select Set Start Frame from the movie menu, select a new zoom and rotation and then select Set End Frame from the movie menu. This allows for customer zooms and rotations.

Frames per second – Controls the speed at which the movie is played. If set too large and the graphics card can not render quick enough the movie will be played as fast as possible without skipping frames.

To save a movie select Save Movie from the movie toolbar. Wait until the movie has completed an entire cycle. A new RGB multi time point dataset is created on the AutoQuant X desktop and an Save As dialog is opened allows you to save the movie as an AVI.

Rock – Play forward and back through the rotation.

Total Frames – Controls the number of steps between the start and end frames of the movie.

Ortho slices and oblique slices can be animated in Start/End frame movies. Move one of these slices to its start position and select Set Start Frame from the movie menu. Move the slice to another location and select Set End Frame from the movie menu. The slice will be animated between the start and end positions during the movie. This can be done in combination with zoom and rotation.

## Windows

### Cascade

This feature places all the open dataset views in overlapping and offset positions from top to bottom of the viewing window.

### Tile Vertically

This function stacks vertically the open datasets in the window. Closed or minimized windows will not be tiled (displayed). To end the tiling, click on each dataset and select the desired size from the zoom box.

### Tile Horizontally

This function stacks horizontally the open datasets in the window. Closed or minimized windows will not be tiled (displayed). To end the tiling, click on each dataset and select the desired size from the zoom box.

### Close All Images

This function allows the closing of all open views, whether active or not, and all docked (datasets are minimized with only part of their title bar showing) views all at once.

### Close All Modules

This function will close all open modules in the application. Any dialogs that are open will be closed at this time.

### Show Dock Windows

This function will display all docked windows in the application: Data Browser, Dialogs, and Operation Status. If any of these is not actually opened, it will not be displayed. For instance, if the Data Browser had been closed, it will not reopen when *Show Dock Windows* is selected.

### Hide Dock Windows

This function will dock all displayed windows in the application: Data Browser, Dialogs, and Operation Status.

## **Window List**

This feature lists all current views on the screen.

## Help

### AutoQuant/5D Viewer Help

This feature provides a help window opened to the table of contents.

### Check for Updates

Media Cybernetics' products incorporate an AutoUpdate feature. It is configured to check for updates once a month if your machine is connected to the internet. If you would like to check for an update before that, select Check for Update from the Help menu. You will be prompted to click OK to search the internet for updates. If you do not have a LAN connection, but are using some other internet connection, you will be prompted to click Configure to configure your internet settings. Any internet connections available on your machine will be listed in the Use Dial Up Connection drop down menu. Select the service you wish to use, then click OK. This will take you back to the original dialog box, at which point you can click OK again. AutoUpdate will now search for any available updates.

### About

This feature shows the version and copyright information about the product.



## Frequently Asked Questions

**Q: How does *AutoQuant* perform an accurate deconvolution without a measured point-spread function (PSF)?**

A: *AutoQuant*'s adaptive blind deconvolution automatically extracts point-spread functions from the raw image dataset. It does this by analyzing the spatial frequencies that make up the raw image volume and creating one (or more) PSF's that are based on this information. Because the performance of a microscope varies across the field of view and (especially) along the optical axis, *AutoQuant* will often automatically derive several PSFs to process a single image volume. This assures that results are accurate, even though the optical system (scope, cover slip, immersion medium, mounting medium, specimen) is not linear. Blindly restored PSFs are also more accurate than empirically measured PSFs because they are free from noise contamination, and are based on the distortion actually present within the dataset, rather than when the PSF was measured.

**Q: Can I process 2D images and time-lapse movies?**

A: Yes. *AutoQuant* contains an iterative blind deconvolution for use on 2D images. It is for researchers who do not have a Z-motor, or, who wish to improve resolving power of their standard 2D images, or, who want to improve resolving power in time-lapse movies.

**Q: Can I process many images as a batch?**

A: *AutoQuant* contains functions that permit you to set up a batch of images that will then be sequentially processed. After setting up the parameters for your deconvolution, you may choose to place the image into a queue that you launch at a later time. This is useful for laboratories who do a lot of imaging and for researchers performing time-lapsed observations of 3D volumes. These laboratories can set up their images to process a batch of images overnight, and have them all ready for analysis in the morning.

**Q: How can I easily inspect my images before and after deconvolution?**

A: Not only does *AutoQuant* contain five different types of deconvolution (adaptive 3D blind, nearest-neighbors, no-neighbors, inverse filtering and 2D deconvolution), but it also provides many different ways to visualize the raw and processed images. *AutoQuant* enables a 3D dataset to be visualized using several different projections and orientations. Each volume can be viewed from XY, XZ and ZY orientations as a maximum, minimum, summed, or voxel gradient projection, or as individual optical slices. No other package currently on the market combines these powerful processing tools with an environment for visualizing the results.

**Q: Can I process images that were acquired with a confocal microscope?**

A: Yes. Confocal microscopes (conventional, two-photon and spinning-disk) are effective at reducing the scatter from deep tissues. In practice, the z resolution is not good compared to the x, y resolution. Resolution along the z-axis is typically 0.5 to 3 microns. This means that structures

are elongated along the z-axis. Applying *AutoQuant* to a confocal image stack improves the z-resolution dramatically - often achieving beyond the theoretical 0.5 micron limit. The signal-to-noise ratio of the deconvolved images is greatly improved, and poor signal-to-noise is otherwise common with confocal microscopes, especially in live cell imaging and where photobleaching is a problem.

**Q: What types of image files are supported by *AutoQuant*?**

A: *AutoQuant* supports most common image formats, including 8, 12, 16 bit TIFF's, Universal Imaging's STK, BioRad's PIC, Leica's native files, BMP's, AVI's, Olympus Fluoview, raw 8, 12, 16 integer, 32 bit floating point data, and others.

**Q: Can *AutoQuant* handle the saturated portions of my image volumes?**

A: Yes. Not only can *AutoQuant* handle images that contain saturated areas, it can often recover information from within these saturated sections.

This is possible in situations where there are non-saturated areas nearby the saturated areas. The blur in the nearby non-saturated region contains information about the saturated region which *AutoQuant* restores. The degree to which this is effective depends on the size of the saturated area and the numerical aperture of the lens.

Of course, according to good laboratory practice it is always best to avoid saturation, but this property of the algorithm makes data collection and experiment design much easier since the avoidance of saturation is now a flexible requirement and no longer an absolute requirement.

**Q: Are noisy, poor signal-to-noise images going to be problem for *AutoQuant*?**

A: *AutoQuant* is able to process extremely noisy images and deliver restored image sets with very little noise. Noisy images are common with a confocal microscope, an intensified camera or whenever low light is used to minimize bleaching.

**Q: How many iterations should I apply? How do I know I have performed sufficient iterations? What happens if I perform too many iterations?**

A: Start with the rule of thumb: With the Use Recommended Expert Settings box checked, for widefield and brightfield use 10 iterations, for confocal, two-photon, and spinning disk, use 10 iterations.

If by inspecting the images you are satisfied with the resolving power, stop. If you want still better resolving power, increase or double the numbers. Repeat this increasing until you are satisfied with the resolving power. You will know you went too far if you notice obvious noise and artifacts, which will be clear because structures will be shown which clearly cannot be present and which do not have remnants seen in the raw images. If you notice such noise and artifacts, cut the number of iterations in half. With some experience, 3 trials should be sufficient to determine the number of iterations necessary for your experimental setup and then all similar samples, having identical conditions (sample type, lens, magnification, z-spacing), should use the same number of iterations. These 3 trials can be done in a few minutes by cropping a small ~64x64 section and experimenting with that section before deconvolving the whole volume.

**Q: How do I know the restored features are real?**

A: By looking at the raw and deconvolved images side-by-side, compare to see if details are visible in both. A structure, if it is real, will appear in the raw image, although it will be more difficult to see due to having low contrast, noise and blur.

**Q: When do artifacts appear?**

A: Artifacts such as “ringing” and “mottling” appear if the wrong parameters have been entered, or if there is excessive noise. Dead or hot pixels can also cause artifacts. Using the Data Correction utilities can eliminate these problems.

**Q: How close together should my optical slices be?**

A: The optimal setting has them spaced equal to the depth of field, for both widefield and confocal. This rule of thumb is flexible if other experimental constraints warrant a compromise.

**Q: What should I do after I load my dataset?**

A: Next, go to Deconvolution – Deconvolution Settings - Standard Settings and fill in any entries that have not been previously entered, including the type of deconvolution you want to perform. Then go to Deconvolution - Start 3D Deconvolution to begin your deconvolution.

**Q: How are saturated pixels handled?**

A: Saturated pixels are handled by a proprietary algorithm which, heuristically, treats them as unknown data points. A special algorithm uses the blur information surrounding the saturated pixels to infer the information that should have otherwise been provided by the saturated pixel.

**Q: When should I use optical density or attenuation correction?**

A: *Use optical density correction* if you notice a flicker in your optical sections. You will notice this flicker by creating a XZ View (View menu --> Single View --> XZ) and a Sum Projection (Visualization menu --> Sum Projection). If it is present, the flicker will be obvious, so when in doubt you may assume there is no flicker. Flicker looks like dark horizontal lines through your dataset. *Use attenuation correction* if you notice that the overall brightness of your picture becomes progressively dimmer as you penetrate deeper into the sample. Your dataset should be in a XZ View and the need for correction will be indicated by having a bright top and dark bottom.

**Q: When do I check “Phase content expected”?**

A: This is mainly used for brightfield samples, which have not been stained and are viewed by closing down the condenser aperture on the microscope. This can create for certain types of transmitted light brightfield samples a phase character. These types of samples cause a refraction of light passing through them, rather than a simple absorption of light. This type of sample is recog-

nized when (1) it is a transmitted light brightfield dataset, and (2) you see regions in the image which are brighter than the background surrounding the sample. By checking “Phase Content Expected” fine detail in the specimen will be preserved.

**Q: How do I deconvolve my dataset faster?**

A: First, try the Inverse Filter under the Deconvolution menu. This usually provides results in a few minutes. It only works with widefield and not with confocal. Secondly, experiment with a cropped ~64x64 section before deconvolving the whole volume. Also set the Deconvolution Performance to Best Medium Fast. Finally, make sure that you have maximized the amount of free memory on your computer.

**Q: How is blind deconvolution different from nearest neighbors and inverse filter? When should I use each algorithm?**

A: *The Blind Deconvolution* is an iterative-constrained algorithm that in most cases (not all) provides the best resolving power, including the resolving power along the z axis. *The Inverse Filter* is a fast linear method that gives remarkable deconvolutions with widefield dataset. It does not work with confocal dataset. With some types of samples, especially those that have fine low contrast staining, it gives even better results than the blind deconvolution. Use the Inverse Filter if speed is the most important consideration or if the sample has fine low contrast detail that is important to see. *The Nearest Neighbor* is the fastest deconvolution. Its main advantage is speed. It provides images typically within seconds. It is a linear method. Use the Nearest Neighbor algorithm for quick feedback or for enhancing pictures for presentations, but be careful about interpretation of the images because it is not as robust against artifacts and noise as are the Blind Deconvolution and Inverse Filter.

**Q: Why do all the features in my deconvolved dataset look darker than before?**

A: The deconvolution will increase the dynamic range of the dataset as a result of removing the blur. When an image is displayed on-screen, it is auto-scaled between the maximum and minimum intensities. Therefore, the background features will appear darker on-screen. Use the brightness and gamma controls to better observe the features in the dataset.

**Q: Why won't my images load into Photoshop?**

A: This is because you may have saved the images in a format that is not compatible with Photoshop. Save the images as TIFF and when asked by the program if you want to “export as multiple files,” answer “Yes.”



## Appendices

### Table of Z Spacings (Z Step Sizes)

**Table 1:**

Numerical Aperture (NA)	Step Size for Oil $\eta = 1.515$	Step Size for Water $\eta = 1.333$	Step Size for Air $\eta = 1.00$
0.2	18.83	16.61	12.37
0.3	8.3	7.33	5.43
0.4	4.6	4.08	2.99
0.5	2.95	2.57	1.87
0.6	2.02	1.76	1.25
0.7	1.458	1.256	0.875
0.8	1.094	0.935	0.625
0.9	0.824	0.713	0.443
1.0	0.663	0.552	0.250
1.1	0.528	0.429	Undefined
1.2	0.424	0.330	Undefined
1.3	0.339	0.242	Undefined
1.4	0.267	Undefined	Undefined
1.5	0.192	Undefined	Undefined

DOF = Depth of Field

Wavelength ( $\lambda$ ) = 0.50 $\mu$ m

$$\Delta Z_{\text{slice}} = \text{DOF} = \frac{\lambda}{4\eta \sin^2(0.5 \sin^{-1}[\text{NA}/\eta])}$$

## Table of Parameters

**Table 2:**

File Name	Numerical Aperture	Refractive Index	Modality	X, Y, and Z Dimensions (Pixels)			X, Y, and Z Spacing (Microns)		
FitcDapi_crop.tif	1.4	1.515	Widefield	90	120	50	0.07	0.07	0.15
FitcDap_crop2d.tif	1.4	1.515	Widefield	90	120	1	0.07	0.07	0.15
Blue_Green.stk	1.4	1.515	Widefield	90	120	50	0.07	0.07	0.15
FitcDapi.decon.tif	1.4	1.515	Widefield	90	120	50	0.07	0.07	0.15
Malaria_Red.tif	1.3	1.518	Widefield	202	243	60	0.1299	0.1299	0.3043
Malaria_Green.tif	1.3	1.518	Widefield	202	243	60	0.1299	0.1299	0.3043
Malaria_Blue.tif	1.3	1.518	Widefield	202	243	60	0.1299	0.1299	0.3043
Pollen.deb	0.7	1.515	Widefield	90	80	110	0.357	0.3636	0.4
Colensc.tif	1.4	1.5	Widefield	256	256	1	2D dataset		
Brain.avz	--	--	MRI	256	256	64	--	--	--
Hip.avz	--	--	CT scan	512	512	90	0.9	0.9	4.8
SmallHip.avz	--	--	CT scan	128	128	120	3.6	3.6	3.6
RotatedHip.avz	--	--	CT scan	128	128	120	3.6	3.6	3.6
Vicera.avz	--	--	CT scan	128	128	120	3.6	3.6	3.6
Skinless.avz	--	--	CT scan	128	128	120	3.6	3.6	3.6
HipMovie.avm	Movie		CT scan	128	128	120	3.6	3.6	3.6
VoxelTLB.avm	Movie		Brightfield	130	130	41	0.279	0.279	1.0
TLBview	1.3	1.515	Brightfield	198	208	64	0.653	0.0653	0.4
Star.0	0.25	1.0	Brightfield	494	348	46	0.6897	0.6897	8.0
Star.lt0	0.25	1.0	Brightfield	494	348	1	0.6897	0.6897	8.0
Star.bs1	0.25	1.0	Brightfield	494	348	1	0.6897	0.6897	8.0
Star.bs2	0.25	1.0	Brightfield	494	348	1	0.6897	0.6897	8.0
Starfish_crop2.deb	0.25	1.0	Brightfield	128	128	54	0.6897	0.6897	8.0
TobaMC.tif	1.4	1.515	Confocal	128	128	32	0.23	0.23	0.4
Neuron.deb	0.8	1.515	Confocal	384	512	64	0.94	0.94	1.0
Neuron_crop1.deb	0.8	1.515	Confocal	128	128	64	0.94	0.94	1.0
RedNuc	1.3	1.33	Confocal	128	512	64	0.07	0.07	0.15

**Table 2:**

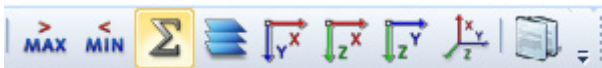
File Name	Numerical Aperture	Refractive Index	Modality	X, Y, and Z Dimensions (Pixels)			X, Y, and Z Spacing (Microns)		
Time Series Data	1.32	1.4502	Widefield	256	256	23	.32	.32	.4783
Test.tif				256	256	1			
Test_parameter.set			A file containing datasets parameters						
Triple label segment(seg			A file containing datasets labels						
Volume & Surface Area.text			A text file of the measurements for Surface Area & Volume of a dataset.						

## Toolbar Icons

### File Toolbar Icons



### Visualization Toolbar Icons



### View Toolbar Icons



## Processing Toolbar Icons



## Analysis Toolbar Icons



## DeconvolutionToolbar Icons



## Zoom Toolbar Icon



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