

The Olympus logo is displayed in white, bold, uppercase letters. A thin orange horizontal line is positioned directly beneath the text. The background of the entire page is a light blue gradient with abstract white curved lines and a pattern of semi-transparent blue circles in the lower half.

OLYMPUS

SIMPLIFIED OPERATION MANUAL

FLUOVIEW FV10i

Note

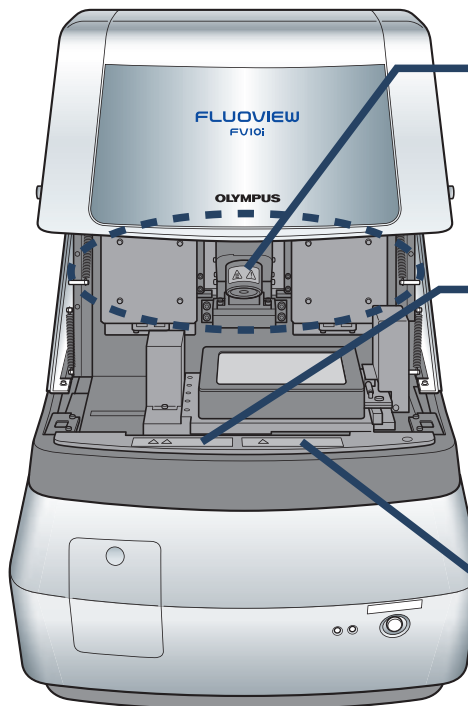
This manual introduces a simplified operating procedure of the product so that even the person who uses it for the first time can complete the operations.

For detailed operating procedures, refer to the separate User's Manuals [Safety] and [Hardware] as well as the Navigation/Online Help of the software.

Retain this manual in an easily accessible place near the work desk for future reference.



During observation and acquisition, the cover is kept closed to avoid the risk on the user. However, when replacing the specimen holder, the cover should be opened so care is required in this operation. Also note that the laser beam is screened by a shutter and therefore safe while the cover is opened.



Caution against high temperature

(FV10i-W only)

This section becomes hot for keeping the incubator temperature.



Caution against finger being caught

This section accommodates motorized modules and the unit should not be operated while the cover is open. Also be careful not to have your hand or finger caught by the cover when closing it.

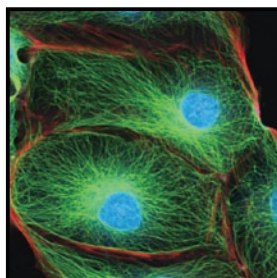


Caution for specimen replacement

Always remove the specimen together with the specimen holder.

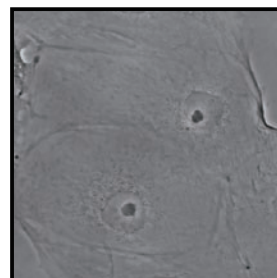
Observation Methods

Fluorescence observation



This method utilizes the fluorescence generated when the specimen is irradiated with strong light to observe the specimen structure or identify the substance according to the presence/absence and color tone of the fluorescence.

Phase contrast observation



This method makes it possible to observe a non-stained specimen (living specimen) or nearly transparent specimen, which is hard to observe in brightfield observation, by enhancing the contrast between brightness and darkness.

FLUOVIEW FV10i

Incubator temperature indicator (FV10i-W only)

The indicator blinks during warm-up and light steadily when the optimum temperature (34 to 38° C) is reached.
The temperature indicator shows “- -” when the heater is OFF.

Cover

Cover lock buttons
(Provided on both sides)

Push and hold the bottoms on the left and right sides to open the cover.

FV10i-W main unit
(FV10C-W2)

FV10i-O main unit
(FV10C-O2)

STATUS LEDs

When both the main and sub switches are ON:

LEDs		Laser	Software
Left	Right		
 Blinking	 Lighting	Initializing	
 Lighting	 Lighting	Standby	Not run
 Lighting	 Blinking		Running

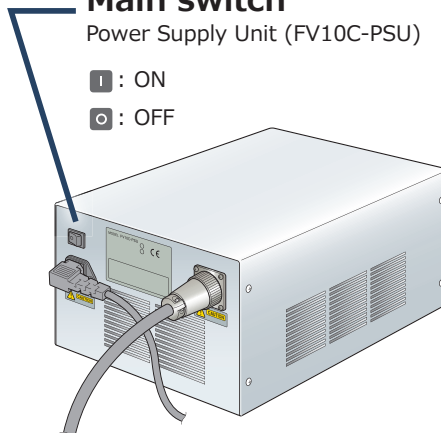
Both the left LED () and right LED () blink when the cover is open.

Sub switch

Main switch

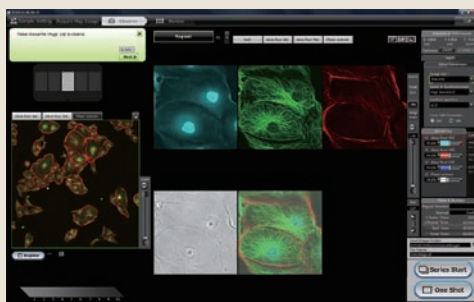
Power Supply Unit (FV10C-PSU)

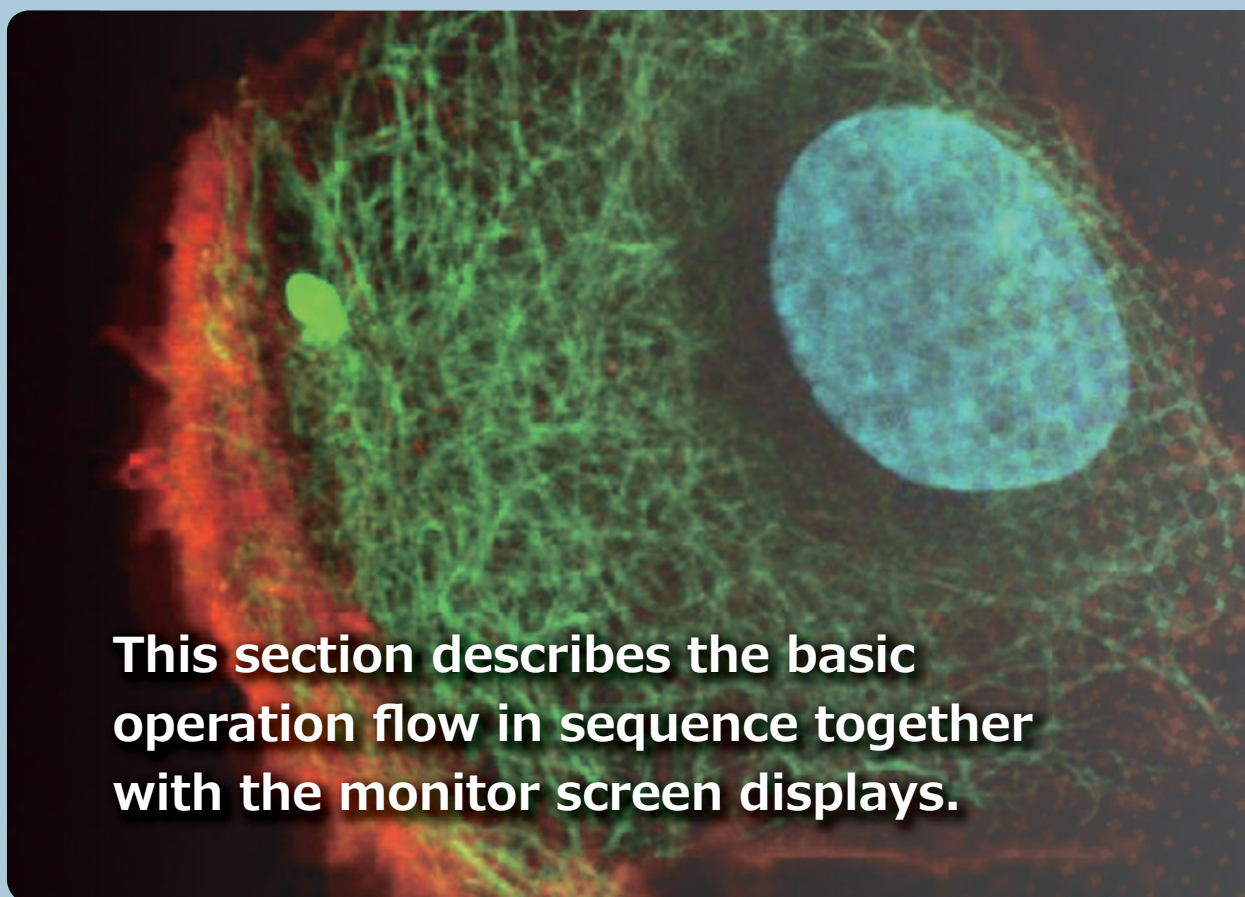
☒ : ON
☐ : OFF



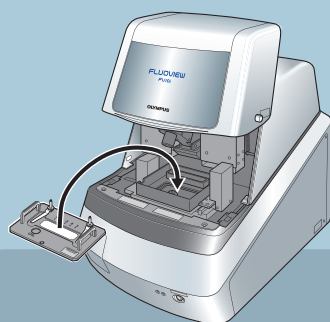
The operations other than those available with the above controls can be adjusted or set from the software. (The operations are easy using the Navigation.)

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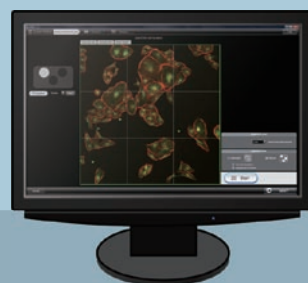
This section describes the basic operation flow in sequence together with the monitor screen displays.



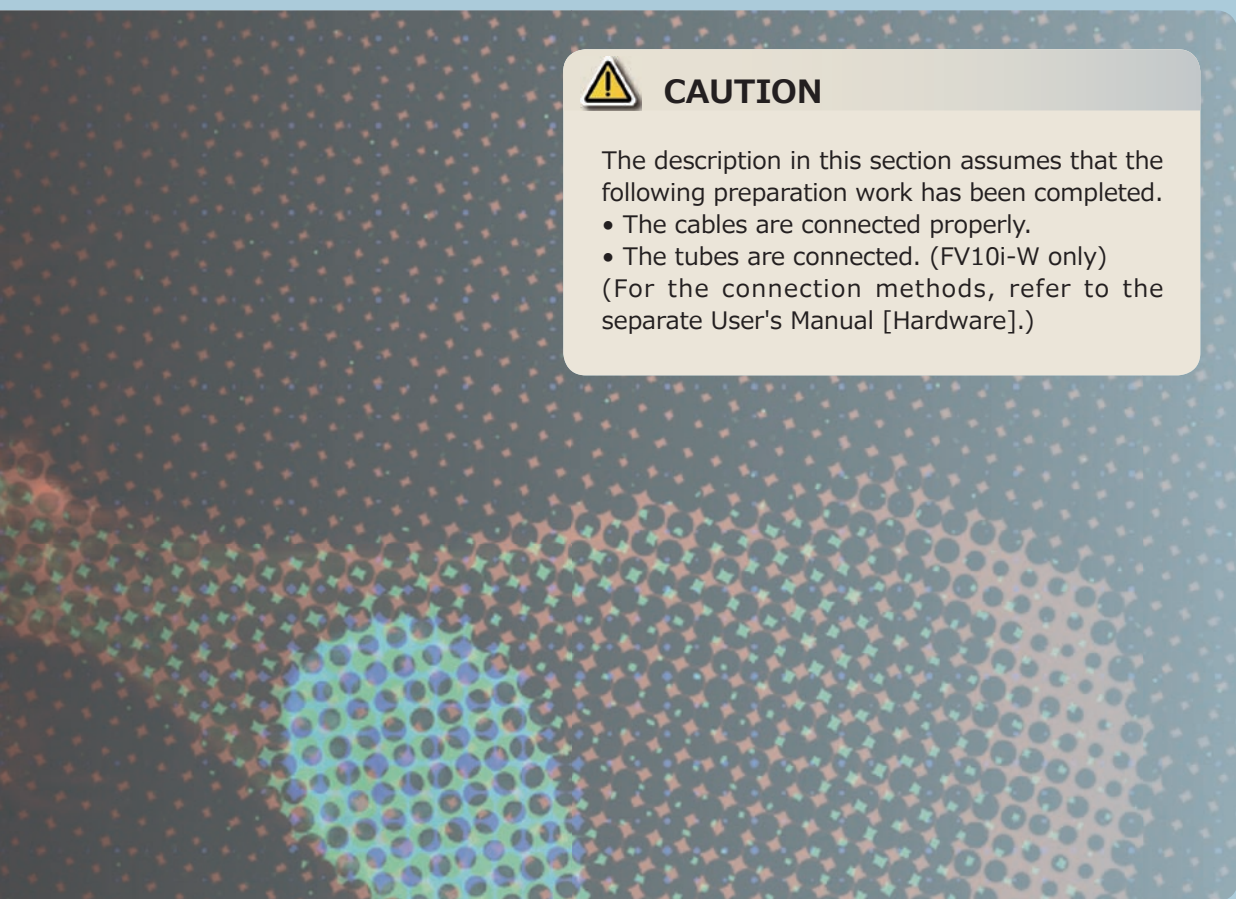
→ p. 6



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CAUTION

The description in this section assumes that the following preparation work has been completed.

- The cables are connected properly.
- The tubes are connected. (FV10i-W only)

(For the connection methods, refer to the separate User's Manual [Hardware].)

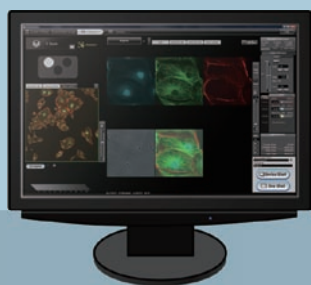


Image Acquisition

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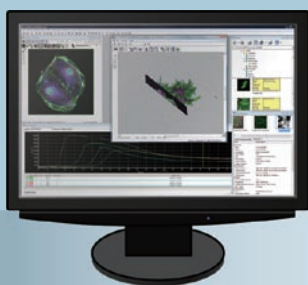
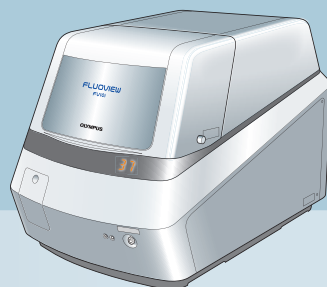


Image Editing/ Analysis

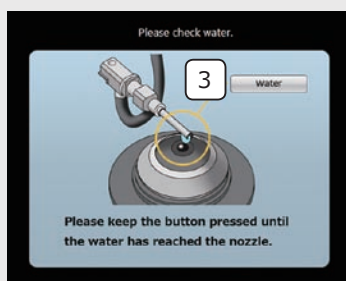
→ p. 13

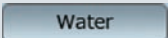
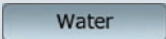


System Termination

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Checking water in the water feed nozzle (FV10i-W only)



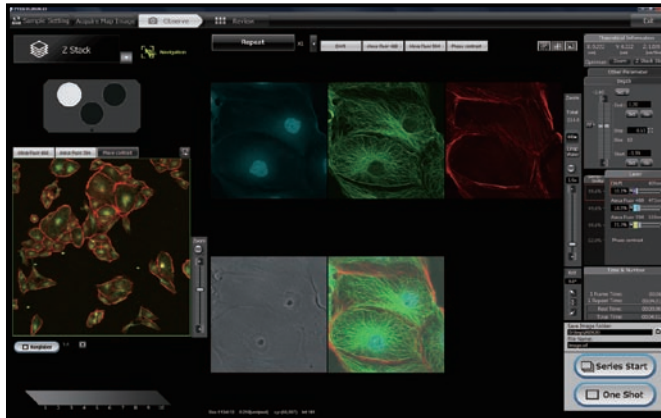
- 1 Open the cover.
- 2 Remove the specimen holder.
- 3 Press and hold the  button. (It may sometimes take as long as 30 seconds to 1 minute until water comes out.)
- 4 When water comes out of the water feed nozzle, release the  button.
- 5 Install the specimen holder in the original position.
- 6 Close the cover.



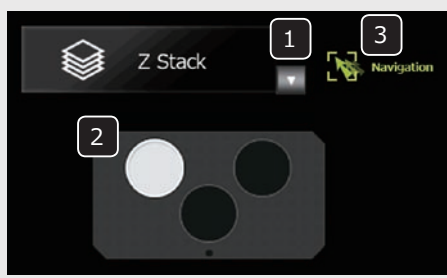
- From the [Setup Dyes] list, select the fluorescent dyes for use in observation and click on the  button. (Up to 4 dyes can be selected.)
- Click on the  button.

Now the software setup is complete.

4. Image Acquisition



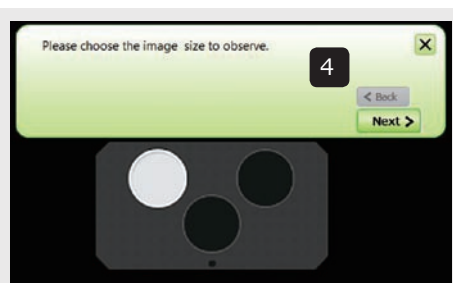
Selecting the acquisition mode



- 1 Click on the ▼ button to select the acquisition mode.
- 2 Double-click on the observation target position (container) that you want to acquire the image.
- 3 Click on the  **Navigation** button to view the Navigation.

The following five acquisition modes are available.

-  **Time Lapse:** Sequential acquisition of the images of a specified area at the specified time intervals.
-  **Z Stack:** Sequential acquisition of the images of a specified area by varying the focused position at constant steps.
-  **Z Stack Time Lapse:** Sequential acquisition combining the Z Stack and Time Lapse modes.
-  **Multi Area Time Lapse:** Time Lapse acquisition while moving across multiple specified areas.
-  **Multi Area Z Stack Time Lapse:** Z Stack/Time Lapse combined acquisition while moving across multiple specified areas.

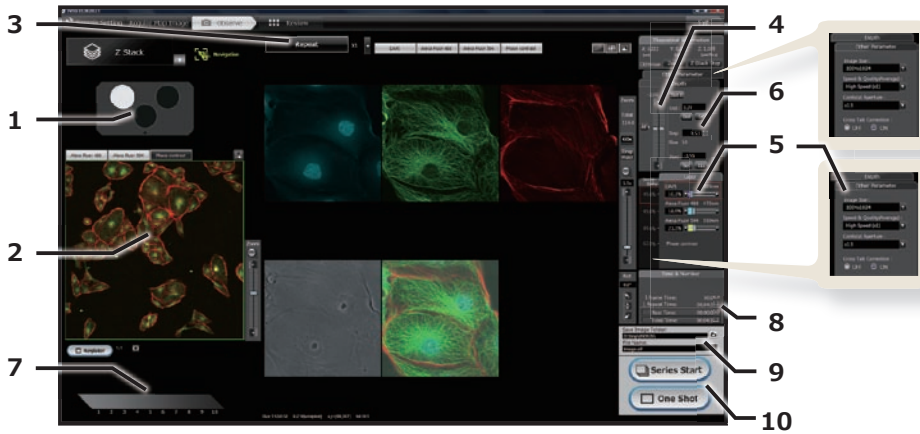


- 4 Hereafter, continue operation by following the instructions given by the Navigation.

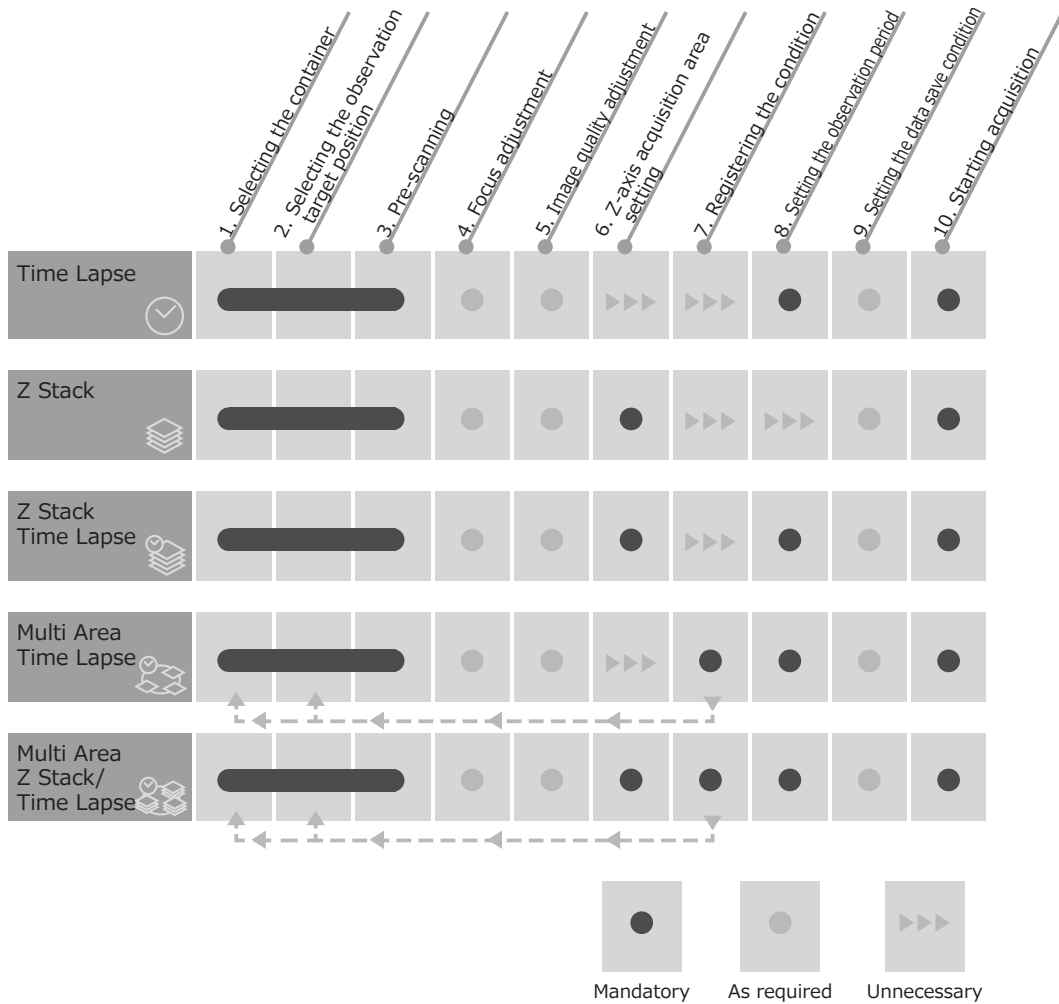
When the operation is completed, click on the  **Next >** button to view the next operated position and the instruction on it.

5. Flow of Image Acquisition

Introduction

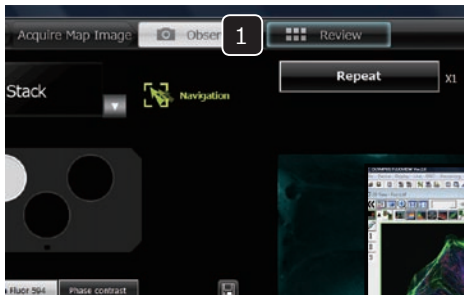


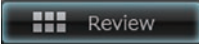
Main Controls



6. Image Editing/Analysis

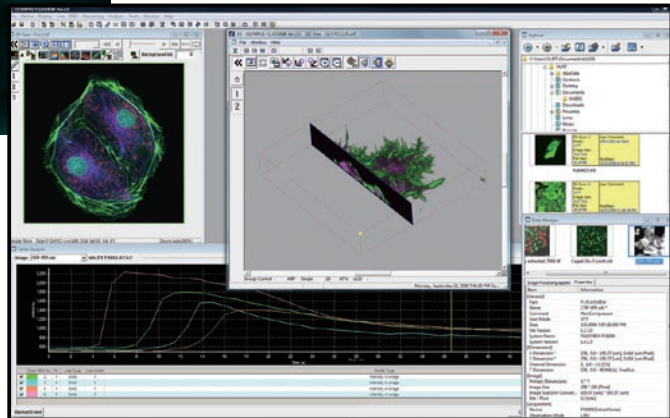
Switching to the Review Software



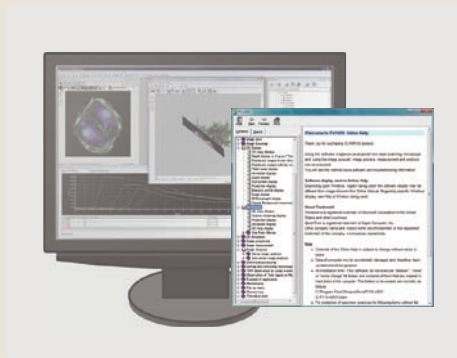
- 1 Click on the  button on the toolbar to switch to the Review software.

The Review software provides the following functions.

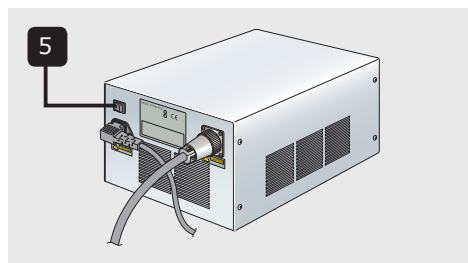
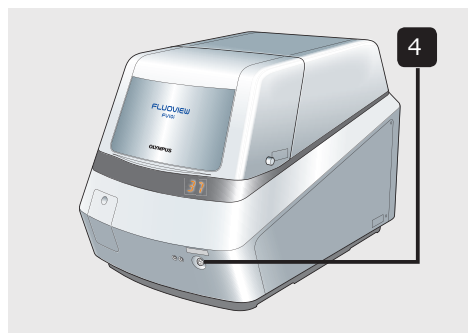
- LUT modification
- Tile image display
- Maximum Intensity Projection image creation
- Annotation addition
- 3D image building
- Time lapse image file editing using Partial Open
- Stitch image creation using MATLAB Viewer
- File export
- Addition of user ID



The software also incorporates various other functions. For details, open the [Help] menu and launch the Online Help.



The screenshot shows the ImageJ software interface. A red circle with the number '1' highlights the 'Exit' button in the top right corner. The 'Theoretical Information' panel on the right displays coordinates (X: 0.222, Y: 0.222, Z: 1.020) and units (µm, µm, µm/step). The 'Depth' panel shows a scale from -2.60 to 1.20 µm. The main image area displays a 3D volume rendering of a sample, with a 'Zoom' slider set to 60x. The 'Drop' button is visible at the bottom of the image area.



- 1 Click on the  button on the toolbar to exit the software.
- 2 Shut down Windows Vista.
- 3 Set the main switch of the display to OFF.
- 4 Press and hold the sub switch on the main unit for 3 seconds to turn it off.
- 5 Set the main switch on the power supply unit to "○" (OFF) as required.

MEMO

Introduction

Main Controls

Observation/Acquisition Flow

