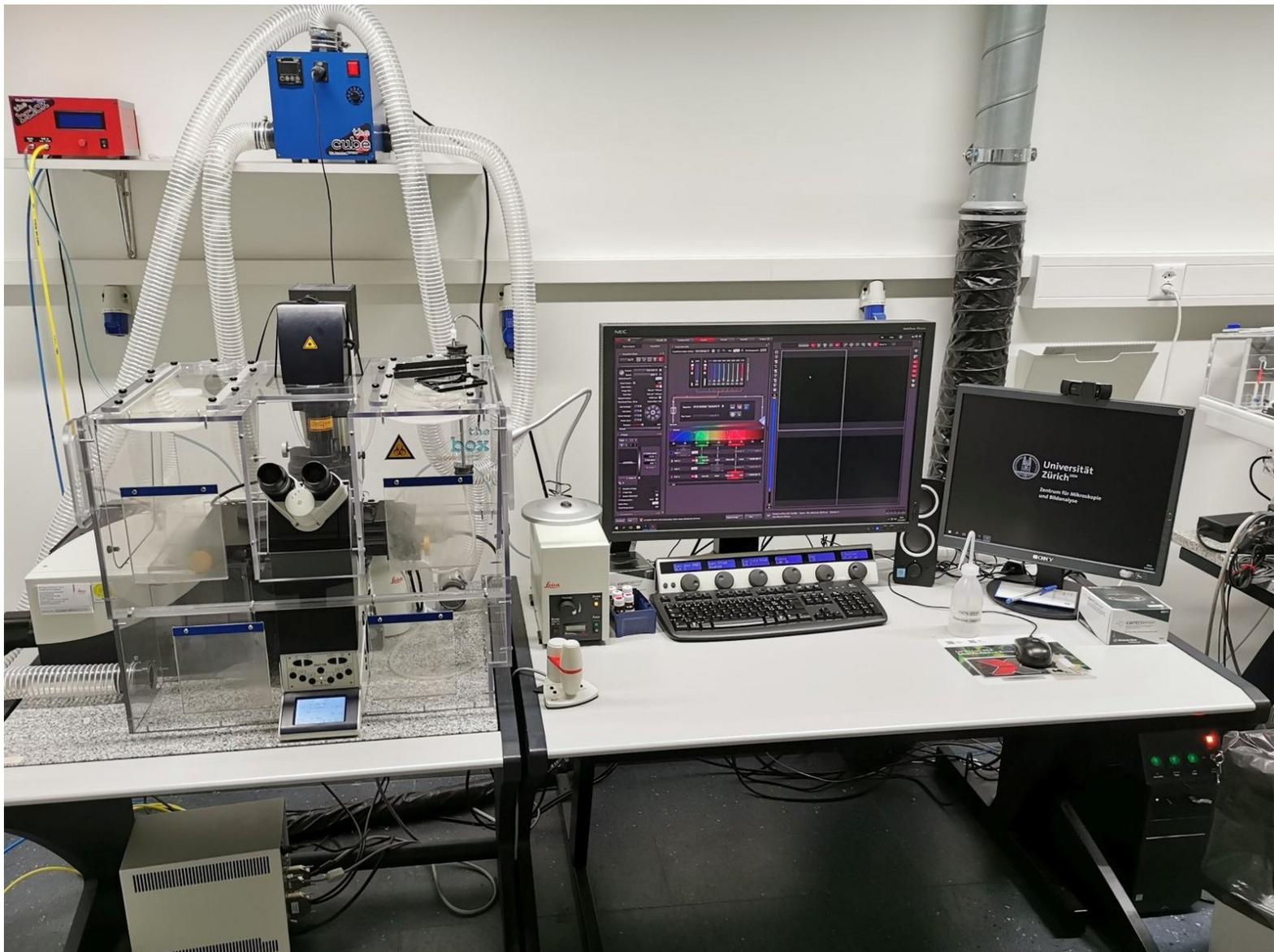


Leica SP8 inverse (Schlieren) - Part 4: Data saving and instrument shut down

How to leave the Leica SP8 confocal laser scanning microscope located at the Schlieren campus, room WAD-12-K-105 after the recording is finished.

Written By: z mbstaff

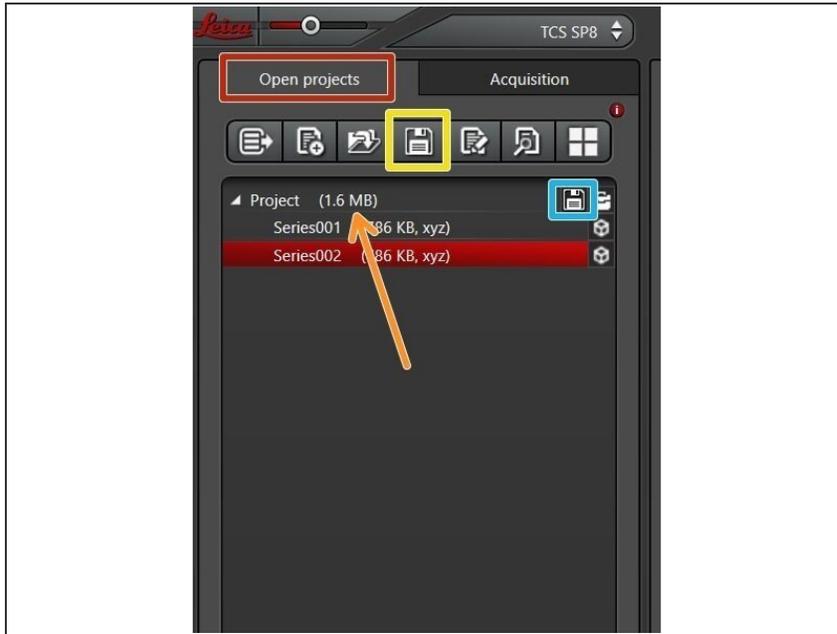


INTRODUCTION

How to save your data and log off or shut down the Leica SP8 confocal laser scanning microscope located at the Schlieren campus, room WAD-12-K-105 once you are done..

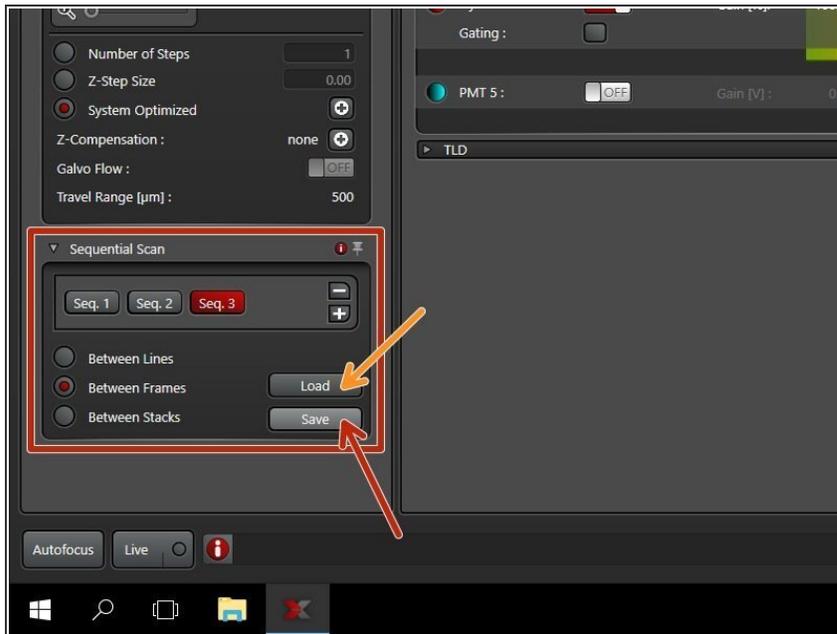
Please find detailed information about the system setup [here](#).

Step 1 — Saving your data



- Go to "**Open Projects**" tab.
- **Save your data** by either
 - right-clicking on the "Project"
 - using the "Save All" icon on top
 - or clicking the save sign behind the "Project"
- ☑ Save your data on your core storage (network path: "**\\files.core.uzh.ch**").
- ⓘ *Please follow our instructions [here](#) on how to access your data.*

Step 2

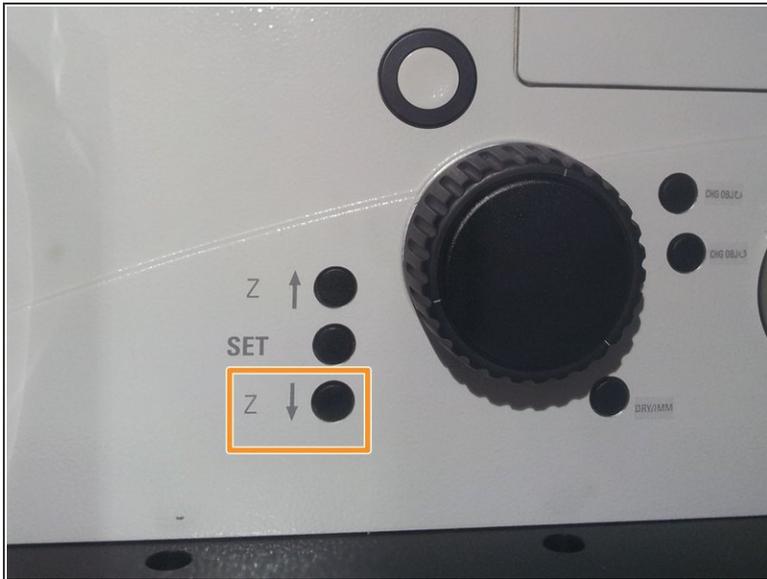


i You can save your settings and reuse them by loading them in your next imaging session. By default the following settings will be loaded: laser intensities, detector settings, averaging and accumulation settings.

- **Save your sequential scan settings** if you want to reuse it (use "**Home**" directory on the server).
- In your next imaging session you just need to **activate "Sequential Scanning"** and click "**Load**" in order to retrieve your scan settings.

i Alternatively, you can also **import** a previously recorded **LIF file**, right click an image and select "**Apply image settings**".

Step 3 — Cleaning



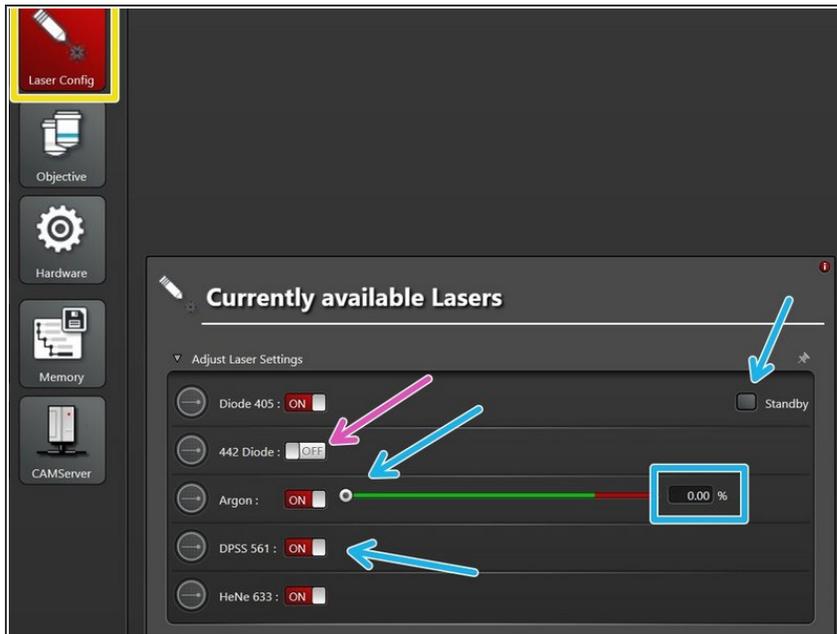
- **Lower the objective** and remove your sample.
 - Clean the immersion objective(s) you have used with the available "**KIMTECH**" wipes and **100% EtOH**.
- ⚠ Always clean the lenses in concentric circles and from the center to the edge. Clean also the sides of the objective(s).**
- ☑ Please also take care that immersion media has been cleaned off from all the other microscope parts which got in contact with.

Step 4 — Check booking system

The screenshot shows the EZbooking interface for 'Light Microscopes' with the object 'CLSM - Leica SP8 inverse (Schlieren)'. The calendar is set to 2021 July. A message banner indicates 'Violet 405 nm laser is back to full intensity! (10.03.21)'. A red arrow points to the booking slots for Monday, July 27, which are highlighted in yellow. The slots for Monday are: 09:30 - 12:00, 12:00 - 14:00, and 14:00 - 18:30. Other days have various time slots, such as 07:00 - 08:30 on Tuesday and 09:00 - 13:00 on Thursday.

- Check the [booking system](#) if there is another user scheduled within the next 2 hours.

Step 5 — Lasers and closing the "LAS X" software

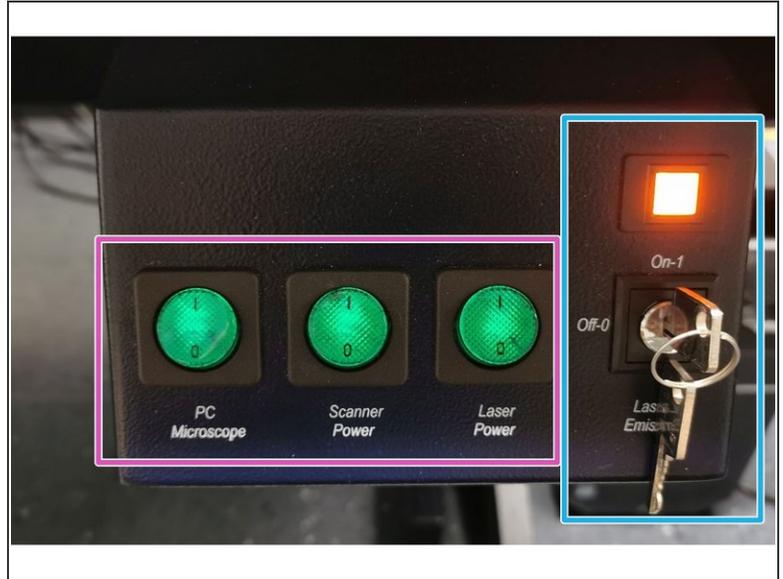
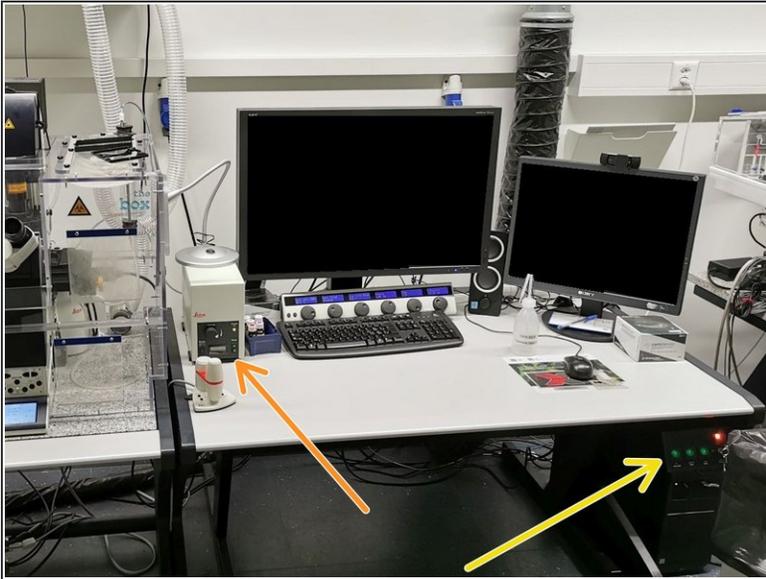


- Go to the "**Configuration**" tab and choose "**Laser Config**".

⚠ Does the booking system indicate another user within the next 2 hours?

- If YES
 - keep the **lasers ON** or put on **standby** and reduce **Argon** output to **0%**
 - **close** the "**LAS X**" software and **Sign-Out** of the Windows profile
- If NO
 - switch **OFF** all lasers
 - **close** the "**LAS X**" software, **shut-down the PC** and follow the next step

Step 6 — Switch OFF hardware



⚠️ Only follow this step if there is no booking within the next 2 hours.

- Switch-OFF the **fluorescence lamp**
- On the **main switch board**
 - turn the "Laser Emission" key to "Off-0"

⚠️ Wait for 5 min to cool down the lasers

- Finally switch OFF the "**Laser Power**", "**Scanner Power**" and "**PC/Microscope**" using the corresponding buttons