Manual for the LabView Integration UI

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1 User Guide

1.1 Prequisites

- The program is not very fast. In case something does not happen instantly, be patient and try to avoid clicking multiple times. This mostly results in unpredictable behavior.
- If the software is not already installed on the computer, refer to the README.md file.

1.2 Starting the software

- 1. Ensure that the following devices are booted and connected to the PC
 - · Olympus Microscope
 - TANGO Stage Controller
 - SOLA Light Engine
 - · Valve Controller
- 2. Start the LabvView project UI_Project.lvproj
- 3. Expand the VIs subfolder and start the latest version of the UI
- 4. Click the Run or Run-Continously button on the top left corner

- Normally, the COM ports can stay unmodified. However, (mainly after Windows Updates) these ports tend to change. Their correctness can be verified in the Device-Manager (Geräte-Manager) by plugging them out and back in.
- 6. The red lamps on the top-left corner turn green upon successful connection to the respective device.

If a lamp stays red:

- · Hit the STOP-Button and start over
- If the lamp is still red, switch the respective device off and on again.
- Ensure the correct COM-Port is selected. (Via Device-Manager or NI-Device Monitor)
- 7. Wait until the status field says "Initialization finished successfully"
- 8. Now, the program is fully operational.

1.3 Operating a Multiring chip

Make sure you started the program according to section 1.2.

1.3.1 Initalization

- 1. Start the Elveflow device and the ESI software
- 2. Disable all 4 valves in the software
- 3. Pressurize the Elveflow with 4 bar to 8 bar by turning the right valve (blue) between the pressure regulators above the microscope
- 4. Wipe the microfluidic top and bottom with Acetone, IPA and ddH20 to remove dust etc. (otherwise it will cause imaging artifacts)
- 5. Fill the control tubing at least (!) 20 cm with ddH2O with a 1 mL syringe and connect it to the respective inlet
- 6. Insert the chip carefully into the microscope
- 7. Ensure that the tubing does not interfere with the other devices, corners or edges and can move freely during stage driving. Move a sufficient length into the microscope to avoid pulling at the inlets.
- 8. Connect a 40 µM Fluorescein solution to the Elveflow
 - Use the small diameter PTFE tubing for the whole length.
 - Apply 10 mbar to 20 mbar at the respective ESI channel to move the fluid to the end of the tubing until a drop starts to form
 - Insert the connector to Inlet 1 of the chip
- 9. Connect ddH₂O or nfH₂O to the Elveflow
 - Apply 10 mbar to 20 mbar at the respective ESI channel to move the fluid to the end of the tubing until a drop starts to form
 - · Insert the connector to Inlet 2 of the chip
- 10. Build a connector from small-diameter PTFE-tubing to big-diameter PTFE-tubing; connect it to a waste container and the outlet of the chip (To minimize the hydraulic resistance and capacitive air effects)
- 11. Start the Labview program

- 12. Click the Live Button, then adjust Filter, Intensity and Exposure settings.
- 13. Open the controle pressure with the left, blue valve on top of the microscope and pressurize the control valves with approx. 1.5 bar by turning the center and left pressure regulator.
- 14. Click the No Flow Button to pressurize all quake valves

1.3.2 Loading the chip

- 15. Check the control inlets for leaks and wait for the channels to fill, control it in the Live mode. Now would be a great time for experienced users to save the different calibration positions.
- 16. Pressurize the **ESI** channels for water and fluorescein to 250 mbar
- 17. Select Reagent 1 and click the **Flush** Button to inject the fluorescein

1.3.3 Adjusting the valve pressure

- 18. Upon arrival, deactivate **Flush** and switch to the *GFP* filter; adjust *exposure* (exit live mode) and *intensity* as well as the histogram sliders
- 19. Use the 10x Objective and move to a valve in the reagent-multiplexer.
 - Increase the pressure until you see two meniski, which indicate that the valve closes completely.
- 20. Repeat the same procedure for the ring pump

1.3.4 Remove air

21. select Reagent 2 and click the Flush Button again to inject the water

For faster flushing, the ESI pressure can be increased temporarily up to 500 mbar

Upon arrival, let it flush as long as air is in the flushing channel between the rings

22. Deactivate the Flush, click All Valves and load all rings with Load

Wait until all rings are partly filled

Activate manually the valve A5 to load another side of the ring

Deactivate A5 and open B5 to close the outlet.

This will pressurize all channel and remove remaining air bubbles.

Test the RemoveAir button. It should do the same.

1.3.5 Performing the pump calibration

- 23. Click the No Flow Button
- 24. Ensure the *Position List* is empty

if not, hit the Clear Last button

- 25. Move to every ring, starting at the bottom (of live mode) and capture one position of each ring with **Save Pos**. (The bottom most ring corresponds to RingNr. 1)
- 26. Exit *Live Mode*, wait some time and hit **Capture Blank** of the calibration panel at the bottom; follow the instructions and wait until the acquisition is finished

- 27. Change to GFP, go into Live and Flush with Fluorescein for 10 s
- 28. Make sure All Valves is still on and load all rings with Load
- 29. Exit Live and Capture Full Intensity
- 30. Go into Live, make sure All Valves is still on and Load all rings with Fluorescein
- 31. Exit **Load** and adjust *Feed* and *Pump* cycles

Best results were achieved with >800 *Pump Cycles* and ~30 *Feed Cycles*

32. Hit Capture Dilutions

After the first dilution cycle, the (a bit under-)estimated remaining time is displayed At finish, an input dialog with 8 refresh ratios is displayed

33. Start MATLAB and run the CalibrationScript.m under lib \rightarrow Calib

Write the according **Refresh Ratios per Pump Cycle** back into Labview This is not ideal, I know...

1.3.6 Creating a program

34. Clear the *Position List* and capture your desired positions.

NOTE: As the focus is always saved with the position, it makes a difference if the positions are captured in 4x, 10x or another magnification

- 35. Save your position list by **Save List** with the suffix .xml in a desired location
- 36. Clear the Microscope Settings by clicking the Clear Button
- 37. Choose your desired acquisition settings (*Filter, Exposure, Intensity*) and **Save Settings**Repeat it for every color you want to acquire
- 38. Save List with the suffix .xml (preferably) at the same location as the positions
- 39. Make sure that the Refresh Ratios are set correctly. This is crucial for any further step!

If you want to change the Refresh Ratios, you can either doubleclick into the respective ratio and change it or hit the **Input Refresh Ratios** to input all ratios again.

- 40. Start the Program Setup
- 41. If you wish to modify a previously created program, click on Load Old Program
- 42. Select the previouly created *Microscope* and *Position Settings* by clicking the **Folder Symbol**If the lists have been loaded correctly, the LED beneath the Folder turns green
- 43. Now you can modify the program by clicking the respective buttons:

Feed: Exchanges a defined fraction of a single ring

Pump: Mixes all rings by actuating the ring pump. (Formula: Time = $\frac{\text{Pump Cycles}}{4 \text{ Hz}}$)

Change Reagent: Flushes the outer channels with the selected reagent for a defined time

Loop Start: Begins a unique Loop with a defined number of iterations

Loop End: Closes a unique Loop with its specified ID

Acquire: Acquires an image at all positions in all specified colors.

44. Hit Save and follow the instructions

The later filename is composed of **YYMMDD_hh_mm_ss_"YOURPREFIX".tif** For simple modifications you can also modify the .xml file accordingly.

1.3.7 Starting a program

- 45. Check if the refresh ratios are still there and valid
- 46. Hit the **Run Program** button
- 47. Select the previously created program.xml file and hit Load
- 48. Check if all values and steps are correct
- 49. Start the continuous loop via the Start button