## **Aurora Start-up Protocol**

- 1. Check sheath and waste tanks, replace if necessary
- 2. Turn ON the PC and log in:

User: My Aurora

Password: Welcome#1

Wait 3 minutes

3. Turn ON the cytometer (make sure there is a tube with  $H_2O$  installed in the SIP)

Wait 3 minutes

4. Launch the software SpectroFlo and log in:

Session: Common user Password: Common user

- 5. Go to Acquisition tab and select the experiment named: Cleaning
- 6. Clean the instrument by running:

5 minutes RINSE

5 minutes CLEAN

5 minutes H2O at flow rate HIGH. Put the flow rate back in LOW (as a preparation for the next steps).

- 7. Select the tab: QC and Set-up
- 8. Wait at least 15 additional minutes before running any sample. This time ensure the proper warming up of the detectors.
- 9. Prepare QC beads:

300ul of PBS 1X + 1 drop of Spectroflo beads (kept in the fridge of the facility)

10.Load the tube and run the beads by pressing start:

The bead lot number we are currently using is: 2003

- 11. Once the QC is done, a message saying, "Passed" should pops up.
- 12.Go to acquisition tab, select or create your experiment and start the acquisition.