

Quick Start Guide

Moxi GO II

Before you Begin:

Your Moxi GO II comes with a partial battery charge and must be run plugged into the supplied power adapter for charging and running tests.

Required Materials:

- Cassettes and diluent (e.g. PBS).
- Biological cell sample.
- Pipette and tips for 60µL aliquot.

Cassette Specifications:

- Cassette size range: 3 - 27µm particles
- Concentration range:
 - * Counts: 10,000 to 1,750,000 cells/ml
 - * Optimal Fluorescence Sensitivity: 100,000 - 500,000 cells/ml

Sample Prep Considerations:

- **Solution Conductivity:** Cells must be suspended in standard, conductive, lab media (e.g. PBS or equivalent)
- **Single-Cell Suspensions:** Cells need to be prepared as single cell suspensions. Clusters/aggregates should be broken apart with mechanical trituration and/or protease dissociation (e.g., Accutase). Samples with large extracellular debris particles or aggregates should be strained/filtered before running.
- **Fluorescent Labeling/Stains:** The Moxi GO II uses a 488nm laser with 525/45nm (e.g. FITC) and EITHER 561nm LP (e.g. R-PE, PI, 7-AAD, PE-Cy5) OR ~650nm/LP (e.g. PI, PE-Cy5) emission filters.
- ORFLO approved/kits reagents, and protocols are strongly recommended.

Data Transfer:

FCS 3.1 test data is available for transfer via USB. To transfer data

- Plug the unit into a PC/Mac with supplied cable.
- Touch the "USB Disk" icon on the Home Screen.
- Touch the "Connect" icon.
- The unit will appear as an external, flash drive.

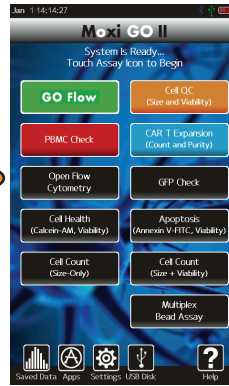
User Manual:

An electronic copy of the complete Moxi GO II user manual can be found at www.GEMBIO.com on the Moxi GO II product page (Under Products | Cell QC & Analysis)

1) Turn Moxi GO II ON



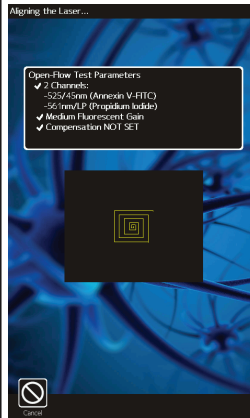
2) Touch Desired Assay



3) Open the Door. Insert Cassette. Close Door. (Note: on older models, tray lever needs to be depressed while inserting the cassette)



4) Auto Laser Alignment. Please Wait.

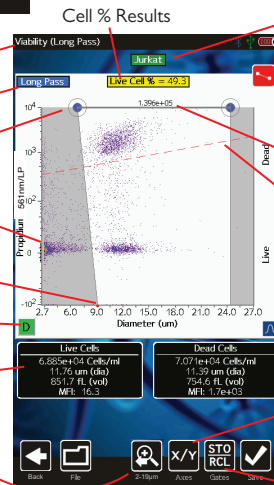


5) Pipette 60µL of Stained Sample Into the Cassette. Close Door. Test Begins.



6) Data Summary

- Application
- PMT Displayed
- Size Gates (Blue)
- Greyed Noise Region
- Toggle to Angle Gate
- Fluorescent Gain Setting
- Lower Gated Population Results
- Press to Change Size Scale Range: 2-26, 2-18, 2-10 µm

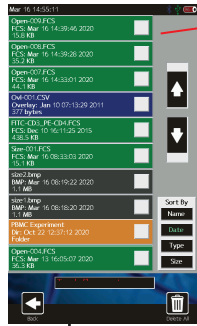
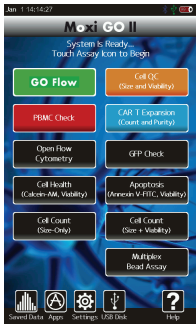


- Touch to Rename File
- Battery/Charging Indicator
- Toggle Between Size (Blue) and Fluorescent (Red) and Noise (Yellow) Gates.
- Total Count Between Size (blue) Gates
- Fluorescent Gate (Red)
- Turn on Histogram Overlays
- Upper Gated Population Results
- Select to Change Axis Display: PMT vs Size, Size/Fluorescence Histograms, PMT vs PMT
- Store/Recall Gate Locations

Dot Plot Results



Opening, Viewing, Comparing, and Printing Data:

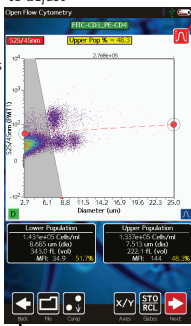


File Colors
 Green = Tests
 Blue = Overlays
 Black = BMPs
 Orange = QC batches/directories

Touch and drag gates to adjust

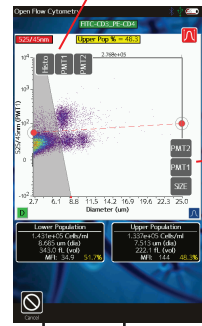
Touch red icon to toggle gate modes

Specify y-axis parameter



X/Y

Specify x-axis parameter



Test Comparisons/Overlays

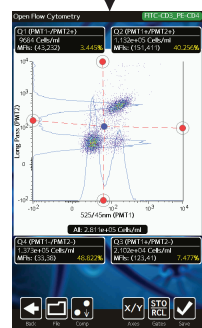
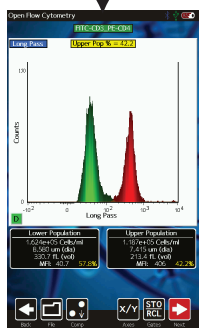
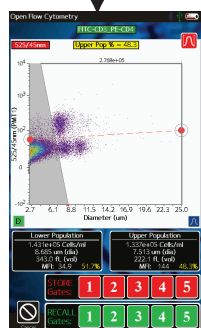
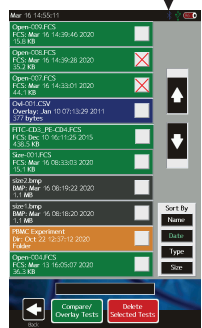


STO RCL

HOLD

PMT2

PMT1

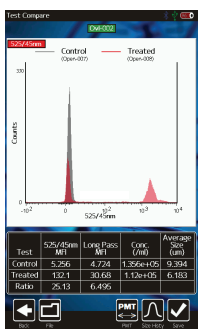
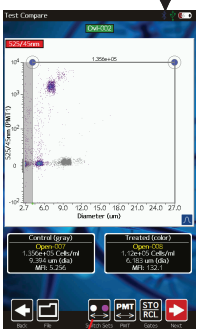


Touch number to store/recall gates into from memory

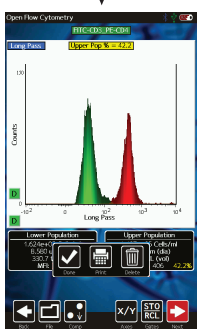
Swipe up/down on histogram to scale

Touch & drag gates to adjust. Note: You can tap blue dot & tap to the side of it to move it with better visibility.

Use blue size gates to define region to be compared



Print Screen (export to .bmp file)



Exact screenshots saved as a .bmp file on the data disk (access through USB connection to the computer)

"Switch Sets" toggles the "control" and "treated" set designations