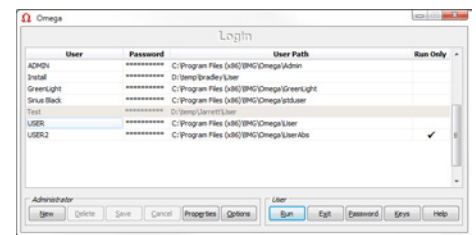


Quick Guide Omega Software

STARTUP

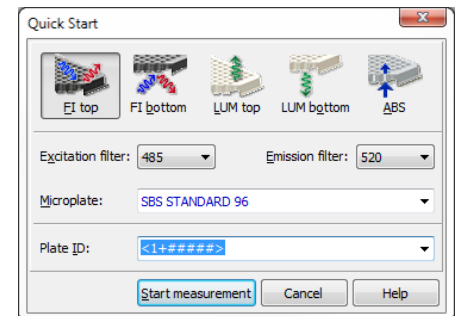
- Turn on the instrument and the computer.
- Start the **Omega Control** software.
- Login with your password or just click 'Run' to login as "User".



To measure a microplate, you can either use the quick start function or you can execute a pre-defined test protocol.

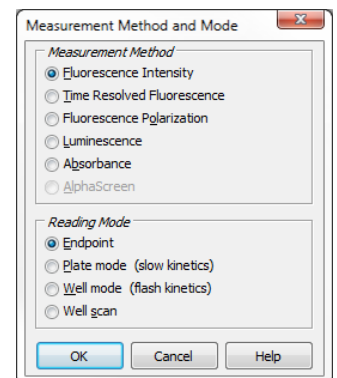
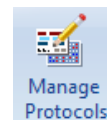
QUICK START

1. To measure a full plate in endpoint mode without defining a test protocol, click the 'Quick Start' button:
2. Select the measurement method. Choose the **excitation** and **emission filters** and the type of **microplate** that will be used.
3. A plate identifier (**Plate ID**) can also be specified (optional).
4. **Start the measurement.**

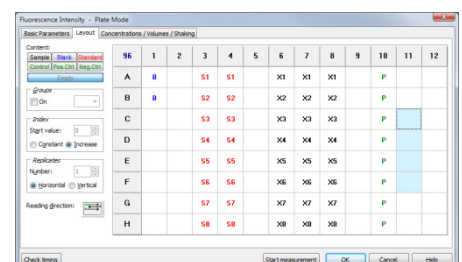
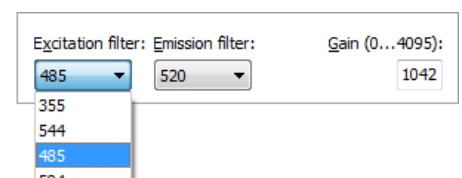
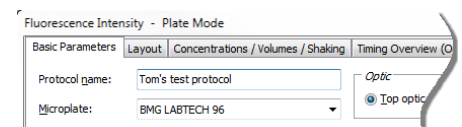


PROTOCOL DEFINITION

1. To create a new **test protocol** or to edit an existing one:
 - Click the 'Manage Protocols' button:
 - Double click the **protocol name** to edit an existing protocol or click 'New' to create a new protocol. Choose the *Measurement Method* (FI, FP, TRF, luminescence, absorbance) and choose the *Reading Mode*:
 - **End point** for single readings
 - **Plate mode** for slow kinetics
 - **Well mode** for fast kinetics
 - **Well scan** for scanning (useful if you use large wells and if the samples are not equally distributed)

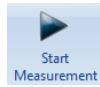


2. Inside the protocol definition window:
 - Enter a **protocol name**.
 - Choose the **microplate** being used (Greiner, Corning, Nunc, etc.).
 - Select between **rapid** or **precise** measurement, depending on your assay.
 - *Plate Mode Kinetics*: Type in the **no. of cycles** (how many times the reader will cycle through the plate).
 - *Well Mode Kinetics*: Type in the **no. of intervals** (how many times the reader will read the well).
 - Choose the **excitation** and **emission filters** to be used.
 - Select the 'Layout' sheet. Enter the position of samples, blanks and standards (if any).
 - If standards and/or reagent dispenser(s) are used, type in the values in the 'Concentrations / Volumes / Shaking' window.
 - Click the 'Check timing' button. This gives you the smallest possible cycle time (*Plate Mode*) or interval time (*Well Mode*). A longer time can be achieved by typing in a higher value in the 'Basic Parameters' sheet.

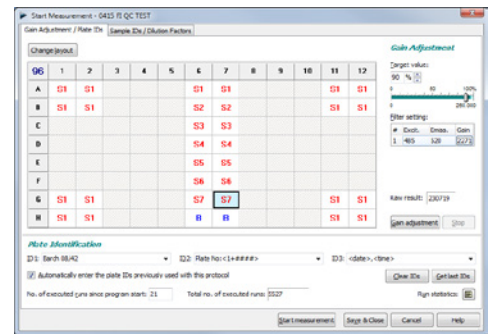


Quick Guide Omega Software

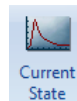
MEASURING (executing pre-defined protocols)



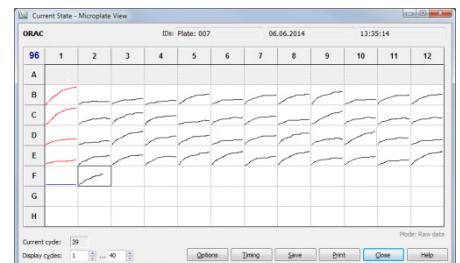
1. Click the 'Start Measurement' button:
2. It is possible to define up to three plate identifiers in the 'Gain Adjustment / PlateIDs' sheet.
3. In the 'Gain Adjustment / PlateIDs' sheet, select the well that will have the highest intensity and click the 'Gain adjustment' button:
 - The **required value** should be 90% in endpoint readings (giving highest values around 260 000 - 10% = 234 000).
 - For kinetic measurements, 10% - 50% could be the required value (this is dependent on the expected increase in the signal).
4. Click the 'Start measurement' button.



RESULTS



1. To see the measurement results during a reading:
 - Click the 'Current State' button. Different display options are available, e.g. curve, spectra...
2. To perform data calculations using the **MARS Data Analysis** software:



- Close the 'Current State' window.
- Click the 'MARS' button:



3. In the 'Manage Test Runs' window:
 - Double click the **test name** of the test run to be analyzed

Test ID	Test Name	ID 1	ID 2	ID 3	Date	Time	Measurement Method	Signal
545	BRACARD 30	peroxide hydrog.	cell bioassay		02.18.2008	14:56:15	Absorbance spectrophotometry	

4. Analyze the measured data:
 - Select the data to be displayed in the working area with the navigation tree (Data Node) on the left side of the main window.
 - Use the standard calculation wizard to perform a quick curve fit calculation; or use the calculation menus to define what is to be calculated and to be displayed.
 - To see a standard curve, open the 'Standard Curve' page. The calculated unknowns are displayed in the 'Microplate View' and the 'Table View'.
 - To remove outliers, simply shade them out in the 'Microplate View' using the toggle function (Ctrl-T).
 - For kinetic measurements (more than one measured cycle or interval), choose the range(s) of interest (**Calc. Start** and **Stop**) and the data values from these ranges can be evaluated using a kinetic calculation.

