

Qubit™ Flex Fluorometer

USER GUIDE

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Revision history: MAN0018186

Revision	Date	Description
B.0	23 January 2020	Changing “A” product skus to new “Q” skus.
A.0	11 October 2019	New user guide.

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About this guide

Overview This user guide describes how to operate the Qubit™ Flex Fluorometer.

User documentation The guides listed below are available with the Qubit™ Flex Fluorometer.

Guide	Pub. No.
<i>Qubit™ Flex Fluorometer User Guide</i>	MAN0018186
<i>Qubit™ Flex Fluorometer Quick Reference Card (QRC)</i>	MAN0018187

Additional resources are available on the Qubit™ Technical Resources page. Go to [thermofisher.com/qubit](https://www.thermofisher.com/qubit) to access protocols, application notes, and tutorials.

Text and keyboard conventions Text and keyboard conventions used in the *Qubit™ Flex Fluorometer User Guide* are listed below. For safety alert words and symbols used in Thermo Fisher Scientific user documentation, see page 4.

Convention	Use
Bold	Bold text indicates user action. For example: Click Run .
►	Right arrow symbol (►) indicates a menu choice, and separates successive commands you execute or select from a drop-down or shortcut menu. For example: Select Settings ► Instrument Settings .

User attention words Two user attention words appear in Thermo Fisher Scientific user documentation. Each word implies a particular level of observation or action as described below.



Note: Provides information that may be of interest or help but is not critical to the use of the product.



IMPORTANT! Provides information that is necessary for proper instrument operation, accurate installation, or safe use of a chemical.

Safety alert words Four safety alert words appear in Thermo Fisher Scientific user documentation at points in the document where you need to be aware of relevant hazards. Each alert word—**IMPORTANT**, **CAUTION**, **WARNING**, **DANGER**—implies a particular level of observation or action, as defined below:



IMPORTANT! – Provides information that is necessary for proper instrument operation, accurate installation, or safe use of a chemical.



CAUTION! – Indicates a potentially hazardous situation that, if not avoided, may result in minor or moderate injury. It may also be used to alert against unsafe practices.



WARNING! – Indicates a potentially hazardous situation that, if not avoided, could result in death or serious injury.



DANGER! – Indicates an imminently hazardous situation that, if not avoided, will result in death or serious injury. This signal word is to be limited to the most extreme situations.

Except for **IMPORTANT!** safety alerts, each safety alert word in a Thermo Fisher Scientific document appears with an open triangle figure that contains a hazard symbol. These hazard symbols are identical to the hazard symbols that are affixed to Thermo Fisher Scientific instruments (see “**Safety symbols**” in Appendix C).

SDSs The Safety Data Sheets (SDSs) for any chemicals supplied by Thermo Fisher Scientific are available to you free 24 hours a day. For instructions on obtaining SDSs, see “**Safety Data Sheets (SDS)**”.



IMPORTANT! For the SDSs of chemicals not distributed by Thermo Fisher Scientific contact the chemical manufacturer.

1. Product information

Product contents

The Qubit™ Flex Fluorometer (Cat. No. Q33327) is shipped with the following components:

Component	Quantity
Qubit™ Flex Fluorometer	1 each
Qubit™ Flex power cord (shipped separately) ^[1]	1 each
USB drive	1 each
Qubit™ Flex LAN cable	1 each
Qubit™ Flex Fluorometer Quick Reference Card (QRC)	1 each
Certificate of Conformity (COC)	1 each
Qubit™ screen cleaning cloth	1 each
Wi-Fi Dongle	1 each

^[1] The power cords for the Qubit™ Flex Fluorometer are not interchangeable with those for the other Qubit™ Fluorometer models. Powering the instrument with an unapproved power cord can irreversibly damage the instrument.

The complete user guide is available for download at thermofisher.com/qubit.

See page 6 for the description and specifications of the Qubit™ Flex Fluorometer.

Upon receiving the instrument

Examine the instrument carefully for damage incurred during transit. Ensure that all parts of the instrument, including the accessories listed above, are included with the product. Damage claims must be filed with the carrier; the warranty does not cover in-transit damage.

See page 10 for instructions to set up the Qubit™ Flex Fluorometer.

Register your instrument

Go to thermofisher.com/qubit to register your instrument. You will be asked to supply the serial number, your name, and your contact details. Registering your instrument ensures that you will receive notifications of software upgrades and information on new assays for use with the Qubit™ Flex Fluorometer.

Product description

Qubit™ Flex Fluorometer The Qubit™ Flex Fluorometer is a benchtop fluorometer for the quantification of DNA, RNA, microRNA, and protein. With the Qubit™ Flex Fluorometer, you can directly measure the fluorescence of up to 8 samples simultaneously using the highly sensitive and accurate fluorescence-based Qubit™ assays.

- Features**
- Fast and highly accurate quantification of DNA, RNA, and protein of up to 8 samples simultaneously in ~3 seconds.
 - High levels of accuracy using only 1–20 µL of sample, even with very dilute samples.
 - Use of dyes selective for dsDNA, RNA, or protein minimizes the effects of contaminants in the sample.
 - Stores results from up to 10,000 samples.
 - 8-inch, state-of-the-art color touchscreen for easy workflow navigation.
 - Instrument indicates samples that are in the extended range or out of range.
 - Saves sample data as a CSV (comma separated value) file.
 - On-board Reagent and Range Calculators provide instructions to prepare Qubit™ working solution using your sample and standard inputs and to select the most accurate assay for your expected concentration range.
 - On-board Molarity and Normalization Calculators allow you to calculate molarity of your samples based on nucleic acid length and determine how to dilute the samples to the same concentration, respectively, using the results from your assays.
 - Allows easy definition and saving of assay preferences.
 - Exports data to a USB drive, to a network drive, or to the Connect™ cloud-based platform.
 - Connects to the local area network via the LAN (RJ-45) port using an Ethernet cable or wirelessly using the supplied Wi-Fi adaptor.
 - Instrument user interface can be personalized to display in the language of your choice including English, French, Spanish, Italian, German, simplified Chinese and Japanese.

Instrument exterior components

Top view




Rear view



- ① **Touchscreen** is the user interface containing the controls for all the functions needed and displays data from the assays.
- ② **Sample chamber** is used to load the Qubit™ Flex Tube Strip containing your samples into the fluorometer for analysis.
- ③ **USB drive ports (Type A)** allow you to transfer and save data to your computer using a USB flash drive or wirelessly to a network drive or a Connect™ account using the Wi-Fi dongle (supplied with the instrument).
- ④ **Power inlet** connects the Qubit™ Flex Fluorometer to an electrical outlet using the supplied power cord and the appropriate plug.
- ⑤ **LAN port (RJ-45)** allows you to connect to the network using an Ethernet cable.

Product specifications

Physical characteristics	Instrument type:	Benchtop fluorometer
	Instrument dimensions:	7.3 in (w) × 11.1 in (l) × 4.1 in (h) (18.6 cm × 28.2 cm × 10.3 cm); rectangular shape
	Weight:	60 oz. (1.7 kg)
	Operating power:	100–240 ±10% VAC, 1.3 A
	Frequency:	50/60 Hz
	Electrical input:	48 VDC, 1.87 A
<div>  IMPORTANT! If the supplied power fluctuates ±10% beyond the rated voltage, a power line regulator may be required. High or low voltages can adversely affect the electronic components of the instrument. </div>		
Operating conditions	Installation site:	Indoor use only
	Altitude:	Between sea level and 2000 m (6500 ft.) above sea level
	Operating temperature:	10–30°C
	Operating humidity:	15–80% (non-condensing)
	Pollution degree:	The instrument has a Pollution Degree rating of II. The instrument may only be installed in an environment that has nonconductive pollutants. Typical environment with a Pollution Degree II rating are laboratories and sales and commercial areas.
Technical specifications	Dynamic range:	4 orders of magnitude
	Processing time:	≤3 seconds/sample
	Light sources:	Blue LED (max 460–480 nm) Red LED (max 620–640 nm)
	Excitation filters:	Blue 456–484 nm Red 612–644 nm
	Emission filters:	Green 513–563 nm Far-Red 671–693 nm
	Detectors:	Photodiodes; measurement capability from 320–1100 nm
	Calibration type:	2- or 3-point standard
	Sample chamber:	Accommodates one Qubit™ Flex Tube Strip
	Tube type:	Qubit™ Flex Tube Strip (8× 0.2-mL thin-wall polypropylene tubes; Cat. No. Q33252)
	Warm-up time:	<35 seconds

Hardware	Display:	8-inch capacitive touchscreen with high resolution color display
	Output ports:	3× USB ports
	Networking capability:	Connection via the LAN (RJ-45) port using an Ethernet cable or wirelessly using the supplied Wi-Fi adaptor
	Power supply:	AC adaptor with country-specific power cords
USB drive	Capacity:	4 Gigabyte

2. Getting started

Set up the Qubit™ Flex Fluorometer

Install the instrument

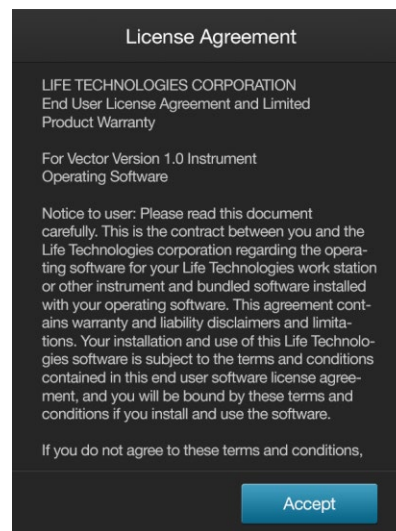
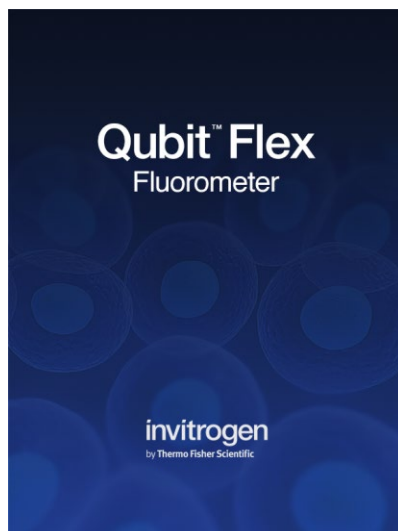
The Qubit™ Flex Fluorometer is a stand-alone instrument that does not require connection to a computer.

1. After unpacking the instrument, place the instrument on a flat, level, dry surface.
2. Plug one end of the supplied power cord into the Qubit™ Flex Fluorometer.
3. Attach the appropriate plug adaptor to the other end of the power cord.
4. Plug the power cord into the electrical outlet. Ensure that the power adaptor plug remains accessible to allow disconnection.



IMPORTANT! Use the power cord plug adapter supplied with the instrument that is appropriate for the electrical outlet configuration in your country. Powering the instrument with an unapproved power cord can irreversibly damage the instrument. Note that the power cords for the Qubit™ Flex Fluorometer are not interchangeable with those for the other Qubit™ Fluorometer models.

5. The instrument automatically powers on, first displaying the splash screen, then the **End User License Agreement (EULA)** screen.



Note: The End User License Agreement (EULA) screen is displayed on the first use of the instrument. On subsequent uses, the **Home screen** (page 14) is displayed after the splash screen.

6. Click **Accept** to accept the terms of the agreement and proceed to “Set language and date/time options” (page 11).



Note: You can also view and export the EULA from the **About Instrument** screen (page 15).

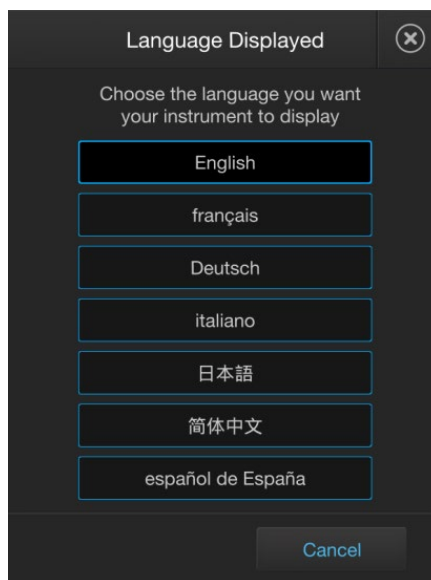
7. To power down the Qubit™ Flex Fluorometer, unplug it.

Set language and date/time options

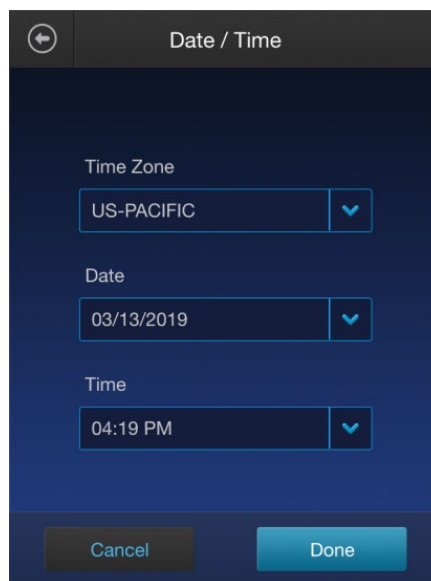
After you accept the EULA, the instrument shows the **Language displayed** and **Date/Time** screens, which allow you to set language and date/time options. If you wish, you can later change the language settings from the **Settings ► Instrument Settings ► Language** screen (page 85).

1. On the **Language displayed** screen, select the **Language** you want your instrument to display, then press **Next**.

Available options are **English, French, German, Italian, Chinese, Japanese, and Spanish**.



2. On the **Date/Time** screen, select the **Time Zone**, set the **Date** and **Time** in the desired format, then press **Next**.



Note: For detailed instructions on how to configure date/time options and to set the date and time, see “Set the date and time”, page 75.

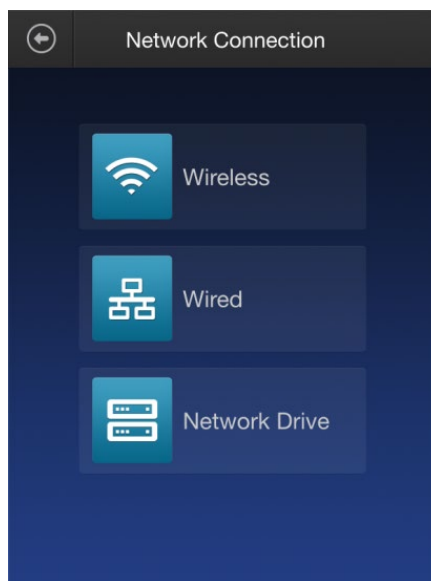
Connect to the network

(Optional) Connect to the network

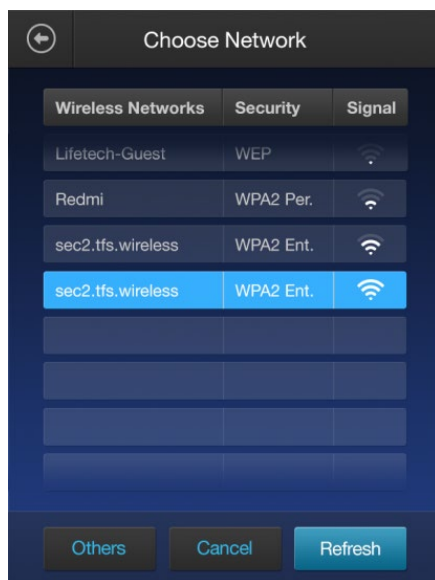
After you set language and date/time options, the instrument displays the Network Connection screen, which allows you to configure network options. If you wish, you can skip this step and connect to the network later from the **Settings ► Instrument settings ► Network connection** screen (page 78).

1. On the **Network Connection** screen, select **Wireless** or **Wired** connection.

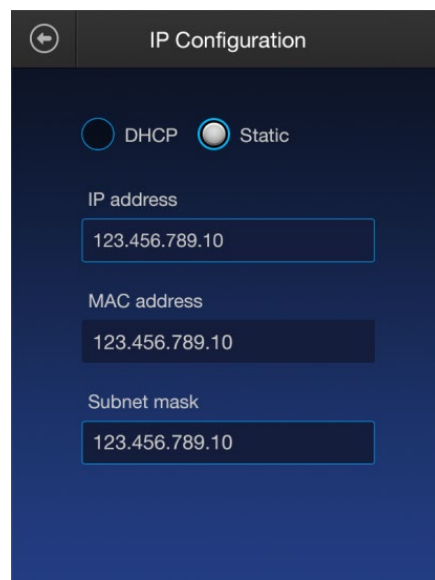
If you wish to use the instrument without joining a network, press **Skip**. You can always join a network and configure network settings later.



2. Depending on your choice, the instrument displays the **Choose Network** or the **IP Configuration** screen (for Wireless and Wired connection, respectively).



Choose Network screen
(for Wireless connection)



IP Configuration screen
(for Wired connection)

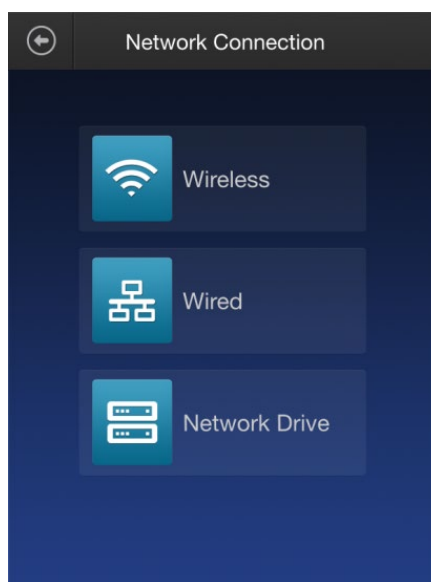
3. For wireless connection, select the network you want to join, then follow the on-screen instructions to configure the network options. When finished, press **Join**.

For wired connection, configure the network connection options, then press **Done**.

For detailed instructions on how to join a network (wireless or wired) and configure network options, see “Network connection”, page 78.

4. On the **Network Connection** screen, click **Network Drive** to map the location on the network where you want to save your Qubit™ Flex files.

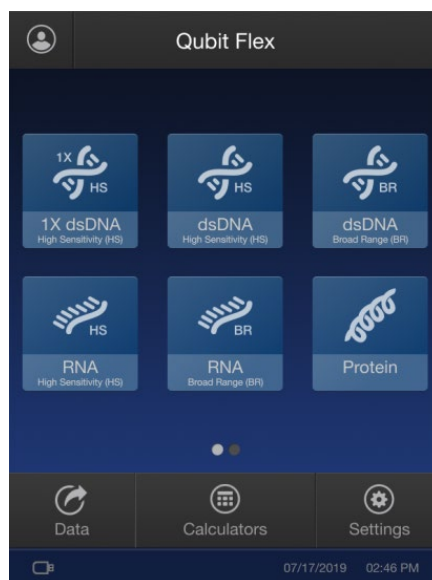
For detailed instructions on how to map a network drive, see “Map a network drive”, page 81.



Note: You must have an established network connection to map a network drive. If you wish, you can map the network drive later.

After instrument setup

Home screen After you have set instrument preferences, the instrument automatically displays the **Home** screen each time it is powered on.



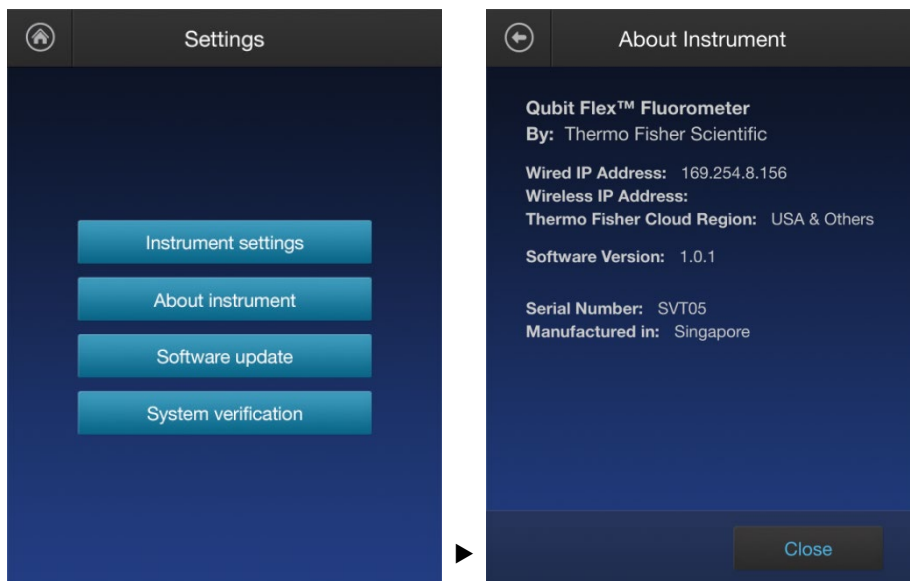
From the Home screen, you can:

- Sign in to your local instrument profile or your Connect™ account.
- Select the assays to perform:
 - 1X dsDNA High Sensitivity (HS)
 - dsDNA High Sensitivity (HS)
 - dsDNA Broad Range (BR)
 - RNA High Sensitivity (HS)
 - RNA Broad Range (BR)
 - Protein
 - Oligo (ssDNA)
 - microRNA
- Access saved data.
- Filter, delete, or export data.
- Configure instrument settings.
- Use the Reagent Calculator to determine the exact volumes of Qubit™ buffer and reagent required to prepare the Qubit™ working solution.
- Use the Range Calculator to determine the best assay to use for your sample.

About Instrument screen The **About Instrument** screen displays information about your Qubit™ Flex Fluorometer, including the currently installed software version.

To access the About Instrument screen:

1. On the **Home** screen, press **Settings**.
2. On the Settings screen, press **About Instrument** to display the About Instrument screen.



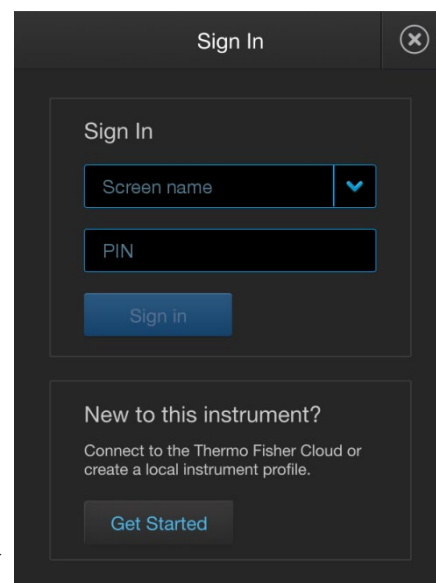
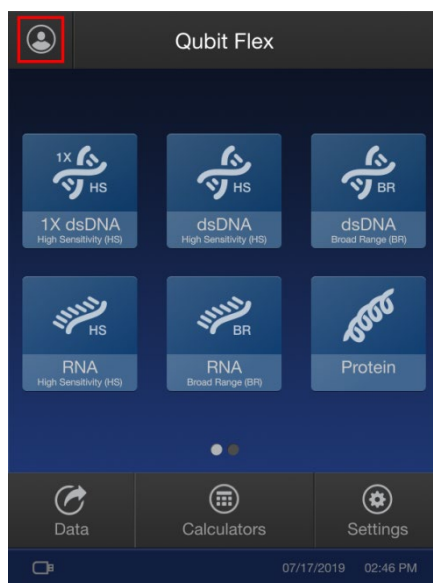
3. Press **Close** or **Back** (⬅️) to return to the Settings screen.

Sign in

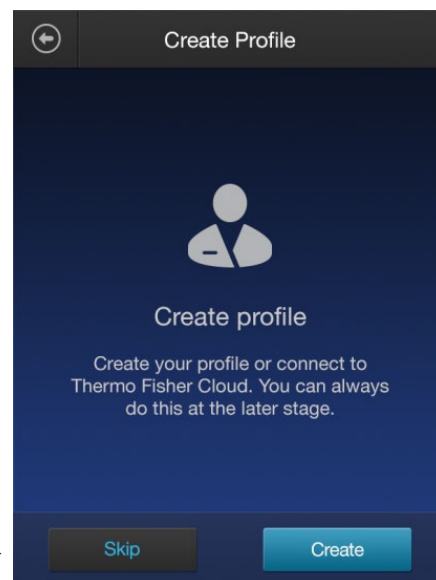
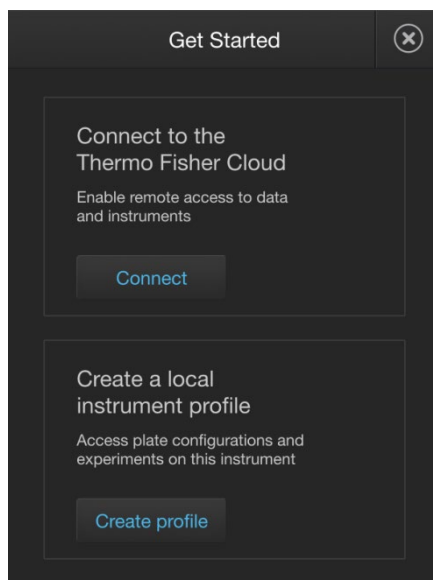
Create a local instrument profile

Qubit™ Flex Fluorometer allows you to create a local instrument profile for each user. A local instrument profile allows you to save to a mapped network location and it is also required to connect to your Connect™ account. If you wish, you can skip this step and create a profile later from the **Profile** screen.

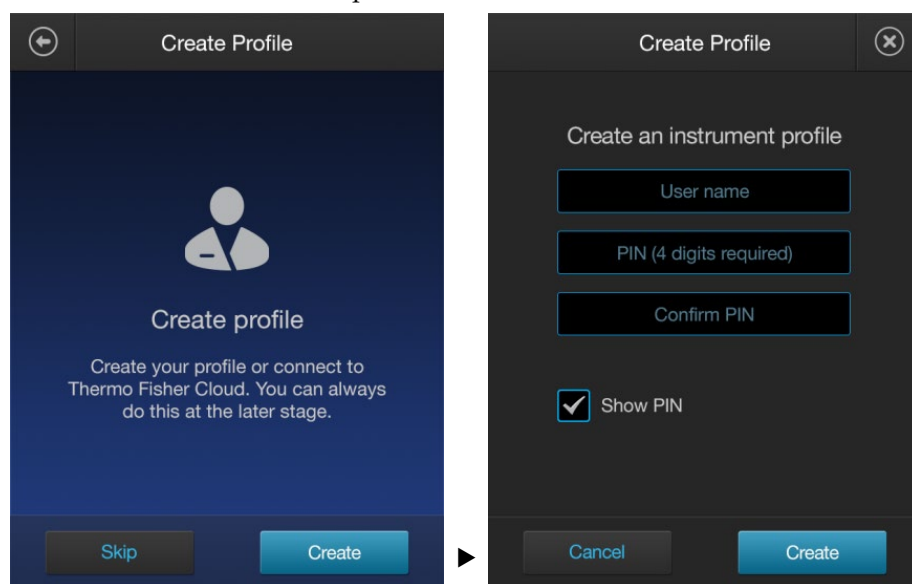
1. On the **Home** screen, press the **Profile** button on the top left corner of the screen to open the Sign In screen.



2. If you are new to the instrument and have not yet created a profile, press **Get Started** to open the Get Started screen, then press **Create Profile**.

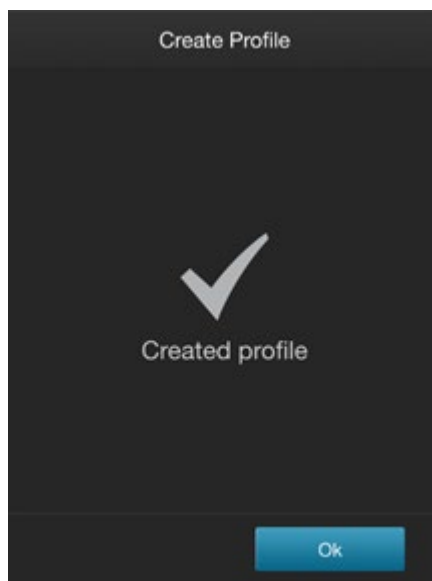


3. On the **Create Profile** screen, press **Create**.



If you wish to use the instrument without creating a local profile, press **Skip**. You can always create an instrument profile later.

4. Press the **User name** field, enter the desired user name for the profile (1–20 alphanumeric characters, no spaces), then press **Done**.
5. Press the **PIN** field, enter a 4-digit PIN, then press **Done**.
6. Enter the PIN in the **Confirm PIN** field, then press **Done**.
7. Press **Create** to create the local instrument profile.



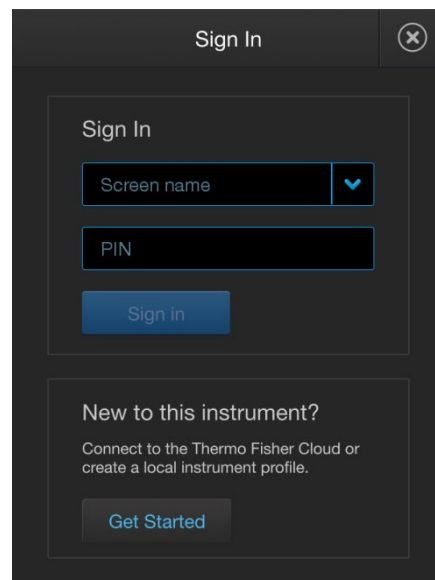
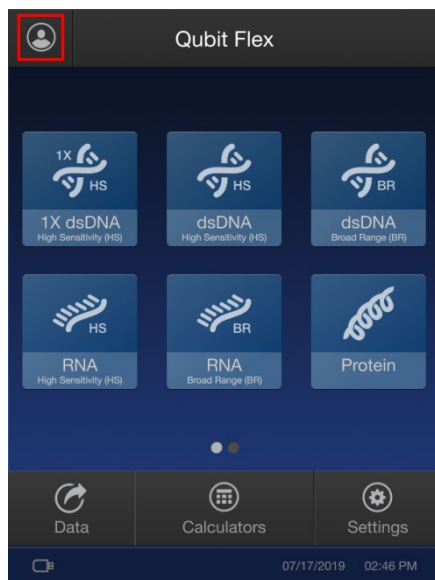
Sign in to your Connect™ account

After you have joined a network, you can also connect to your Connect™ account, Thermo Fisher's cloud-based platform, to store and access your data files.

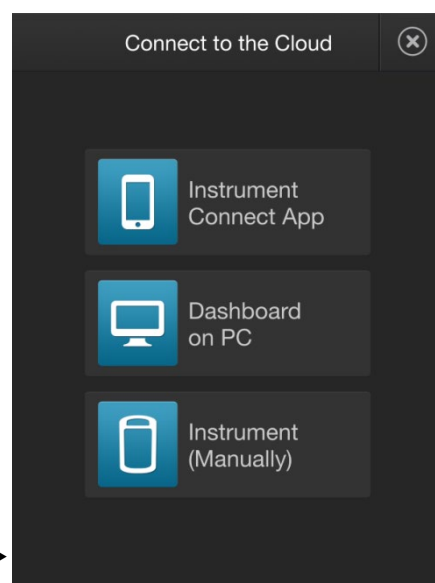
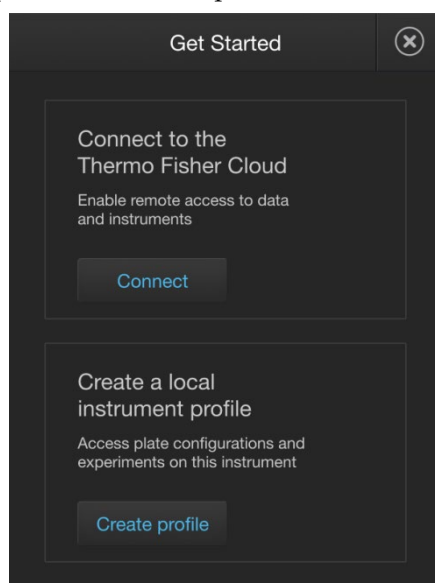


Note: To connect to the Thermo Fisher Cloud, you must have a Connect™ account or create one. To create your Connect™ account online or to sign in to your existing account, go to thermofisher.com/cloud.

1. Ensure that you are connected to the network on your Qubit™ Flex instrument (page 12).
2. On the **Home** screen, press the **Profile** button on the top left corner of the screen to open the Sign In dialog.

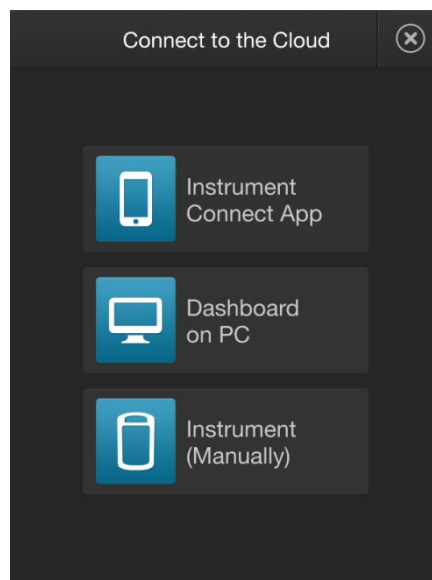


3. On the Sign In screen, press **Get Started** to open the Get Started screen, then press **Connect** to open the Connect to the Cloud screen.



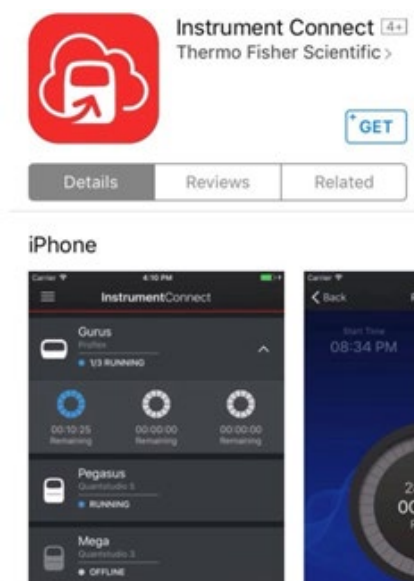
4. Connect to the Cloud screen offers three methods to sign in to your Thermo Fisher Connect™ account:

- **Instrument Connect App** on your mobile phone (Step 5, page 19)
- **Dashboard on PC** (Step 6, page 20)
- **Instrument (Manually)** (Step 7, page 21)

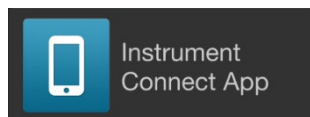


5. To connect to your Thermo Fisher Connect™ account with the **Instrument Connect App** on your mobile phone:

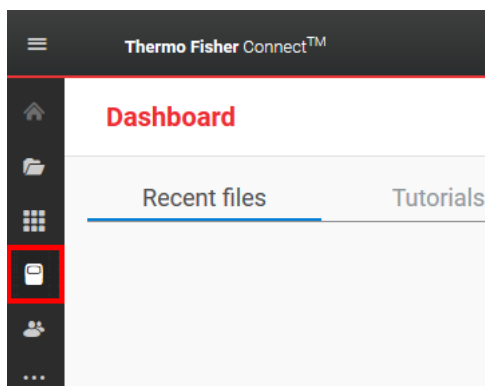
- a. Download the **Instrument Connect Mobile App** from the application store on your mobile phone.



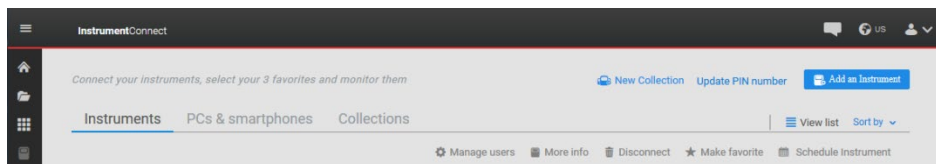
- b. Press **Instrument Connect App** on the Connect to the Cloud screen, then follow the steps on the Qubit™ Flex instrument. When finished, go to Step 8 (page 22).



6. To connect to your Thermo Fisher Connect™ account with **Dashboard on PC**:
 - a. Go to **thermofisher.com/cloud** and sign in to your Thermo Fisher Connect™ account.
 - b. On the Connect™ dashboard, press the **Instrument Connect** button.



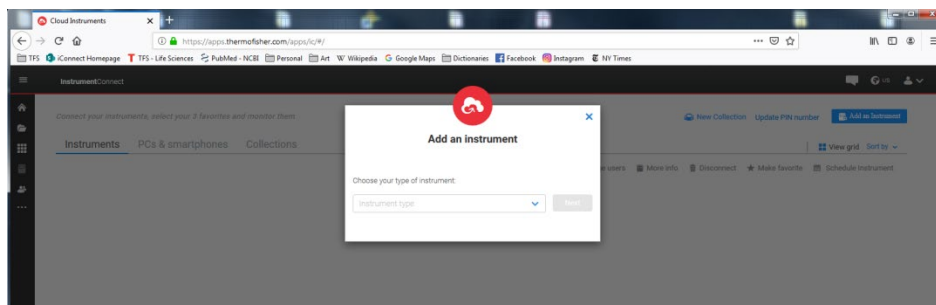
Instrument Connect screen opens.



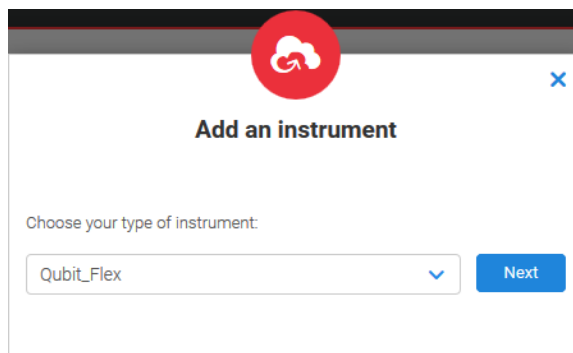
- c. On the Instrument Connect screen, press **Add an instrument**.



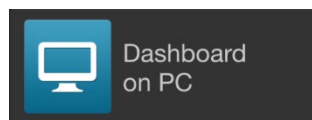
Add an instrument dialog opens.



- d. From the instrument type dropdown, select **Qubit_Flex**, then press **Next**.



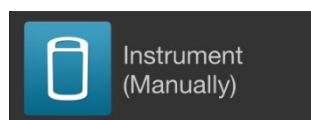
- e. Press **Dashboard on PC** on Connect to the Cloud screen (on the Qubit™ Flex instrument; see page 19) to display the linking code.



- f. Enter the linking code displayed on the Qubit™ Flex instrument into the Add an instrument dialog, then press **Send**.

A white dialog box with a red circular close button (X) in the top right corner. The title "Add an instrument" is centered at the top. Below the title, the text "Insert linking code:" is followed by a smaller line of text "Get the code from your instrument". There is a text input field with the placeholder text "Type your linking code here" and a grey "Send" button to its right.

- g. When finished, go to Step 8 (page 22).
7. To connect to your Thermo Fisher Connect™ account with **manually with the Qubit™ Flex instrument**:
- a. Press **Instrument (Manually)** on the Connect to the Cloud screen (on the Qubit™ Flex instrument; see page 19).



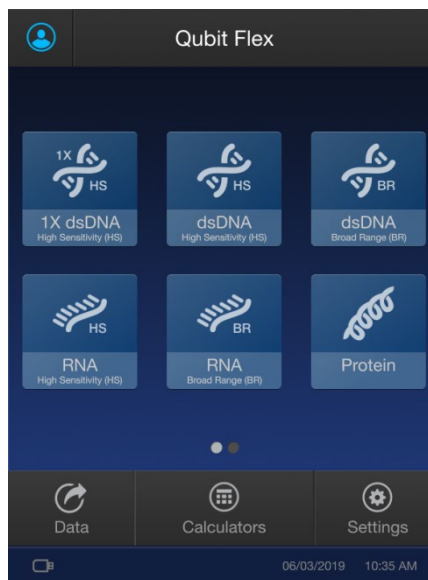
- b. Enter your **User name** and **Password** for your Thermo Fisher Connect™ account, then press **Connect**.

A dark grey dialog box with a white circular close button (X) in the top right corner. The title "Connect to the Cloud" is at the top. Below the title, the text "Connect to your Thermo Fisher Cloud account" is centered. There are two text input fields, one labeled "User name" and one labeled "Password". At the bottom, there are two buttons: a grey "Cancel" button and a blue "Connect" button.

8. When you have signed in to your Thermo Fisher Connect™ Account, the Profile button on the Home screen becomes blue.



When signed in, you can export your data to your Connect™ account.



Guidelines for using the Qubit™ Flex Fluorometer

Recommendations To obtain the best results, follow the recommendations below. For more information, see “Critical Qubit™ Assay considerations”, page 99.

- Do **not** operate the instrument in direct sunlight.
- Wear gloves during sample handling.
- Use the instrument at room temperature only (22–28°C).
- Bring all kit reagents to room temperature and insert all assay tubes into the instrument only for as much time as it takes for the instrument to measure the fluorescence.
- Do not hold the assay tubes in your hand before performing a measurement.
- Make sure that you have calibrated the Qubit™ Flex Fluorometer using the appropriate standards.
- The assay volume must be 200 µL for an accurate read.
- Take care not to create air bubbles when mixing the sample or standard with the working solution.
- Incubate the tubes for the Qubit™ DNA and RNA assays for 2 minutes after mixing the sample or standard with the working solution.
- Incubate the tubes for the Qubit™ protein assays for 15 minutes after mixing the sample or standard with the working solution.
- If you are performing multiple readings of a single tube, remove the tube from the instrument and let it equilibrate to room temperature for 30 seconds before taking another reading.

Note: Multiple readings of RNA samples are not recommended.

- Visit [thermofisher.com/qubit](https://www.thermofisher.com/qubit) for additional application notes, technical notes, citations, software updates, and a list of validated Qubit™ assays that have been tested using the Qubit™ Flex Fluorometer.

Assay tubes for the Qubit™ Flex Fluorometer Only thin-wall, clear 0.2-mL PCR tube strips are appropriate for use in the Qubit™ Flex Fluorometer. For best results, we recommend using Qubit™ Flex Tube Strips (Cat. No. Q33252).

3. Perform assays

Before you begin

- Materials needed**
- A Qubit™ assay kit appropriate for quantifying your samples (see page 101 for available Qubit™ assay kits and ordering information)
 - DNA, RNA, or protein samples in Qubit™ Flex Tube Strips
 - Appropriate standards for your assay in Qubit™ Flex Tube Strips
 - Single channel pipette (1–20 µL), multichannel pipette (200 µL)
 - Qubit™ Flex Reservoir (Cat. No. Q33253) or other suitable sample reservoir



Note: For instructions on the preparation of the assay standards, see the instructions that accompany the assay you are using or the *Qubit™ Flex Fluorometer Quick Reference Card (QRC)* (Pub. No. MAN0018187).

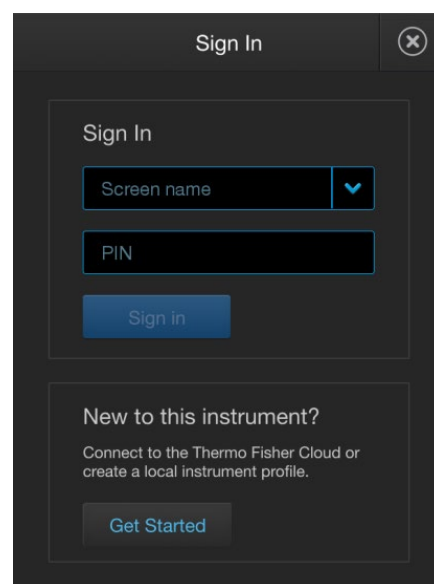
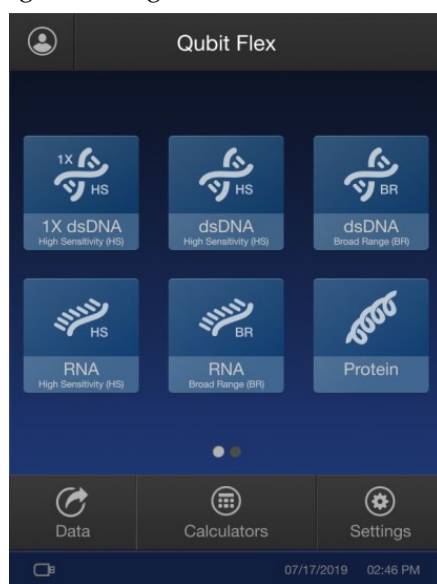
- (Optional) USB drive, USB cable, or Ethernet cable for data transfer, supplied with the instrument or available separately



Note: You can also transfer your data to a network location or your Connect™ account wirelessly, if you have set up a wireless connection.

Sign in to your profile

1. Press the **Profile** button on the top left corner of the screen to open the Sign In dialog.



2. If you are new to the instrument and have not yet created a local instrument profile or signed in to your Thermo Fisher Connect™ account, press **Get Started**.

Get Started

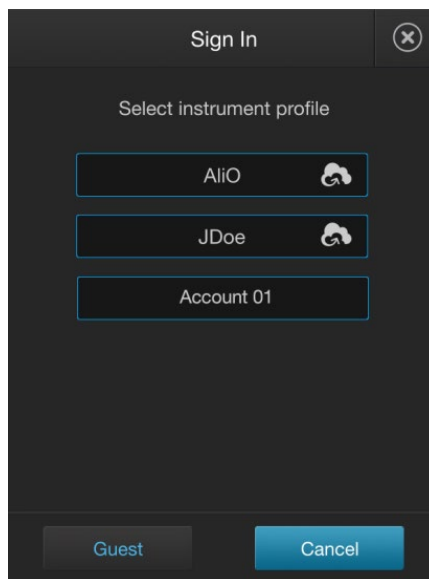
To create a local instrument profile, see page 16.

To sign in to your Thermo Fisher Connect™ account, see page 18.

Otherwise, go to Step 3 (page 25).

- Press **Screen name**, then select your instrument profile from the available options.

Screen name 



Note: The Connect icon next to a screen name indicates that the profile has an associated Connect™ account.

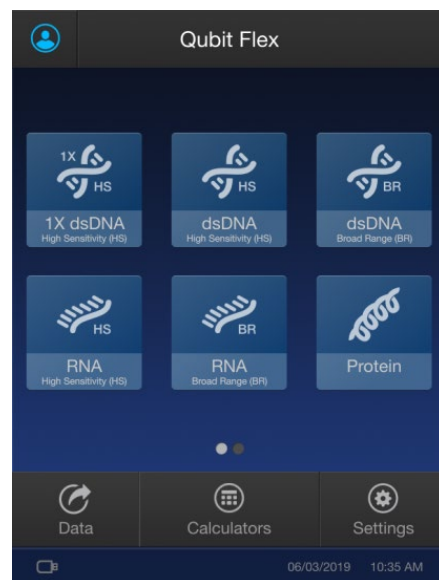
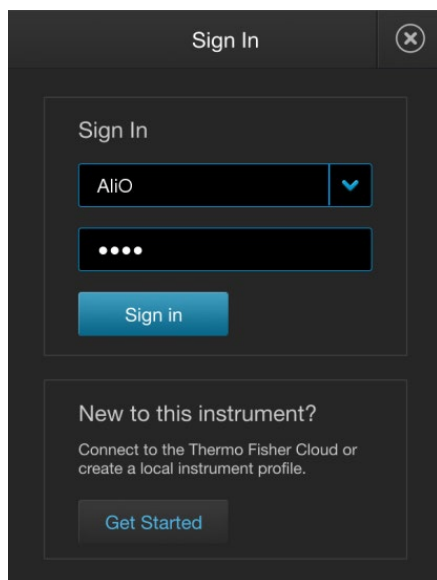
When the Connect icon is blue, the profile is signed in to the associated Connect™ account.



- Press **PIN**, enter the PIN for your profile, then press **Enter**.

PIN

- Press **Sign in** to sign in to your account and return to the Home screen. The blue profile button indicates that you have signed in to your account.

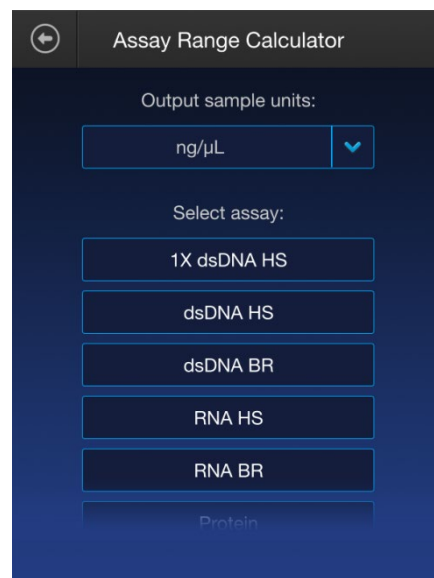
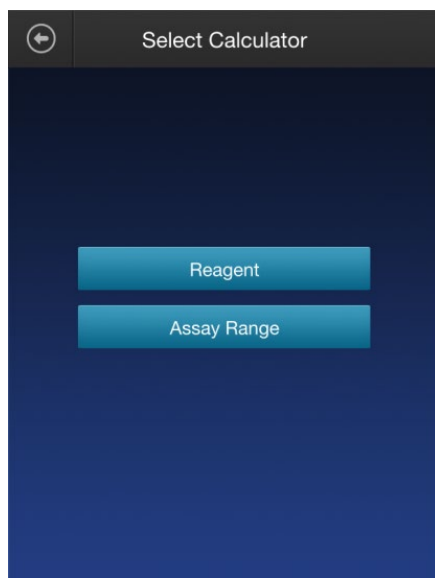
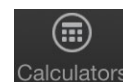


(Optional) Use the Assay Range Calculator to determine the assay range

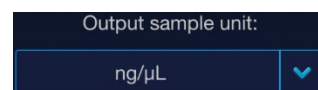
The on-board Assay Range Calculator displays the core sample concentration range for which the selected assay is most accurate, as well as the extended low and high ranges based on your sample volume. Knowing the assay range can help you determine which Qubit™ assay provides the most accurate quantification based on your sample volume and estimated sample concentration.

Use the Assay Range Calculator

1. On the **Home** screen, press **Calculators**.
2. On the **Select Calculator** screen, press **Assay Range** to open the Assay Range Calculator.



3. Press **Output sample unit**, then select the **units** in which you wish to view the assay range.



4. Select the **Assay** for which you wish to view the assay accuracy range.

The screenshot shows the 'Assay Range Calculator' interface. At the top, there is a back arrow and the title 'Assay Range Calculator'. Below this, the 'Output sample units:' are set to 'ng/μL'. The 'Select assay:' section lists several options: '1X dsDNA HS', 'dsDNA HS', 'dsDNA BR', 'RNA HS', 'RNA BR', and 'Protein'. The '1X dsDNA HS' option is currently selected.

5. Enter the **sample volume** to be used directly in the sample volume text box. You can also use the + and – buttons or adjust the sample volume wheel.

The screenshot shows the 'Assay Range Calculator' interface with the '1X dsDNA HS' assay selected. The 'Enter sample volume used:' section shows a sample volume of '15 μL'. Below this is a circular volume wheel with '+' and '-' buttons. At the bottom, there is a table showing the 'Range' and 'Sample Concentration (ng/μL)' for the selected assay. The 'Core' range is highlighted in blue, and a horizontal bar at the bottom shows the concentration range '0.01 - 6.7'.

Range	Sample Concentration (ng/μL)
Extended low	0.007 - < 0.01
Core	0.01 - 6.7
Extended high	> 6.7 - 8

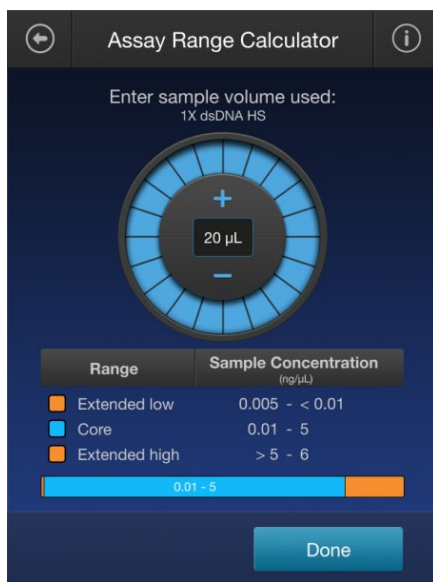
The Assay Range Calculator displays the Core sample concentration range for the selected assay and the Extended low and high ranges based on your input.

Range	Sample Concentration (ng/μL)
Extended low	0.007 - < 0.01
Core	0.01 - 6.7
Extended high	> 6.7 - 8



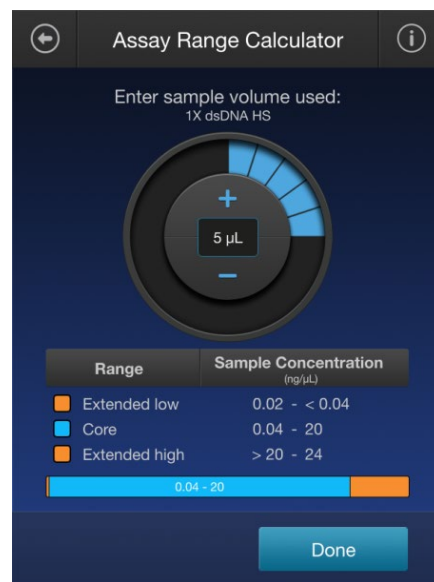
Note: Samples with concentrations within the Core range of the assay will have <15% relative error for the given sample volume. Samples with concentrations within the extended range will have <25% relative error for the given sample volume.

6. Increase or decrease the sample volume to observe how changes in the sample volume affect Core and Extended accuracy ranges for the assay.



Range	Sample Concentration (ng/µL)
Extended low	0.005 - < 0.01
Core	0.01 - 5
Extended high	> 5 - 6

dsDNA HS Assay range
for 20 µL sample volume



Range	Sample Concentration (ng/µL)
Extended low	0.02 - < 0.04
Core	0.04 - 20
Extended high	> 20 - 24

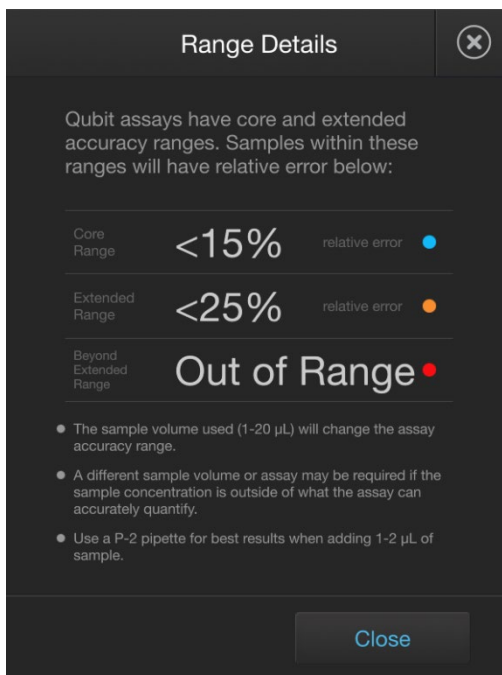
dsDNA HS Assay range
for 5 µL sample volume



Note: The sample volume used (1–20 µL) changes the assay accuracy range. For highest accuracy, use the maximum sample volume that would keep the concentration measurements within the core range.

Note that a different sample volume or assay may be required if the sample concentration is outside of what the assay can accurately quantify.

7. Press the **Information** icon on the header bar to view the Range Details (relative errors for the core and extended ranges) and guidelines for obtaining best assay results.



8. Press **Close** to return to the Assay Range Calculator.
9. (*Optional*) If desired, repeat the procedure for another assay to determine whether it would provide more accurate results in the expected concentration range.
10. Press **Done** to return to the Home screen.

Use the Reagent Calculator to prepare Qubit™ Working Solution

Use the on-board Reagent Calculator to determine the amount of Qubit™ dye and buffer required to prepare the Qubit™ Working Solution for your samples and standards.

Use the Reagent Calculator

1. On the **Select Calculator** screen, press **Reagent** to open the Reagent Calculator.

The first screenshot shows the 'Select Calculator' screen with a back arrow, the title 'Select Calculator', and two buttons: 'Reagent' and 'Assay Range'. A right-pointing arrow indicates a transition to the second screenshot.

The second screenshot shows the 'Reagent Calculator' screen. It has a back arrow and the title 'Reagent Calculator'. Below the title are two input fields: 'How many samples?' and 'How many standards?' (with a note '(0, 2 or 3 standards)'). Below these is a checkbox labeled 'Include overage' (with a note '(3 extra tubes)'). At the bottom is a numeric keypad with digits 1-9, 0, a backspace button (X), and an 'Enter' button.

2. Enter the total number of samples and standards that you plan to run.

This screenshot shows the 'Reagent Calculator' screen with the following values entered: '16' in the 'How many samples?' field, '2' in the 'How many standards?' field, and the 'Include overage' checkbox is checked. The numeric keypad is visible at the bottom.

3. (Optional) Select **Include overage**, if you want to include reagents for three additional tubes (600 µL) in the total calculated volume.

4. Press **Enter** to calculate the amount of Qubit™ dye and buffer required to prepare the Qubit™ Working Solution with these inputs.

The screenshot shows a mobile application interface titled "Reagent Calculator". At the top left is a back arrow icon. Below the title, there are two input fields: "How many samples?" with the value "16" and "How many standards?" with the value "2". Below the standards field is a checkbox labeled "Include overage (3 extra tubes)" which is checked. A horizontal line separates the inputs from the results section. The results section is titled "Results:" and contains a text box that says "Add 35µL dye to 6965µL buffer for a total volume of 7000µL". At the bottom right is a blue button labeled "Done".



Note: You can change the total number of tubes that you plan to run or the overage selection on this screen.

5. Press **Done** to return to the **Select Calculator** screen.
6. Press the **Back** button to return to the **Home screen** or press **Assay Range** to open the Assay Range Calculator (page 26).

Run standards for assay calibration

For each assay, you can run new standards to calibrate the assay on the Qubit™ Flex Fluorometer or use the values from the previous calibration. For more information, see “Qubit™ Flex Fluorometer calibration”, page 100.

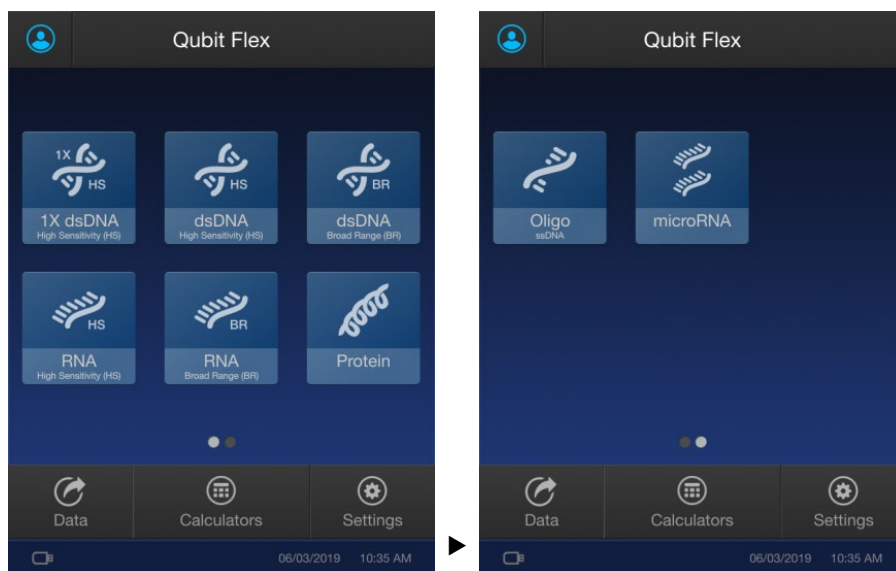


IMPORTANT! Be sure to use the appropriate standards for your assay. For best results, run new standards each time you perform an assay.

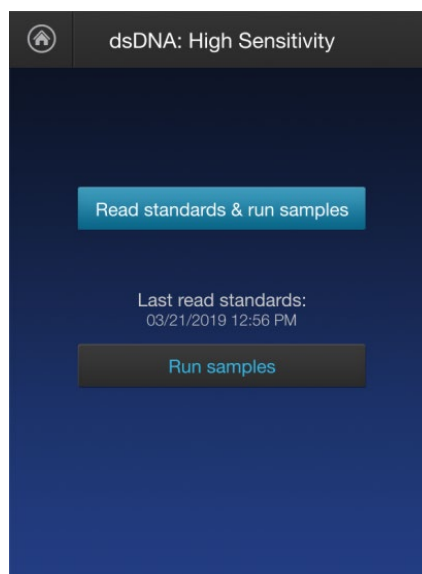
Run new standards

1. On the **Home screen**, press to select the **Assay** to perform.

To view the next screen of available assays, swipe to the left. To return to the previous page, swipe to the right.

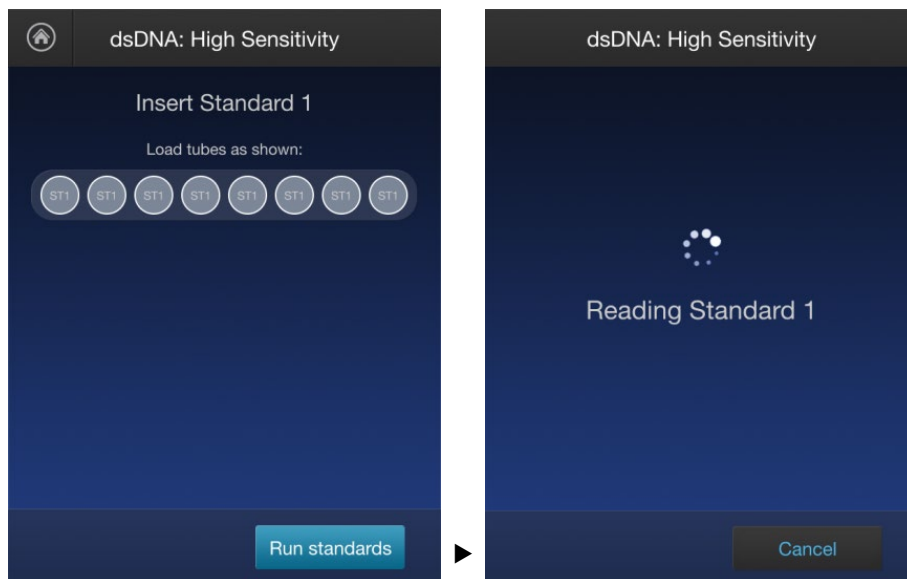


2. When prompted, press **Read standards & run samples** to read new standards.

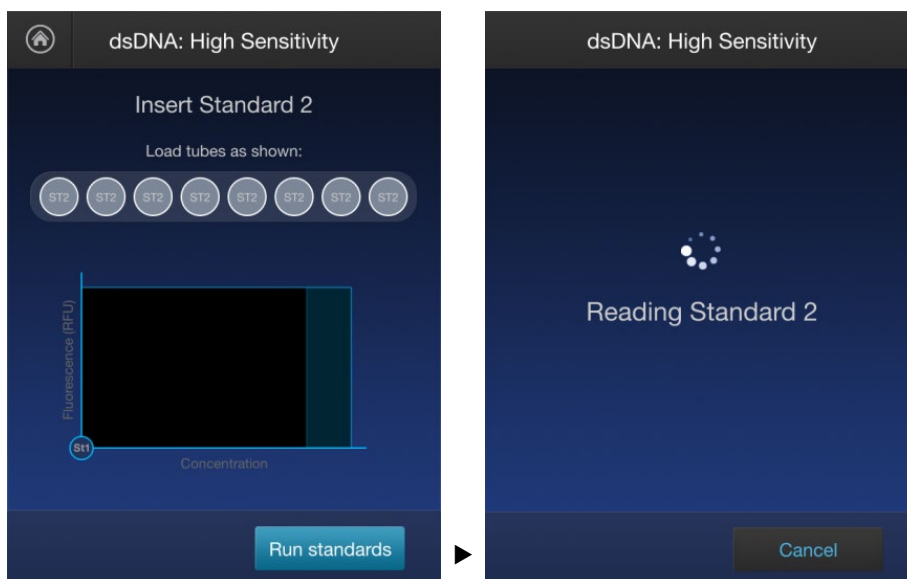


Note: To apply the previous calibration to your assay, press **Run samples**. See “Read samples”, page 36.

3. When prompted, load the Qubit™ Flex Tube Strip containing Standard #1 into the sample chamber, then press **Run standards**. The reading takes ~3 seconds.



4. When prompted, insert Standard #2, then press **Run standards**.



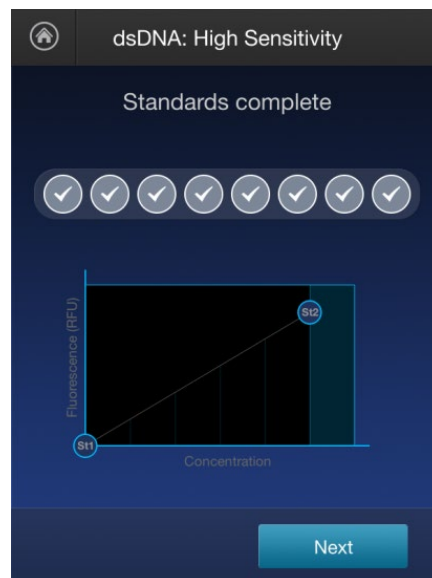
5. *For Qubit™ protein assays only:* When prompted, insert Standard #3, then press **Run standards**.

The calibration is complete after Standard #2 is read (or after Standard #3 for the Qubit™ protein assay) and the software displays the results (see “Calibration results”, page 34).

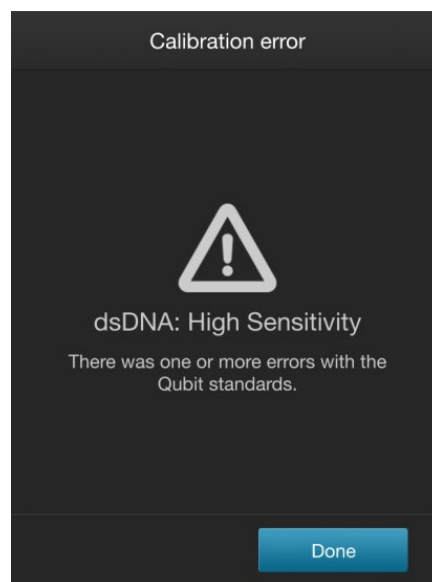
6. If your calibration is successful, press **Next** to proceed to “Read samples”, page 36.

- Calibration results**
- If the calibration is successful, **Standards complete** screen with the **Fluorescence vs. Concentration graph** is displayed.

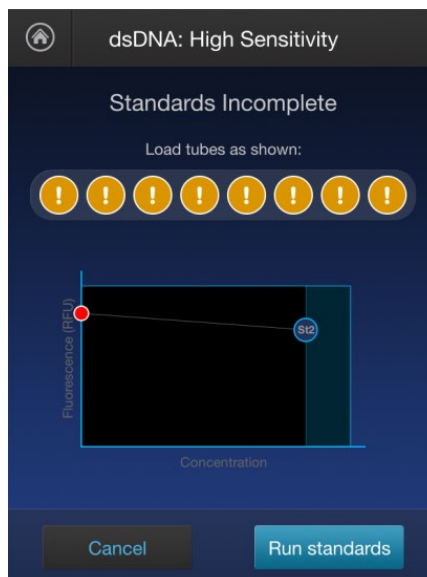
In the Fluorescence vs. Concentration graph, the standard data points are connected by a line and open circles represent correct standards.



- If the calibration is not successful, **Calibration error** message is displayed. If you receive the Calibration error message, you can re-run the standards (see “Re-run standards after calibration error”, page 35).



- (Optional) Re-run standards after calibration error**
1. In the **Calibration error** screen, press **Done**.
 2. If you wish to re-run the standards, or run new standards, prepare a fresh set of standards, then load Standard #1 into the instrument.



3. Press **Run standards**, then repeat the calibration procedure (page 25).

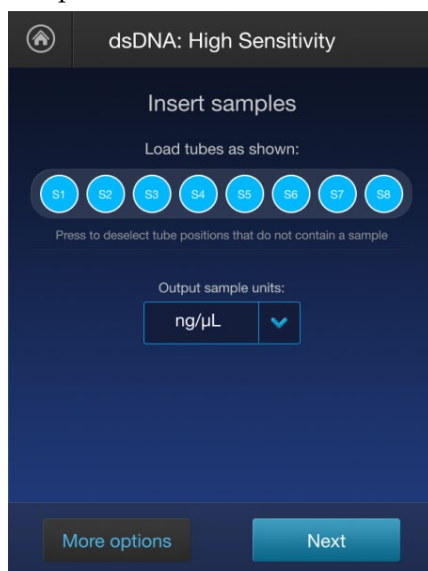
Read samples

- Before you begin**
- Calibrate the Qubit™ Flex Fluorometer as described on page 25. (Run the appropriate standards or accept the values from the previous calibration.)
 - Prepare the samples. Refer to the instructions provided with the assay.

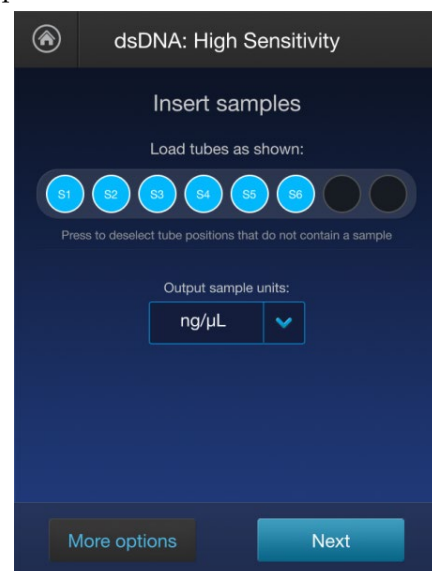


Note: Incubate the samples for the appropriate amount of time after mixing them with the working solution (2 minutes for the Qubit™ DNA and RNA assays, 15 minutes for the Qubit™ protein assay).

- Insert samples**
1. When prompted, load the tube strip containing the samples as shown in the **Insert samples** screen. If you have fewer than 8 samples, press to deselect the tube positions that do not contain a sample.

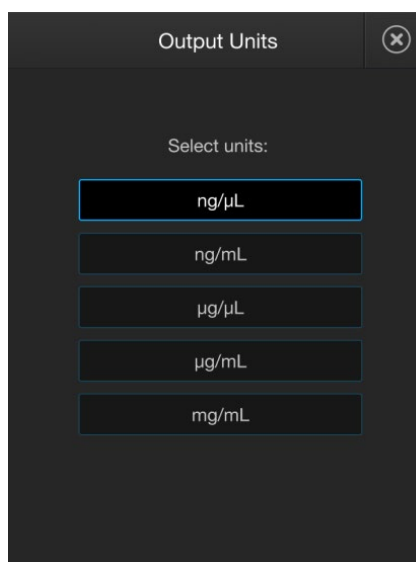
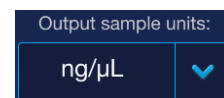


All 8 tubes contain samples



No sample in positions S7 and S8

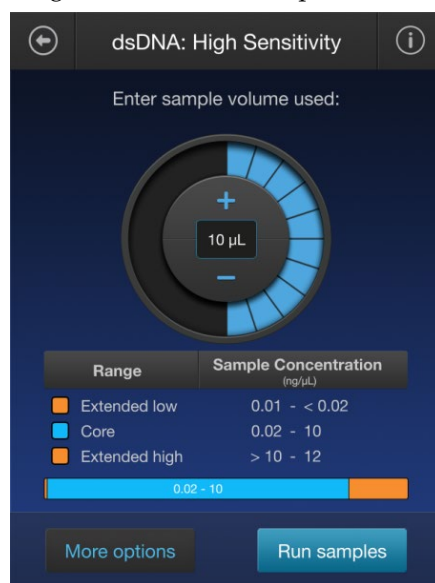
2. Press **Output sample units** to open the **Output Units** screen, then select the desired units.



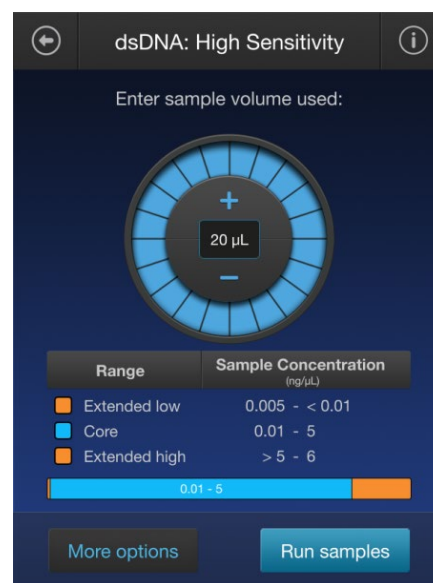
3. Press **Next** to go to the Sample volume screen.
4. In the **Sample volume** screen, enter the **sample volume** added to the assay tube (between 1 and 20 μL).

You can enter the volume directly in the sample volume text box, use the + and – buttons, or adjust the sample volume wheel.

When you enter the sample volume, the assay range information on the screen automatically changes to reflect the new core and extended accuracy ranges based on the sample volume.



dsDNA HS Assay range
for 10 μL sample volume



dsDNA HS Assay range
for 20 μL sample volume



Note: The sample volume used (1–20 μL) changes the assay accuracy range. For highest accuracy, use the maximum sample volume that would keep the concentration measurements within the core range. If the sample concentration is outside of what the assay can accurately quantify, a different sample volume or assay may be required.

(Optional) Enter
Assay kit lot #,
Add Tags,
Add Sample IDs

1. Press **More options** to open More Options screen, where you can:

- Enter assay kit lot # (Step 2, page 38)
- Add Tags to your sample run (Step 3, page 39)
- Add Sample IDs (Step 6, page 40)

The information you have entered will be available on the Data Details of your samples (page 62).

More options

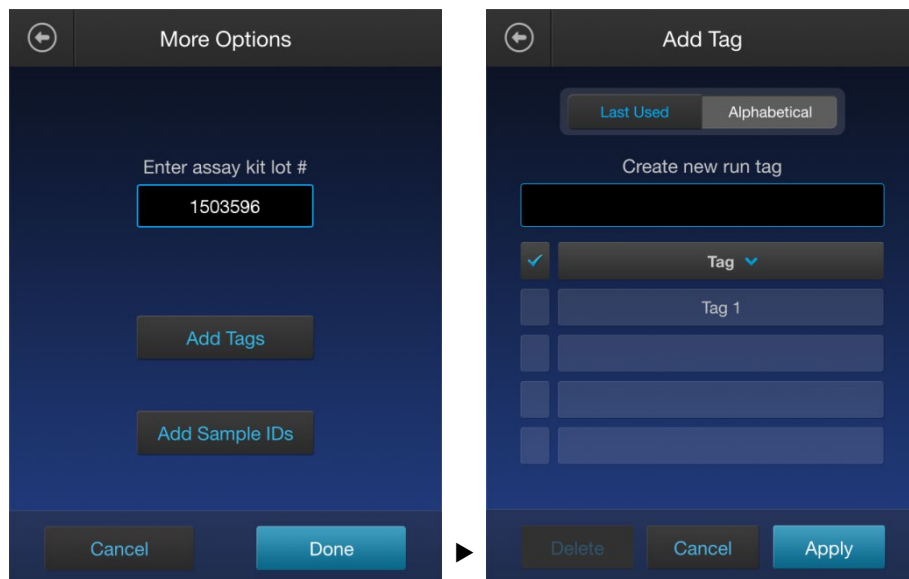


Note: You can open the More Options screen from the Insert Samples (page 36) or the Sample Volume (page 37) screens.

2. To enter an assay kit lot number, press the **Enter assay kit lot #** text box, enter the assay kit lot number, then press **Enter**.

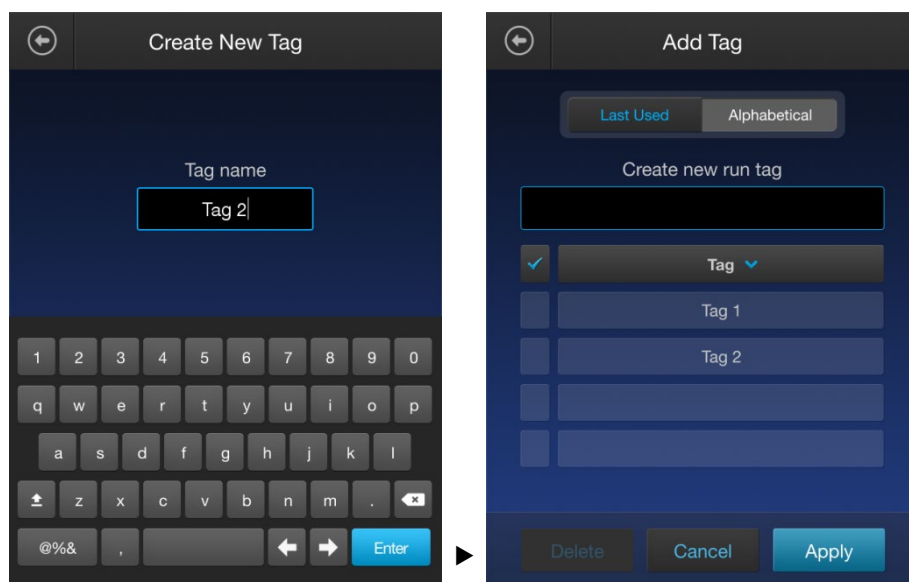
Enter assay kit lot #

- To add a tag to your samples in the run, press **Add Tags** on the More Options screen to open the Add Tag screen.



- To create a new tag, press the **Create new run tag** text box to open the Create New Tag screen, enter the new tag, then press **Enter**.

The new tag will be added to the list of available tags on the Add Tag screen.

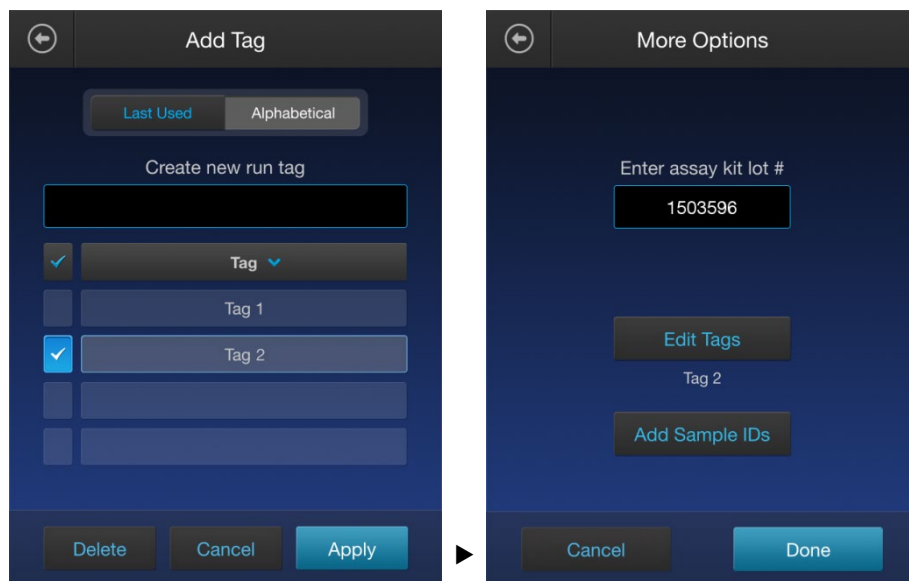


Note: To filter the list of available tags for the last used tag, press **Last Used**.

To display all existing tags alphabetically, press **Alphabetical**.

To sort the list of available tags alphabetically in ascending or descending order, press the Tag column header.

5. Select the desired tag from the list of available tags, then press **Apply** to add the selected tag to your samples and return to the More Options screen. The tag you have applied to your sample run is displayed on the More Options screen and the Add Tags button changes to Edit Tags.

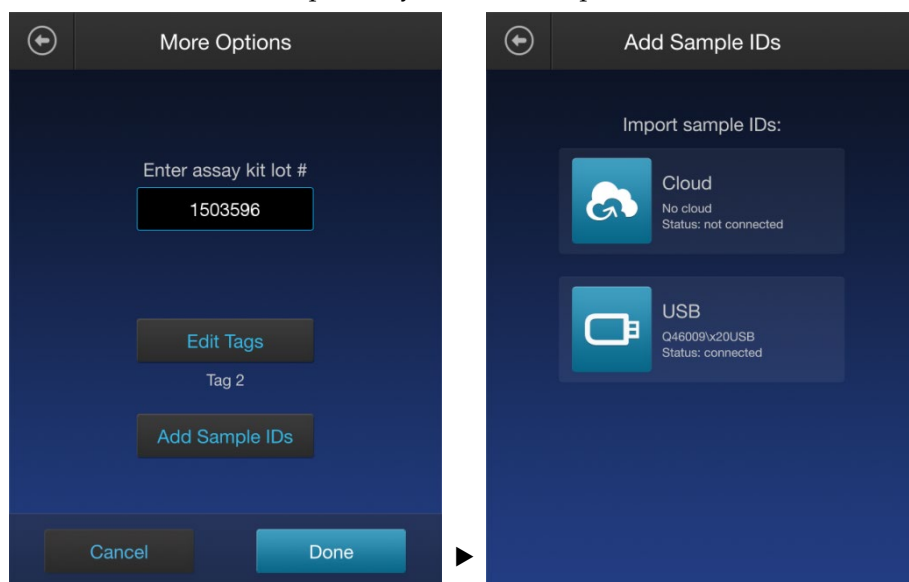


Note: To return to the More Options screen without applying a tag to your samples, press **Cancel**.

To delete an existing tag, select the tag from the list of available tags, then press **Delete**.

To change the tag applied to your sample run, press **Edit Tags** on the More Options screen.

6. To add sample IDs to your samples, press **Add Samples IDs**, then select **Cloud** (your Connect™ account; see page 18 for sign in instructions) or **USB** for the location of the sample IDs you want to import.





Note: The file containing the sample IDs must be in CSV (comma separated value) format and filled out like the example below: first “Plate Barcode” then “Well” and “Sample Id”.

	A	B	C
1	Plate Barcode	Well	Sample Id
2	96W207	A1	GS2072-2
3	96W208	A2	GS2072-3
4	96W209	A3	GS2072-4

7. Select the file containing the sample IDs from the list of available files, then press **Apply**.

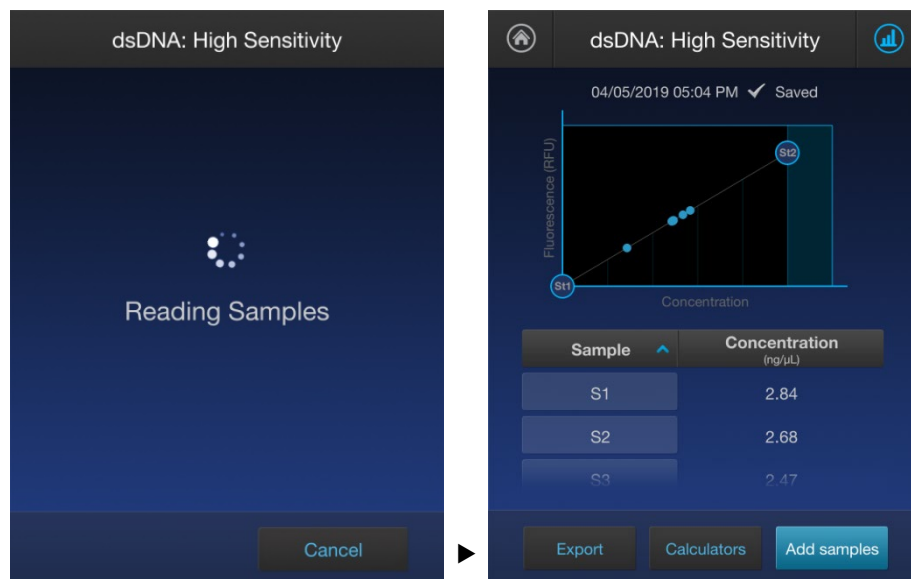
The left screenshot shows the 'Add Sample IDs' screen with a table containing one row: 23/09/2019, Sample ID.csv, 8. The right screenshot shows the same screen with a table containing 8 rows of sample information.

Sample	Sample ID	Plate ID	Well
S1	GS2072-2	96W207	A1
S2	GS2072-3	96W208	A2
S3	GS2072-4	96W209	A3
S4	GS2072-5	96W210	A4
S5	GS2072-6	96W211	A5
S6	GS2072-7	96W212	A6
S7	GS2072-8	96W213	A7
S8	GS2072-9	96W214	A8

8. Press **OK** at the confirmation page.
9. When finished entering assay kit lot number and applying tags and sample IDs, press **Done** at the More Options screen. The assay screen displays the new information added to your samples at the bottom of the screen.
To go back to the assay screen without applying the new information, press **Cancel**.

The left screenshot shows the 'Add Sample IDs' screen with a large checkmark and the text 'Sample IDs successfully applied'. The right screenshot shows the 'dsDNA: High Sensitivity' screen with the 'Insert samples' section, showing 8 sample positions (S1-S8) and a dropdown menu for 'Output sample units' set to 'ng/μL'.

- Run Samples**
1. Press **Run samples**. The reading takes approximately 3 seconds and the results are displayed in graph view in the Results screen (see “Results”, page 43).



2. To display the results in list view, press the **Graph** button to unselect it. The Results screen lists the concentration of each original sample using the output units selected at the beginning of the assay.

Sample	Concentration (ng/μL)
S1	2.84
S2	2.68
S3	2.47
S4	2.43
S5	2.47
S6	2.45
S7	2.45
S8	1.43



Note: By default, the Results screen displays the measurements in graph view. However, the graph settings are “sticky”, so that if you close the graph, the next time anyone runs an assay, the graph view is hidden and the results are shown in list form.

3. To run more samples, press **Add samples**, and repeat the procedure.

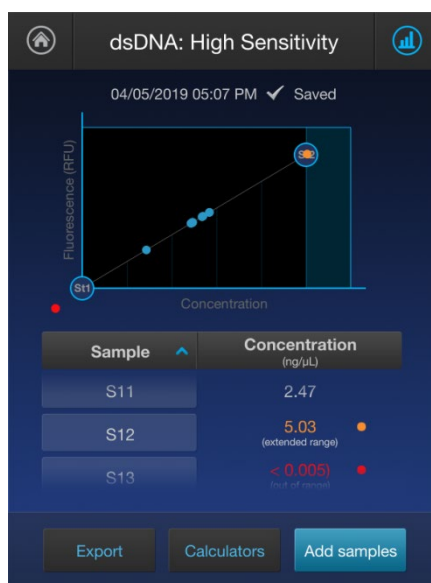
Results

- View results** 1. The instrument automatically displays the Results screen after the completion of each sample run.

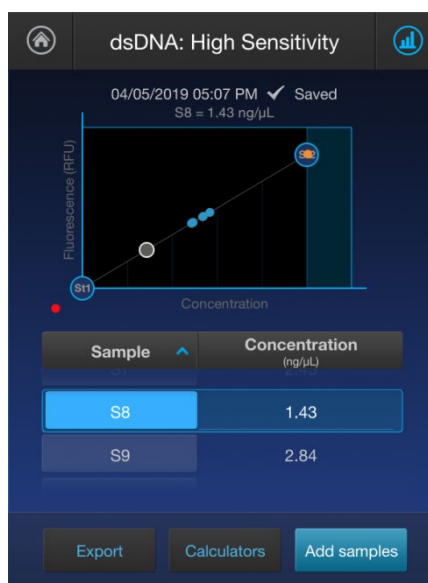
By default, the results are displayed in graph view, which shows the Fluorescence vs. Concentration graph and lists the concentration of each original sample below the graph.

In the graph:

- Open circles represent correct standards.
- Blue circles represent samples that fall within the assay's core range.
- Orange circles represent samples that fall within the assay's extended range.
- Red circles represent samples that fall outside the assay's range.



2. To view a sample on the Fluorescence vs. Concentration graph, press the desired sample on the sample list. The selected sample is displayed as a gray circle on the graph.



3. To display the results in list view, press the **Graph** button to hide the graph.



The Results screen shows the concentration of each original sample in a list form, using the output units selected at the beginning of the assay.

Sample	Concentration (ng/μL)
S1	2.84
S2	2.68
S3	2.47
S4	2.43
S5	2.47
S6	2.45
S7	2.45
S8	1.43

Sample	Concentration (ng/μL)
S7	2.45
S8	1.43
S9	2.84
S10	2.7
S11	2.47
S12	5.03 (extended range)
S13	< 0.005 (out of range)
S14	< 0.005 (out of range)

- If the concentration of a sample is within the assay's extended range, the concentration value is displayed in orange, and an "extended range" message and an orange circle are displayed next to the concentration value.
 - If the concentration of a sample is outside of the assay's range, an "out of range" message and a red circle are displayed next to the sample.
4. To display the results in graph view again, press the **Graph** button.



(Optional) Use the Molarity Calculator to determine sample molarity

The on-board Molarity Calculator allows you to calculate the molarity of your samples based on nucleic acid length and their measured concentration.

Use the Molarity Calculator

1. On the **Results** screen, press **Calculators**, then select **Molarity** to open the Molarity Calculator.

Calculators

Calculator

Select a calculator:

Molarity

Normalization

Molarity Calculator

Desired units: to

Molecular weight: g/mol

☐ Auto-populate DNA length

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44	<input type="text"/>	<input type="text"/>
S2	2.44	<input type="text"/>	<input type="text"/>
S3	2.57	<input type="text"/>	<input type="text"/>
S4	2.46	<input type="text"/>	<input type="text"/>

Export Cancel Calculate

2. On the **Molarity Calculator** screen, press the **Desired units** fields to select the **input** and **output** units.

Desired units: to

Units Selection

Choose the units you want to use

Input units

Units Selection

Choose the units you want to use

Output units



Note: The Qubit™ Flex Fluorometer auto-populates the Molecular weight (MW) depending on the Qubit™ assay performed (for example, for the dsDNA HS assay, it uses a default value of 660 g/mol for the average molecular weight of one DNA base pair).

Molecular weight: 660 g/mol

To change the auto-populated MW value, press the **Molecular weight** field and enter the desired average molecular weight of your sample.

3. Press **Length (bp)** field for Sample 1 (S1), enter the length (bp) of Sample 1, then press **Enter**.

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44		

← Molarity Calculator

Desired units: ng/μL to nM

Molecular weight: 660 g/mol

☐ Auto-populate DNA length

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44	200	
S2	2.44		
S3	2.57		
S4	2.46		

Export Cancel Calculate

4. If all your samples have the same length, select **Auto-populate DNA length**.

☒ Auto-populate DNA length

← Molarity Calculator

Desired units: ng/μL to nM

Molecular weight: 660 g/mol

☒ Auto-populate DNA length

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44	200	
S2	2.44	200	
S3	2.57	200	
S4	2.46	200	

Export Cancel Calculate

- Press **Calculate** to calculate the molarity of your samples based on the assay results and DNA length in the output units that you have selected.

Calculate

Molarity Calculator

Desired units: to

Molecular weight: g/mol

☒ Auto-populate DNA length

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44	<input type="text" value="200"/>	18.5
S2	2.44	<input type="text" value="200"/>	18.5
S3	2.57	<input type="text" value="200"/>	19.5
S4	2.46	<input type="text" value="200"/>	18.6

Sample	Concentration (ng/μL)	Length (bp)	Molarity (nM)
S1	2.44	<input type="text" value="200"/>	18.5
S2	2.44	<input type="text" value="200"/>	18.5
S3	2.57	<input type="text" value="200"/>	19.5
S4	2.46	<input type="text" value="200"/>	18.6



Note: When you press Calculate, the instrument saves the data from molarity calculations with the sample data in the CSV file.

- To export your results, press **Export**. The instrument exports the complete CSV file with all sample data, including the molarity calculation results.
To go back to the Calculator screen, press the **Back** button.

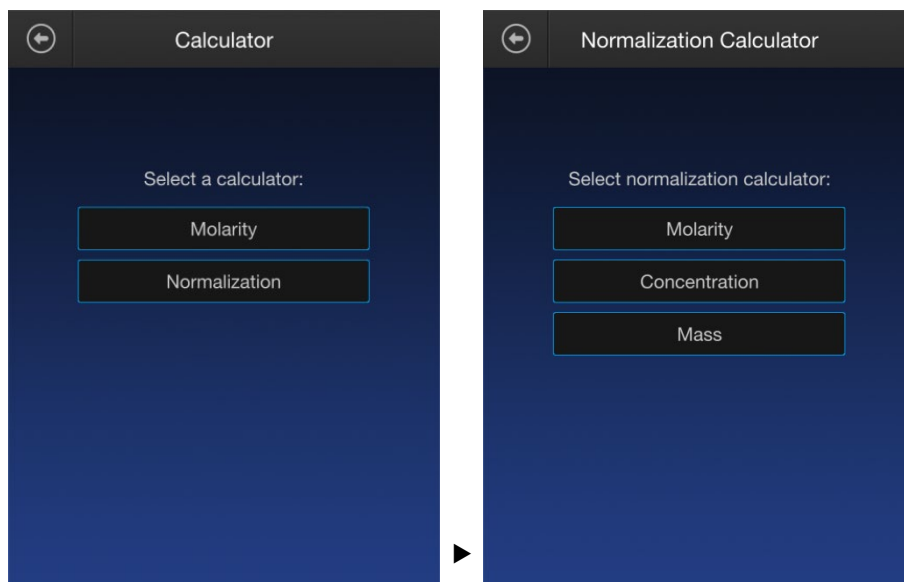
(Optional) Use the Normalization Calculator to determine how to dilute the samples to the same molarity, concentration, or mass

The on-board Normalization Calculator helps you to normalize your samples of variable concentration to the same molarity, concentration, or mass using the results from your assay.

Select the Normalization Calculator

1. On the **Results** screen, press **Calculators**, then press **Normalization**.

Calculators



2. On the **Normalization Calculator** screen, select:
 - **Molarity** to determine how to dilute your samples to the same final mass and volume (page 49).
 - **Concentration** to determine how to dilute your samples to the same final concentration (page 52).
 - **Mass** to determine how to dilute your samples to the same final mass and volume (page 55).



Note: The option to normalize your samples based on molarity is available only if you have run the Molarity calculator (page 45) on your samples.

Normalize your samples to the same molarity

1. On the **Normalization Calculator** screen, select **Molarity**.

The first screenshot shows the 'Normalization Calculator' screen with the prompt 'Select normalization calculator:'. Below this prompt are three buttons: 'Molarity', 'Concentration', and 'Mass'. The second screenshot shows the same screen after 'Molarity' has been selected. The 'Final sample molarity:' field now contains 'nM' and the 'Final sample volume:' field contains 'μL'. A numeric keypad is visible at the bottom of the second screenshot.

2. Enter the **Final sample mass** and select **units**.
3. Enter the **Final sample volume** and select **units**, then press **Enter**.

The third screenshot shows the 'Normalization Calculator' screen with the 'Final sample molarity:' field set to 'nM' and the 'Final sample volume:' field set to '10 μL'. The fourth screenshot shows the same screen with a 'Calculate' button appearing below the input fields.



Note: The minimum allowed sample volume on the Normalization Calculator is 5 μL .

- Press **Calculate** to see how much sample and buffer to mix to achieve the desired final sample concentration and volume.

Calculate

Normalization Calculator

Final Molarity: 5 nM Final Volume: 10 μ L

Sample	Add sample (μ L)	Add buffer (μ L)
S1	2.7	7.3
S2	2.7	7.3
S3	2.6	7.4
S4	2.7	7.3
S5	2.7	7.3
S6	2.7	7.3
S7	2.7	7.3
S8	2.7	7.3

Page 1 of 3

Export Done

Final Molarity: 5 nM Final Volume: 10 μ L

Sample	Add sample (μ L)	Add buffer (μ L)
S1	2.7	7.3
S2	2.7	7.3



Note: When you press Calculate, the instrument saves the data from normalization calculations with the sample data in the CSV file.

- Press the **right arrow** to view page 2 of results, which displays the required sample:buffer dilution before mixing ("Required Dilution", if applicable) and the sample concentration after the dilution ("Diluted conc.").
If dilution is not required before mixing, then "N/A" is displayed in the Required Dilution and Diluted conc. columns for the sample.

Normalization Calculator

Final Molarity: 5 nM Final Volume: 10 μ L

Sample	Required Dilution (sample:buffer)	Diluted conc. (nM)
S1	N/A	N/A
S2	N/A	N/A
S3	N/A	N/A
S4	N/A	N/A
S5	N/A	N/A
S6	N/A	N/A
S7	N/A	N/A
S8	N/A	N/A

Page 2 of 3

Export Done

- Press the **right arrow** again to view page 3, which displays the actual sample concentration (“Concentration”).

Normalization Calculator	
Final Molarity: 5 nM Final Volume: 10 µL	
Sample	Concentration (nM)
S1	18.5
S2	18.5
S3	19.5
S4	18.6
S5	18.6
S6	18.7
S7	18.6
S8	18.4

Page 3 of 3

Export Done

- Press the **left arrow** to go back to the previous page.
- To export your calculations, press **Export**. The instrument exports the complete CSV file with all sample data, including the normalization calculation results.
- Otherwise, press **Done** to close the Normalization calculator and go back to the Calculator screen.



Note: If your sample needs further dilution before mixing to achieve the desired final molarity, the required sample:buffer dilution is indicated in the Add sample column (in red) and in the Required Dilution column (on page 1 and 2 of calculation results, respectively).

Final Molarity: 100 pM Final Volume: 20 µL		
Sample	Add sample (µL)	Add buffer (µL)
S1	1.1 (1:9)	18.9
S2	1.1 (1:9)	18.9

Final Molarity: 100 pM Final Volume: 20 µL		
Sample	Required Dilution (sample:buffer)	Diluted conc. (pM)
S1	1:9	1850
S2	1:9	1850

If dilution is not required before mixing, then the Required Dilution and Diluted conc. columns display “N/A” for the sample.

Normalize your samples to the same concentration

1. On the **Normalization Calculator** screen, select **Concentration**.

The first screenshot shows the 'Normalization Calculator' app with the option 'Concentration' selected. The second screenshot shows the input fields for 'Final sample concentration' (set to 1 ng/μL) and 'Final sample volume' (set to 200 μL), with a numeric keypad below.

2. Enter the **Final sample concentration** and select **units**.
3. Enter the **Final sample volume** and select **units**, then press **Enter**.

The first screenshot shows the input fields for 'Final sample concentration' (set to 1 ng/μL) and 'Final sample volume' (set to 200 μL). The second screenshot shows the 'Calculate' button.



Note: The minimum allowed sample volume on the Normalization Calculator is 5 μL.

- Press **Calculate** to see how much sample and buffer to mix to achieve the desired final sample concentration and volume.

Calculate

Normalization Calculator

Final Concentration: 1 ng/μL Final Volume: 200 μL

Sample	Add sample (μL)	Add buffer (μL)
S1	82.0	118
S2	82.0	118
S3	77.8	122
S4	81.3	119
S5	81.6	118
S6	81.0	119
S7	81.6	118
S8	82.3	118

Page 1 of 3

Export Done

Final Concentration: 1 ng/μL Final Volume: 200 μL

Sample	Add sample (μL)	Add buffer (μL)
S1	82.0	118
S2	82.0	118



Note: When you press Calculate, the instrument saves the data from normalization calculations with the sample data in the CSV file.

- Press the **right arrow** to view page 2 of results, which displays the required sample:buffer dilution before mixing ("Required Dilution", if applicable) and the sample concentration after the dilution ("Diluted conc.").
If dilution is not required before mixing, then "N/A" is displayed in the Required Dilution and Diluted conc. columns for the sample.

Normalization Calculator

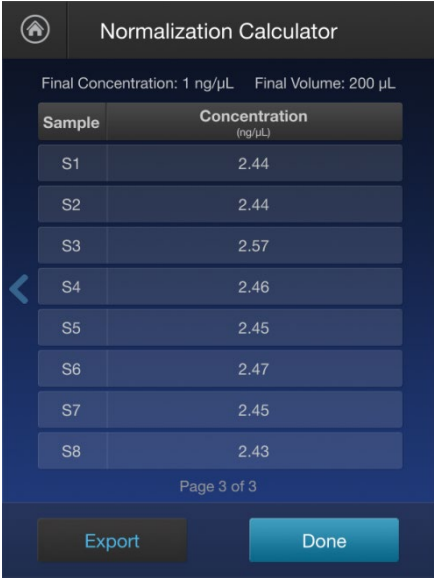
Final Concentration: 1 ng/μL Final Volume: 200 μL

Sample	Required Dilution (sample:buffer)	Diluted conc. (ng/μL)
S1	N/A	N/A
S2	N/A	N/A
S3	N/A	N/A
S4	N/A	N/A
S5	N/A	N/A
S6	N/A	N/A
S7	N/A	N/A
S8	N/A	N/A

Page 2 of 3

Export Done

6. Press the **right arrow** again to view page 3 of results, which displays the actual sample concentration (“Concentration”).



Final Concentration: 1 ng/μL Final Volume: 200 μL	
Sample	Concentration (ng/μL)
S1	2.44
S2	2.44
S3	2.57
S4	2.46
S5	2.45
S6	2.47
S7	2.45
S8	2.43

Page 3 of 3

Export Done

7. Press the **left arrow** to go back to the previous page.
8. To export your calculations, press **Export**. The instrument exports the complete CSV file with all sample data, including the normalization calculation results.
9. Otherwise, press **Done** to close the Normalization calculator and go back to the Calculator screen.



Note: If your sample needs further dilution before mixing to achieve the desired final concentration, the required sample:buffer dilution is indicated in the “Add sample” column (in red) and in the “Required Dilution” column (on page 1 and 2 of calculation results, respectively).

Final Concentration: 2 ng/mL Final Volume: 200 μL		
Sample	Add sample (μL)	Add buffer (μL)
S1	1.1 (1:6)	199
S2	1.1 (1:6)	199

Final Concentration: 2 ng/mL Final Volume: 200 μL		
Sample	Required Dilution (sample:buffer)	Diluted conc. (ng/mL)
S1	1:6	349
S2	1:6	349

If dilution is not required before mixing, then the Required Dilution and Diluted conc. columns display “N/A” for the sample.

Normalize your samples to the same mass and volume

1. On the **Normalization Calculator** screen, select **Mass**.

The first screenshot shows the 'Normalization Calculator' app with the option 'Mass' selected. The second screenshot shows the input fields for 'Final sample mass' (1 ng) and 'Final sample volume' (10 µL).

2. Enter the **Final sample mass** and the desired **units**.
3. Enter the **Final sample volume** and the desired **units**, then press **Enter**.

The first screenshot shows the input fields for 'Final sample mass' (1 ng) and 'Final sample volume' (10 µL). The second screenshot shows the 'Calculate' button.



Note: The minimum allowed sample volume on the Normalization Calculator is 5 µL.

- Press **Calculate** to see how much sample and buffer to mix to achieve the desired final sample concentration and volume.

Calculate

Normalization Calculator

Final Mass: 1 ng Final Volume: 10 μ L

Sample	Add sample (μ L)	Add buffer (μ L)
S1	1.2 (1:2)	8.8
S2	1.2 (1:2)	8.8
S3	1.2 (1:2)	8.8
S4	1.2 (1:2)	8.8
S5	1.2 (1:2)	8.8
S6	1.2 (1:2)	8.8
S7	1.2 (1:2)	8.8
S8	1.2 (1:2)	8.8

Page 1 of 3

Export Done

Final Mass: 1 ng Final Volume: 10 μ L

Sample	Add sample (μ L)	Add buffer (μ L)
S1	1.2 (1:2)	8.8
S2	1.2 (1:2)	8.8



Note: When you press Calculate, the instrument saves the data from normalization calculations with the sample data in the CSV file.

- Press the **right arrow** to view page 2 of results, which displays the required sample:buffer dilution before mixing ("Required Dilution", if applicable) and the sample concentration after the dilution ("Diluted conc.").
If dilution is not required before mixing, then "N/A" is displayed in the Required Dilution and Diluted conc. columns for the sample.

Normalization Calculator

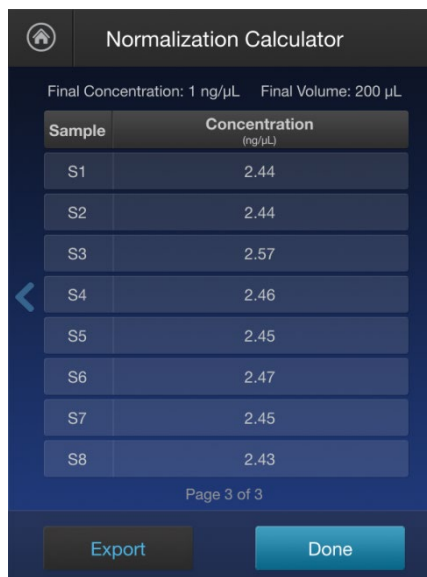
Final Mass: 1 ng Final Volume: 10 μ L

Sample	Required Dilution (sample:buffer)	Diluted conc. (ng/ μ L)
S1	1:2	0.813
S2	1:2	0.813
S3	1:2	0.857
S4	1:2	0.82
S5	1:2	0.817
S6	1:2	0.823
S7	1:2	0.817
S8	1:2	0.81

Page 2 of 3

Export Done

- Press the **right arrow** again to view page 3 of calculation results, which displays the actual sample concentration (“Concentration”).



Normalization Calculator	
Final Concentration: 1 ng/μL Final Volume: 200 μL	
Sample	Concentration (ng/μL)
S1	2.44
S2	2.44
S3	2.57
S4	2.46
S5	2.45
S6	2.47
S7	2.45
S8	2.43

Page 3 of 3

Export Done

- Press the **left arrow** to go back to the previous page.
- To export your calculations, press **Export**. The instrument exports the complete CSV file with all sample data, including the normalization calculation results. Otherwise, press **Done** to close the Normalization calculator and go back to the Calculator screen.



Note: If your sample needs further dilution before mixing to achieve the desired final mass and volume, the required sample:buffer dilution is indicated in the “Add sample” column (in red) and in the “Required Dilution” column (on page 1 and 2 of calculation results, respectively).

Final Mass: 1 ng Final Volume: 10 μL		
Sample	Add sample (μL)	Add buffer (μL)
S1	1.2 (1:2)	8.8
S2	1.2 (1:2)	8.8

Final Mass: 1 ng Final Volume: 10 μL		
Sample	Required Dilution (sample:buffer)	Diluted conc. (ng/μL)
S1	1:2	0.813
S2	1:2	0.813

If dilution is not required before mixing, then the Required Dilution and Diluted conc. columns display “N/A” for the sample.

4. Manage data

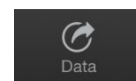
Overview The Qubit™ Flex Fluorometer can save data for up to 10,000 samples.

For the saved data, the Qubit™ Flex Fluorometer allows you to:

- View detailed data for each sample (page 58).
- Rename data files (page 63).
- Export data as a CSV (comma separated value) file to a USB drive, directly to your computer, or to your Connect™ account (page 64).
- Delete data files (page 70).

View detailed sample data

- View list of data sets**
1. On the **Home** screen, press **Data**. The Data screen opens and displays the list of data sets that are saved in the instrument.



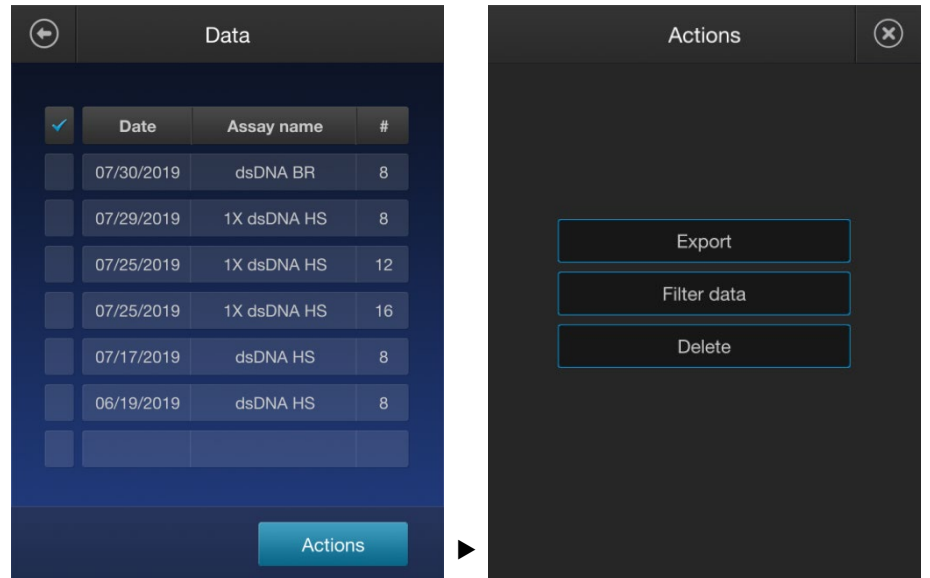
The screenshot shows the 'Data' screen of the Qubit Flex Fluorometer. At the top is a header bar with a back arrow icon on the left and the word 'Data' in the center. Below the header is a table with four columns: a selection column with checkboxes, 'Date', 'Assay name', and '#'. The table contains seven rows of data, with the last row being empty. At the bottom of the screen is a blue button labeled 'Actions'.

<input checked="" type="checkbox"/>	Date	Assay name	#
<input type="checkbox"/>	07/30/2019	dsDNA BR	8
<input type="checkbox"/>	07/29/2019	1X dsDNA HS	8
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	12
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	16
<input type="checkbox"/>	07/17/2019	dsDNA HS	8
<input type="checkbox"/>	06/19/2019	dsDNA HS	8
<input type="checkbox"/>			

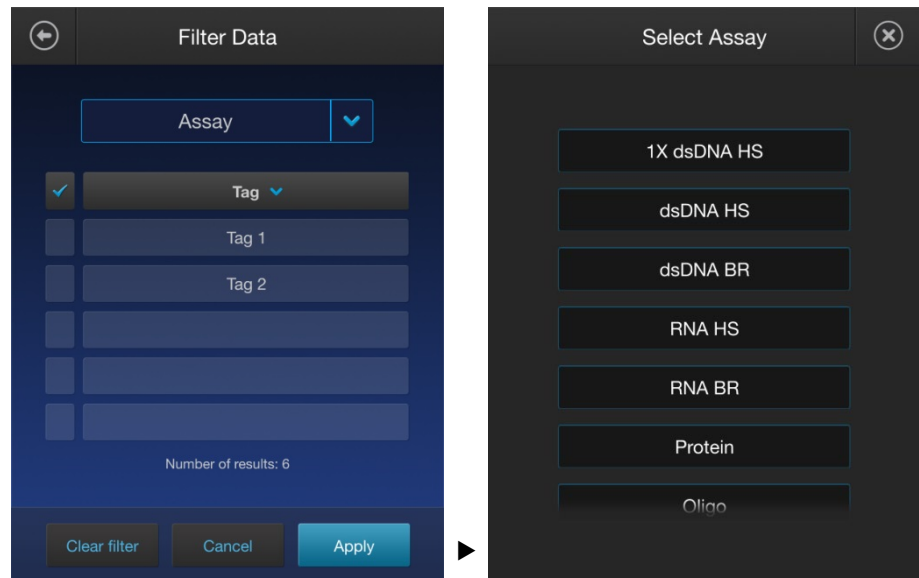
2. By default, the data sets are arranged by date in descending order. To sort the data sets, press the appropriate category in the header row:
 - To sort the data sets by date in ascending order, press **Date**.
To sort the data sets by date in descending order, press **Date** again.
 - To sort the data sets by Assay name in descending order, press **Assay name**.
To sort the data sets by Assay name in ascending order, press **Assay name** again.
 - To sort the data sets by the number of samples in descending order, press **#**.
To sort the data sets by the number of samples in ascending order, press **#** again.

(Optional) Filter data sets

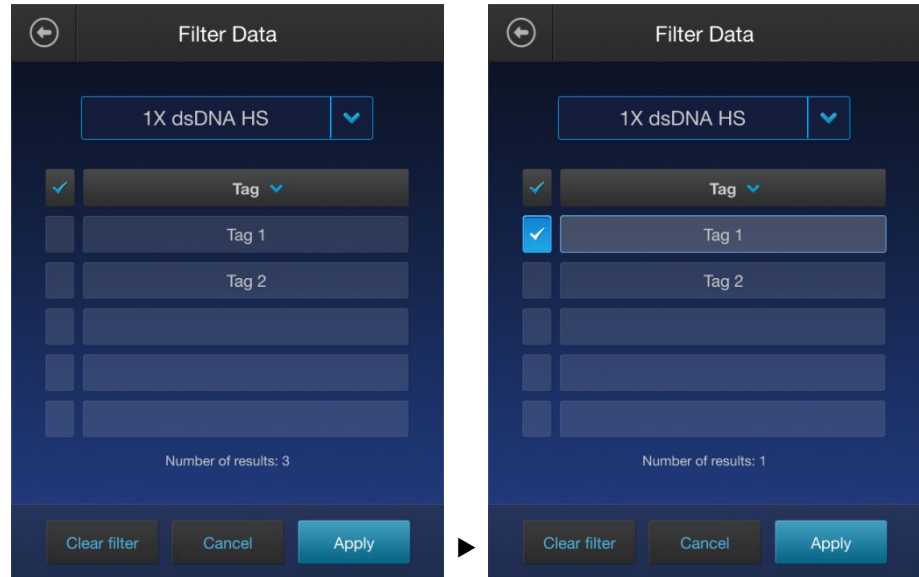
1. To filter data sets by Assay or Tag, press **Actions** to open the Actions screen, then select **Filter data**.



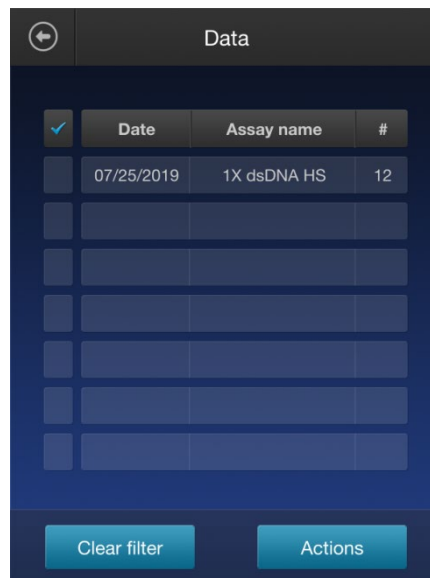
2. On the Filter Data screen, press **Assay**, then select the Assay of interest.



3. If you had applied a tag to the assay (page 38), select the **Tag** from the list. Otherwise, go to step 4.

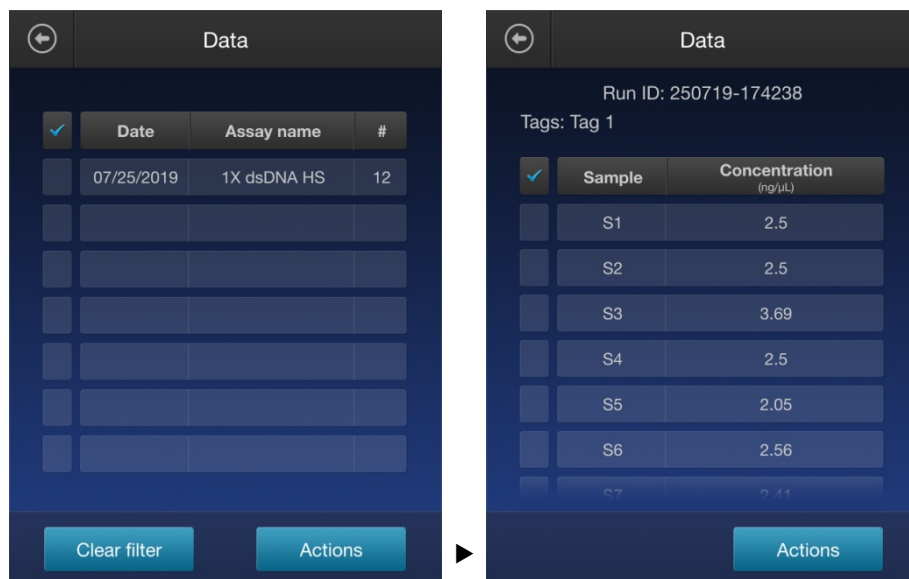


4. Press **Apply** to filter the data list by the assay and tag you have selected. Only the data sets that satisfy the filter criteria are displayed in the Data screen.

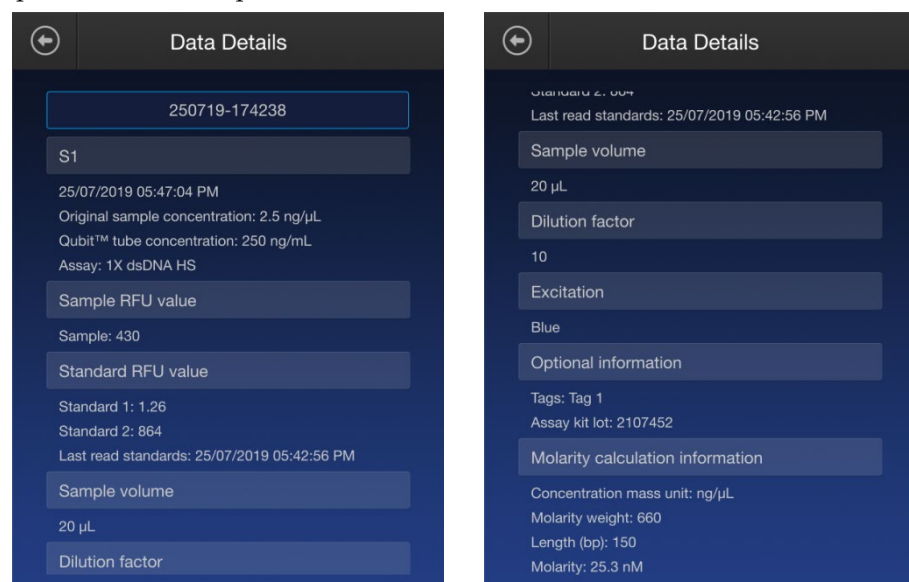


Select data set and view detailed sample data

1. On the Data screen (filtered or not filtered), press the **data set of interest**. The Data set screen opens and displays a list of samples in that run.



2. To view the sample details, press the **sample of interest**. A Data details screen opens. To view sample details that do not fit in the screen, scroll down.



Information in the detailed sample data

←

Data Details

①

250719-174238

②

S1

③

25/07/2019 05:47:04 PM

④

Original sample concentration: 2.5 ng/μL

⑤

Qubit™ tube concentration: 250 ng/mL

⑥

Assay: 1X dsDNA HS

⑦

Sample RFU value

⑧

Sample: 430

⑧

Standard RFU value

⑧

Standard 1: 1.26

⑧

Standard 2: 864

⑨

Last read standards: 25/07/2019 05:42:56 PM

⑩

Sample volume

⑩

20 μL

⑩

Dilution factor

←

Data Details

Standard 2: 864

Last read standards: 25/07/2019 05:42:56 PM

Sample volume

20 μL

Dilution factor

⑪

10

Excitation

⑫

Blue

Optional information

⑬

Tags: Tag 1

Assay kit lot: 2107452

Molarity calculation information

⑭

Concentration mass unit: ng/μL

Molarity weight: 660

Length (bp): 150

Molarity: 25.3 nM

- ① Run ID

② Sample name

③ Assay date

④ Original sample concentration

⑤ Qubit™ tube sample concentration

⑥ Assay name

⑦ Sample RFU* value
- ⑧ RFU values for the standards

⑨ Date of last read standards

⑩ Sample volume

⑪ Dilution factor

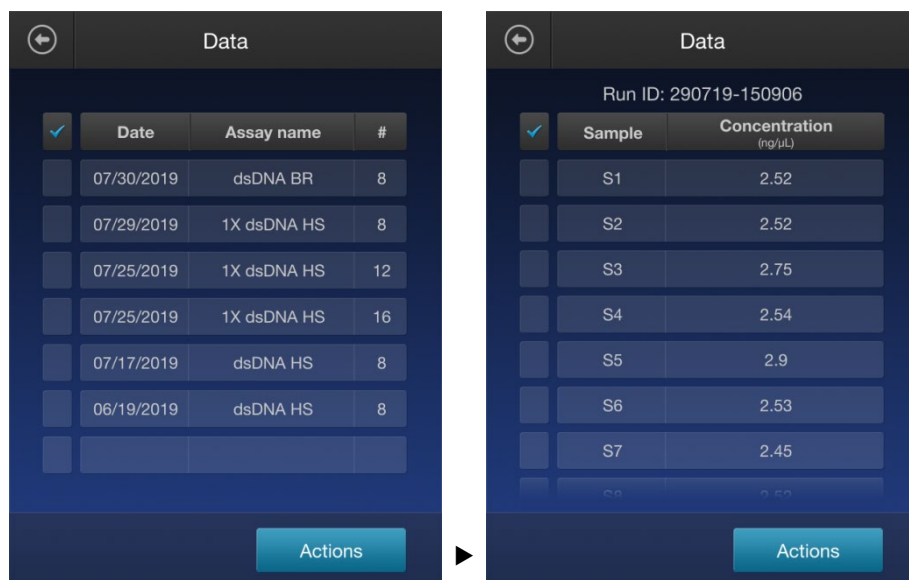
⑫ Excitation channel

⑬ Optional information (Tags, Reagent lot etc.)

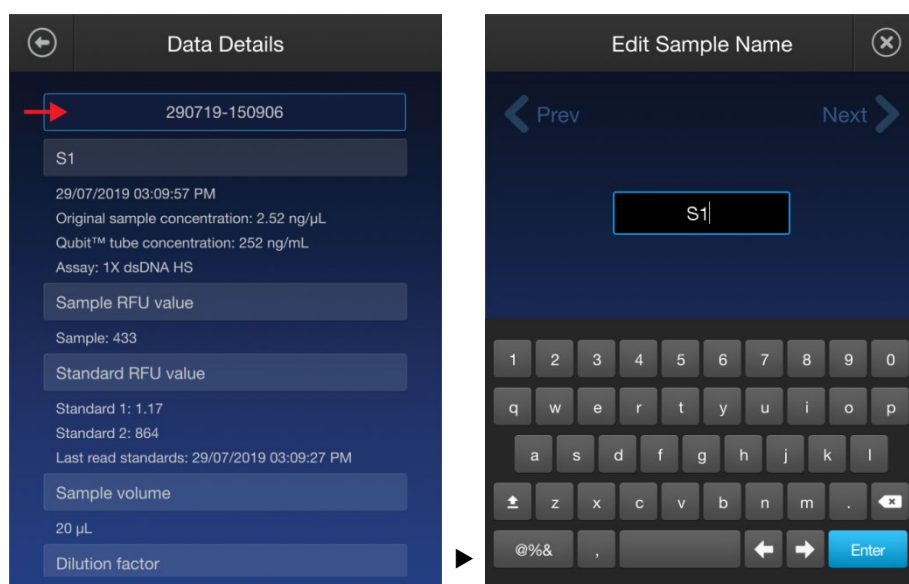
⑭ Molarity calculation information (units, nucleic acid length, MW, molarity)
- *RFU: Relative Fluorescence Units

Edit sample name

- Edit sample name**
1. On the Data screen, select the **data set of interest**, then select the **sample** you want to rename.



2. On the Data details screen, press the **Sample set #** field (indicated by red arrow). Edit Sample Name screen opens.



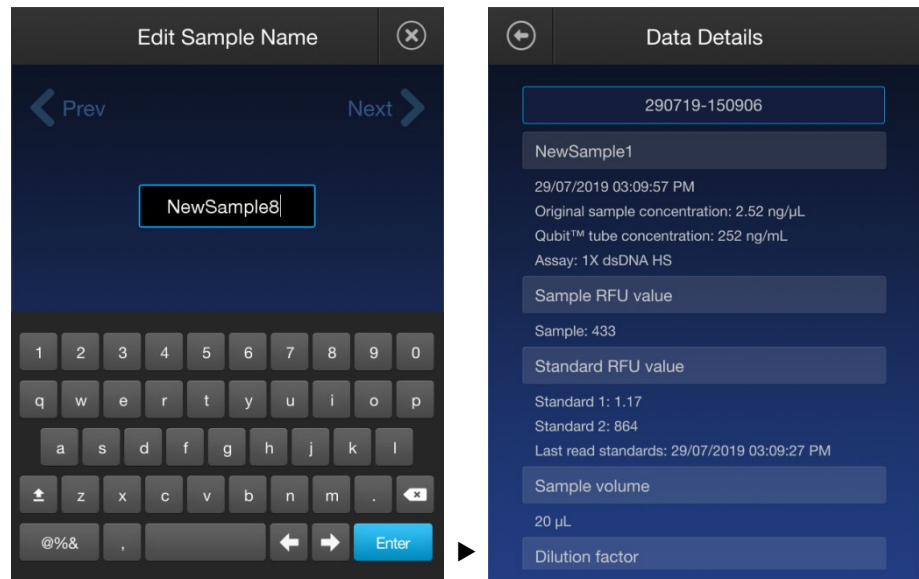
3. Enter the desired sample name, then press **Enter**. Data Details screen reappears and displays the new sample name.

The image shows two screenshots of a mobile application interface. The left screenshot is titled 'Edit Sample Name' and features a text input field containing 'NewSample1'. Below the input field is a QWERTY keyboard with an 'Enter' button highlighted in blue. Navigation buttons 'Prev' and 'Next' are visible at the top. The right screenshot is titled 'Data Details' and displays a list of assay parameters: '290719-150906', 'NewSample1', '29/07/2019 03:09:57 PM', 'Original sample concentration: 2.52 ng/μL', 'Qubit™ tube concentration: 252 ng/mL', 'Assay: 1X dsDNA HS', 'Sample RFU value', 'Sample: 433', 'Standard RFU value', 'Standard 1: 1.17', 'Standard 2: 864', 'Last read standards: 29/07/2019 03:09:27 PM', 'Sample volume', '20 μL', and 'Dilution factor'.

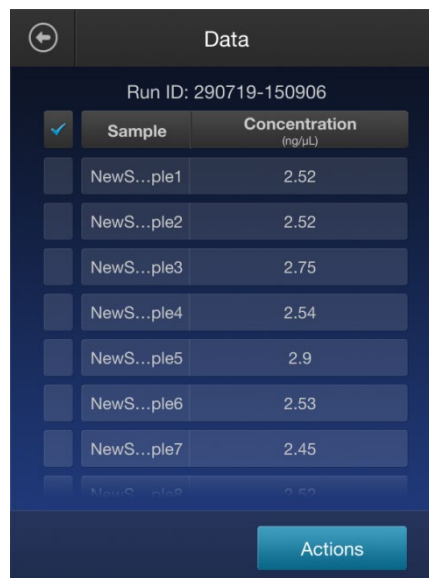
4. If you wish to rename all of the samples in the data set, press the **Next** button to go the next sample (instead of pressing **Enter** at step 3), then enter the new name for that sample.

The image shows two screenshots of the 'Edit Sample Name' screen. The left screenshot shows the input field with 'S2' and the keyboard. The right screenshot shows the input field with 'NewSample2' and the keyboard. Both screens have 'Prev' and 'Next' navigation buttons at the top.

- Repeat for all remaining samples. When finished renaming all the samples, press **Enter**. Data Details screen reappears and displays the new sample name.



- Press the **Back** button to return to the Data screen for the assay. All of the samples display the new sample names.

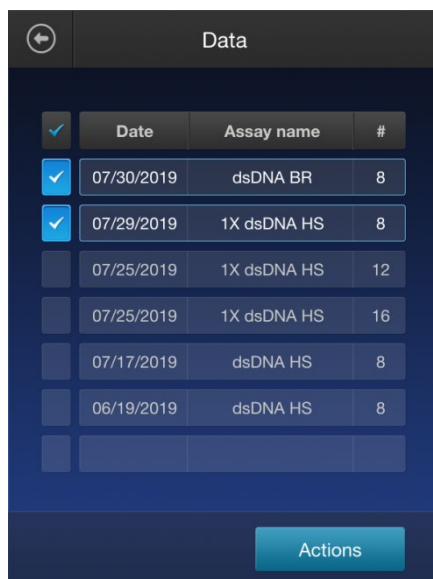


Export data

Introduction The Qubit™ Flex Fluorometer is designed for standalone use; it does not require an external computer. However, to archive data and generate reports, you can export the numeric data stored in the CSV file to a computer using a USB flash drive, or save to your Connect™ account or a network drive wirelessly or via the Ethernet cable. You can then view the file in any spreadsheet program.

- Export data**
1. On the **Home screen**, press **Data** to open the Data screen.
 2. To export entire data sets, press the **check box** to the left of each data set that you wish to export. You can select multiple data sets.

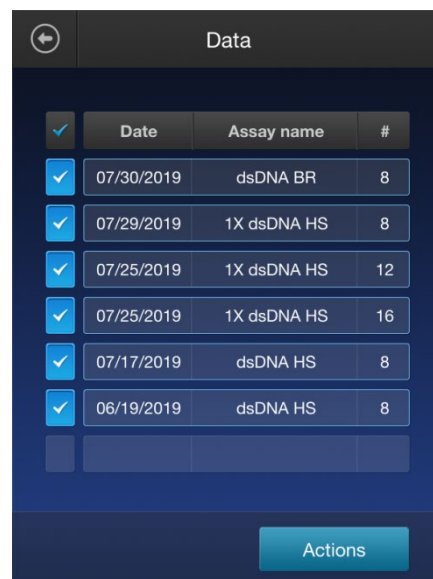
To select all data sets to export, press the blue **check** icon on the header row.



The screenshot shows the 'Data' screen with a table of data sets. The first two rows are selected, indicated by blue checkmarks in the selection column. The table has columns for Date, Assay name, and #. An 'Actions' button is at the bottom right.

	Date	Assay name	#
<input checked="" type="checkbox"/>	07/30/2019	dsDNA BR	8
<input checked="" type="checkbox"/>	07/29/2019	1X dsDNA HS	8
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	12
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	16
<input type="checkbox"/>	07/17/2019	dsDNA HS	8
<input type="checkbox"/>	06/19/2019	dsDNA HS	8
<input type="checkbox"/>			

Two data sets selected

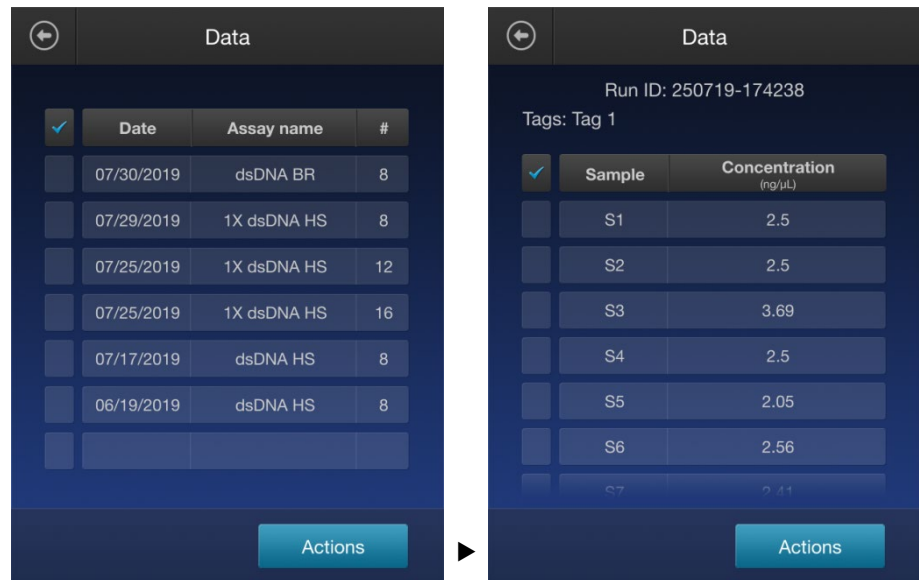


The screenshot shows the 'Data' screen with a table of data sets. All rows are selected, indicated by blue checkmarks in the selection column. The table has columns for Date, Assay name, and #. An 'Actions' button is at the bottom right.

	Date	Assay name	#
<input checked="" type="checkbox"/>	07/30/2019	dsDNA BR	8
<input checked="" type="checkbox"/>	07/29/2019	1X dsDNA HS	8
<input checked="" type="checkbox"/>	07/25/2019	1X dsDNA HS	12
<input checked="" type="checkbox"/>	07/25/2019	1X dsDNA HS	16
<input checked="" type="checkbox"/>	07/17/2019	dsDNA HS	8
<input checked="" type="checkbox"/>	06/19/2019	dsDNA HS	8
<input type="checkbox"/>			

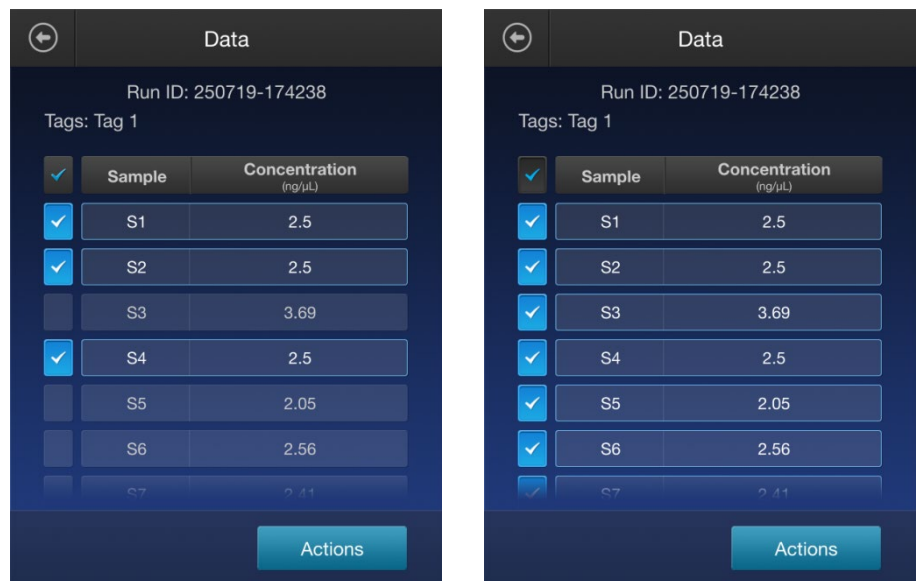
All data sets selected

- To export only individual data entries from a data set, press the **data set of interest** to view individual samples in the data set.



- Press the **check box** to the left of the samples that you wish to export. You can select multiple samples to export.

To select all samples in the data set to export, press the blue **check** icon on the header row.

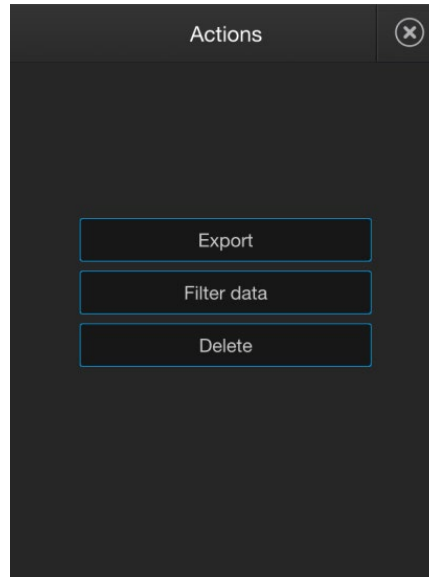


Three samples in the data sets selected

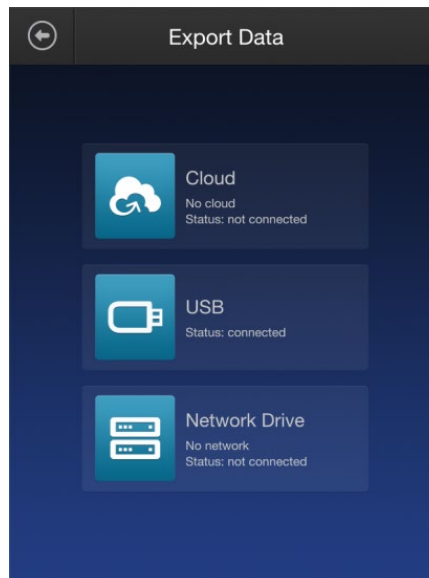
All samples in the data set selected

5. After you have selected the data sets or the samples, press **Actions**, then select **Export**.

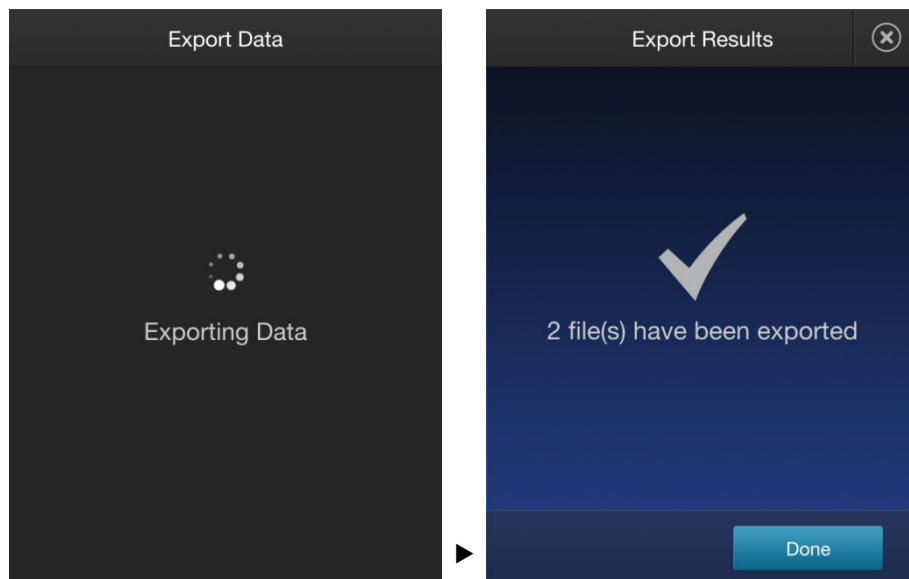
Actions



6. In the **Export data** screen, select the **Export method**. Available options are **Cloud** (i.e., your Connect™ account), **USB**, and **Network Drive**.
- To export data to a USB drive, insert the USB drive into the Qubit™ Flex Fluorometer.
 - To export data to your Connect™ account or a network drive, ensure that the instrument is connected to the network wirelessly or via an Ethernet cable.



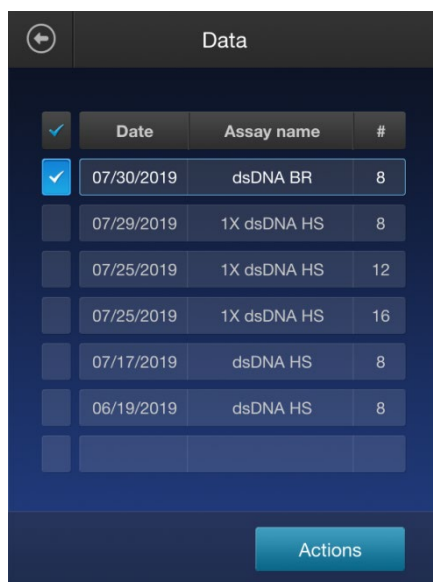
7. Press **Export** to export the data. The numeric data is automatically saved as a CSV file. You can open the CSV file using any spreadsheet program.



Delete data

- Delete data files**
1. On the **Home screen**, press **Data**.
 2. On the Data screen, press the **check box** to the left of each data set you wish to delete. To select all data sets, press the blue **check** icon on the header row.

To delete only individual sample files from a data set, press the **data set of interest** to view individual samples in the data set, then press the **check box** to the left of the samples you wish to delete.



<input checked="" type="checkbox"/>	Date	Assay name	#
<input checked="" type="checkbox"/>	07/30/2019	dsDNA BR	8
<input type="checkbox"/>	07/29/2019	1X dsDNA HS	8
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	12
<input type="checkbox"/>	07/25/2019	1X dsDNA HS	16
<input type="checkbox"/>	07/17/2019	dsDNA HS	8
<input type="checkbox"/>	06/19/2019	dsDNA HS	8
<input type="checkbox"/>			

Actions

Select data sets to delete



Run ID: 250719-174238
Tags: Tag 1

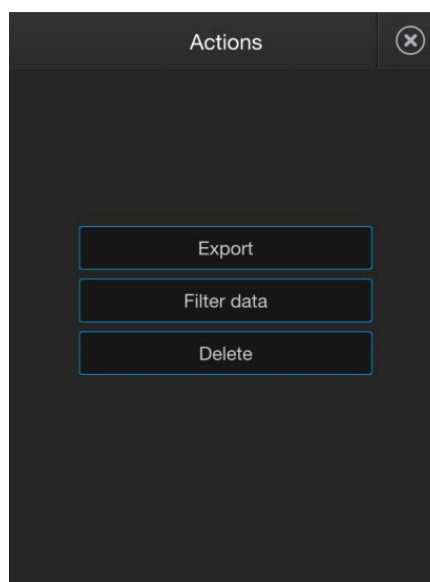
<input checked="" type="checkbox"/>	Sample	Concentration (ng/μL)
<input checked="" type="checkbox"/>	S1	2.5
<input checked="" type="checkbox"/>	S2	2.5
<input type="checkbox"/>	S3	3.69
<input checked="" type="checkbox"/>	S4	2.5
<input type="checkbox"/>	S5	2.05
<input type="checkbox"/>	S6	2.56
<input type="checkbox"/>	S7	2.11

Actions

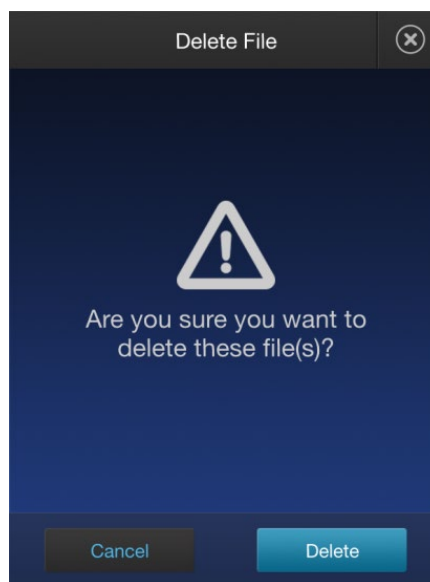
Select Sample files to delete

3. After you have selected the data sets or the samples, press **Actions**, then select **Delete**.

Actions



4. Press **Delete**. A warning screen appears.



5. Press **Delete** to permanently delete the sample data or data set.
6. Press **Cancel** to return to the screen previously viewed without deleting any data.

5. Configure instrument settings

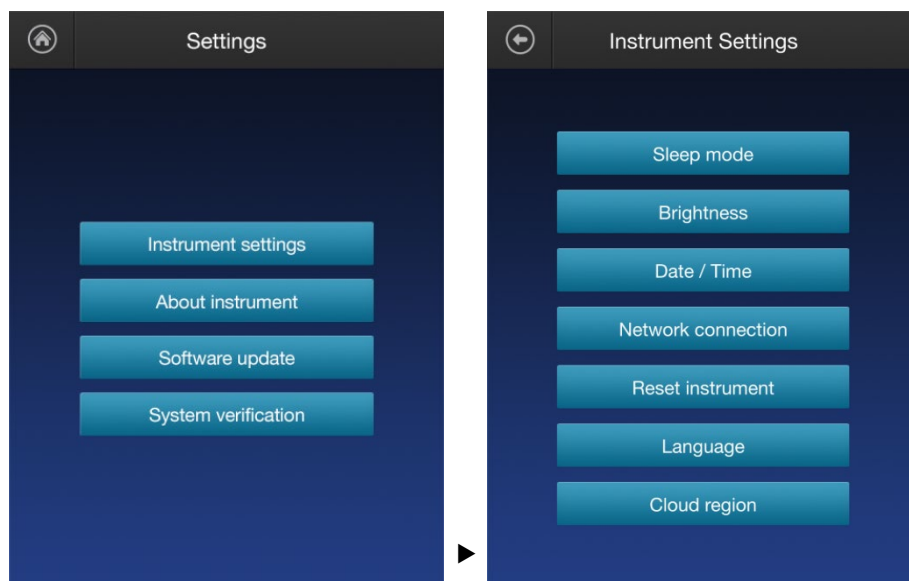
Instrument settings

You can configure the following instrument settings for the Qubit™ Flex Fluorometer from the **Settings** ► **Instrument Settings** screen:

- Sleep mode (page 73)
- Brightness (page 74)
- Date/Time (page 75)
- Network Connection (page 78)
- Reset instrument (page 84)
- Language (page 85)
- Cloud region (page 86)

Access the Instrument Settings screen

1. On the Home screen, press **Settings**.
2. On the **Settings** screen, press **Instrument settings** to display the Instrument settings screen.

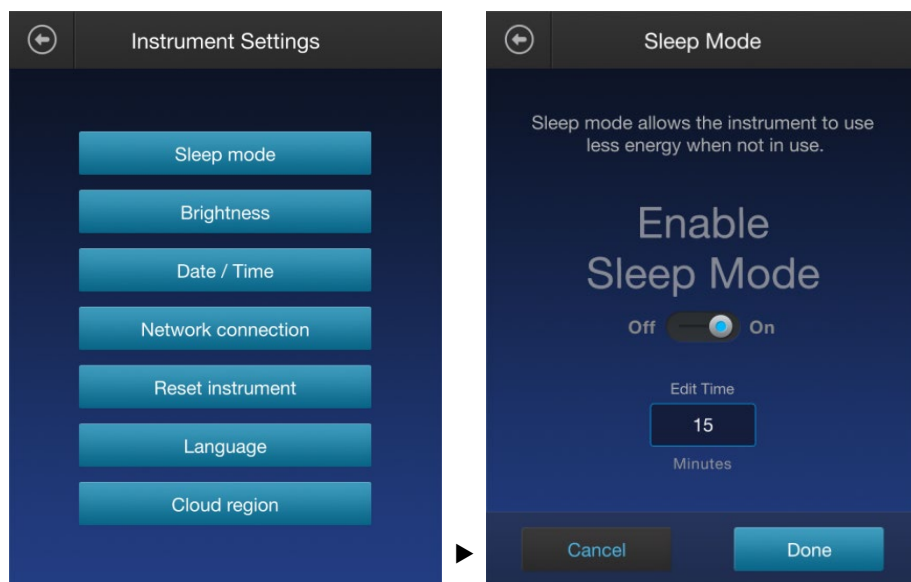


Sleep mode

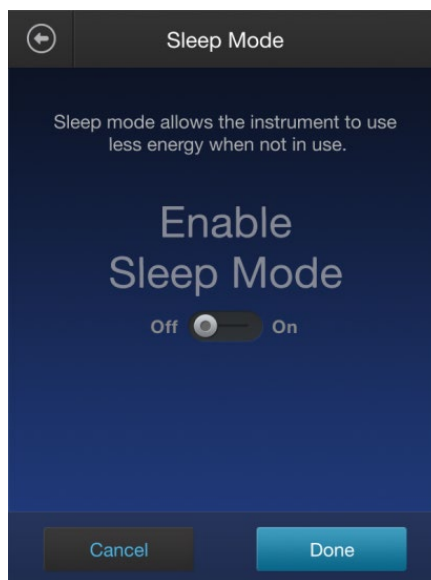
Adjust the sleep mode

The Qubit™ Flex Fluorometer has a sleep mode (i.e., automatic standby) that is triggered by inactivity. The system default is 10 minutes of inactivity before the instrument goes into sleep mode.

1. On the **Instrument Settings** screen (page 72), press **Sleep Mode**.



2. To change the time of inactivity before the instrument goes into sleep mode, press **Edit Time** field, then enter the time between 1 minute and 60 minutes.
3. To disable the sleep mode, toggle the **Enable Sleep Mode** switch to the **Off** position.

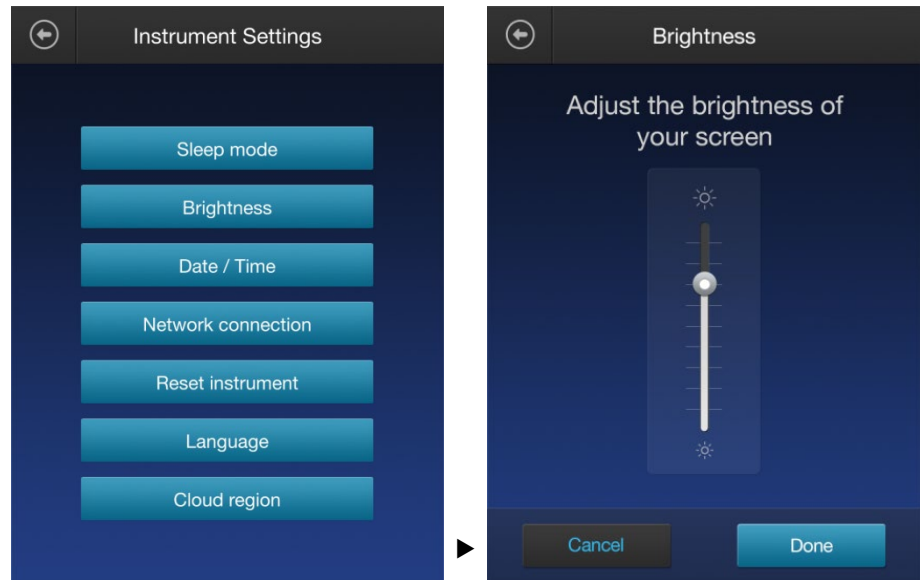


4. Press **Done** to save the changes and return to the Instrument Settings screen. Press **Cancel** or **Back** (⬅️) to return to the Instrument Settings screen without saving the changes.

Brightness

Adjust screen brightness

1. On the **Instrument Settings** screen (page 72), press **Brightness**.

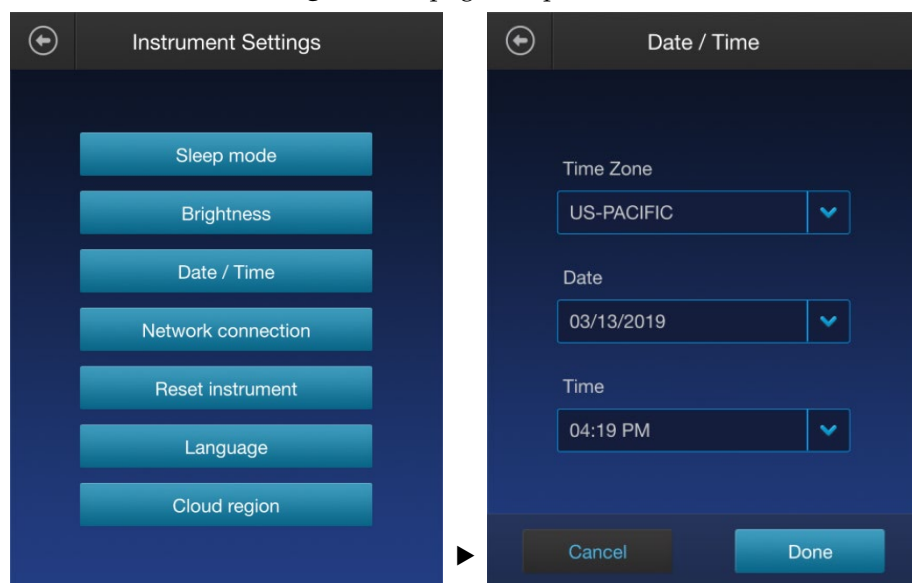


2. Move the **Brightness slider** up or down to adjust the brightness of the display.
3. Press **Done** to save the changes and return to the Instrument settings screen.
Press **Cancel** or **Back** (⬅️) to return to the Instrument settings screen without saving the changes.

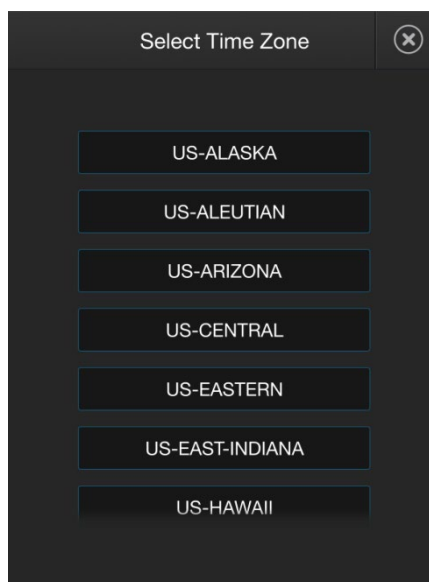
Date and Time

Set the date and time

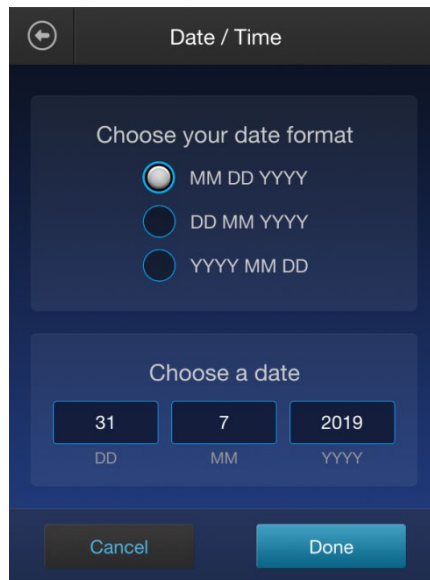
1. On the **Instrument Settings** screen (page 72), press **Date/Time**.



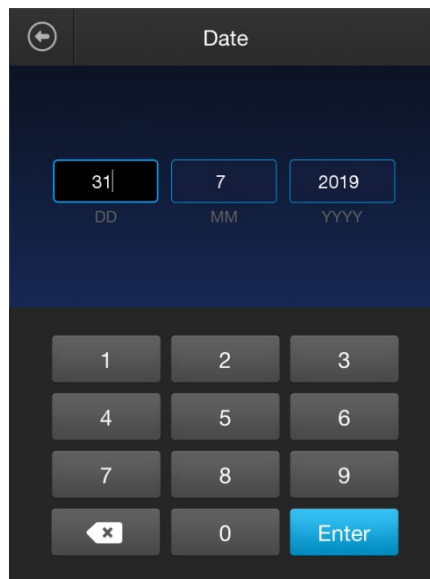
2. Press **Time Zone**, then select the time zone for your location from the list.



3. Press **Date**, then choose **MM DD YYYY**, **DD MM YYYY**, or **YYYY MM DD** for the date format.

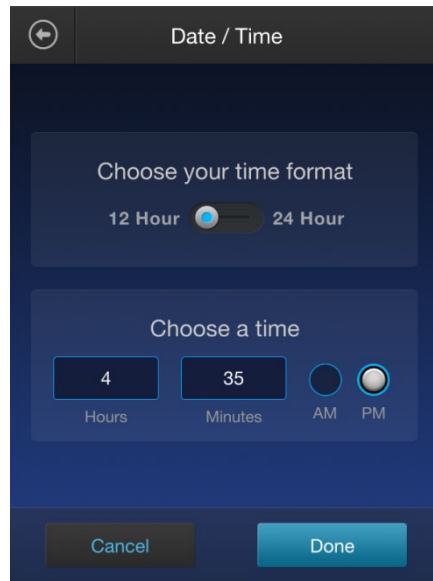


4. To set the date, press the **DD**, **MM**, and **YYYY** fields to enter the Day, Month, and Year.

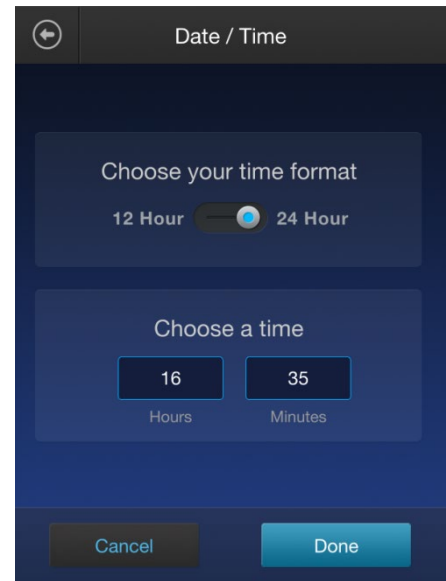


5. Press **Enter** when finished entering the date, then press **Done**.

6. Press **Time**, then choose **12 Hour** or **24 Hour** for the time format.

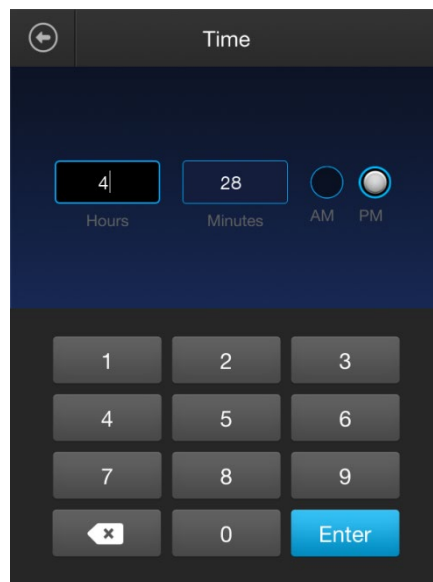


12-Hour format

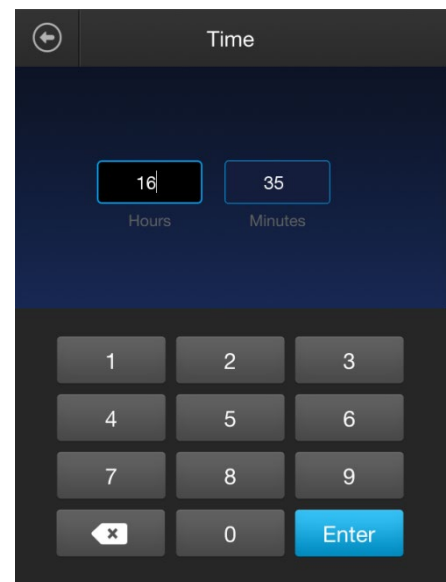


24-Hour format

7. To set the time, press the **Hours** and **Minutes** fields to enter the Hours and Minutes. If you have chosen the 12 Hour format, select **AM** or **PM**.



12-Hour format



24-Hour format

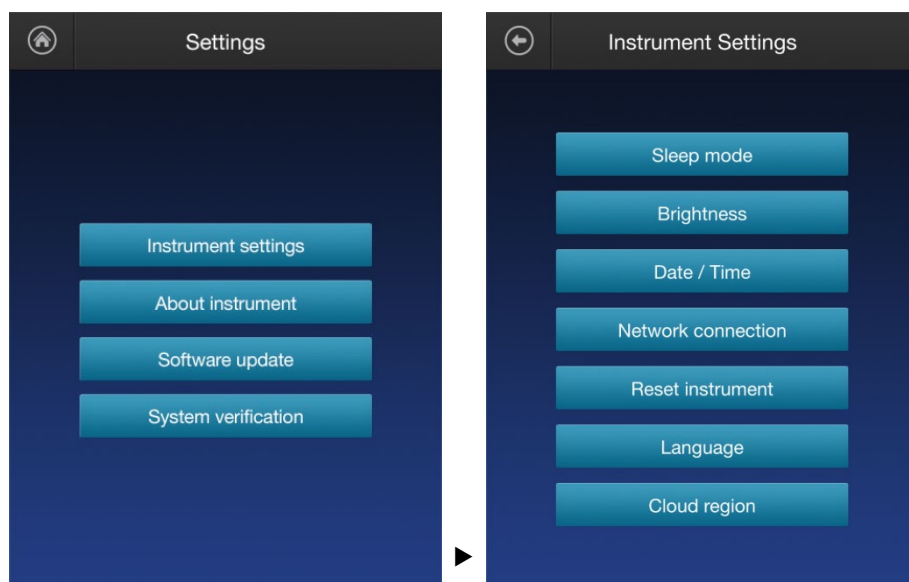
8. Press **Enter** when finished entering the time, then press **Done**.

Network connection

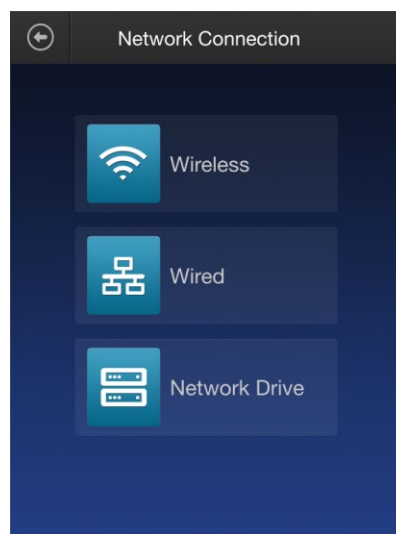
Access the Network Connection screen

Network Connection screen allows you to connect to an available wireless network using the supplied Wi-Fi adaptor, or to configure and join a local area network via the LAN (RJ-45) port using an Ethernet cable. After you have joined a network, you can also connect to Thermo Fisher's Connect™ cloud-based platform to store and access your data files.

1. To access the Network Connection screen, press **Settings** ► **Instrument Settings**, then select **Network connection**.



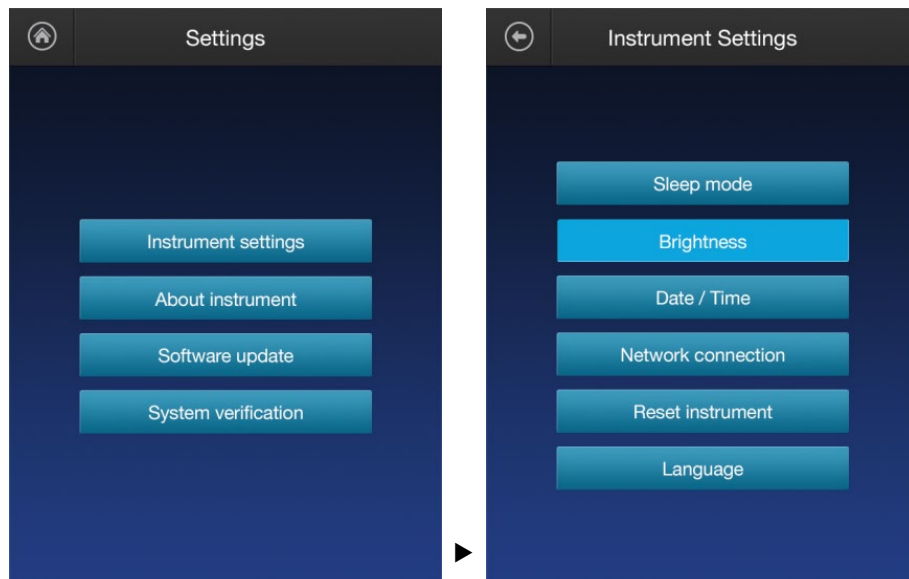
2. The Network Connection screen opens.



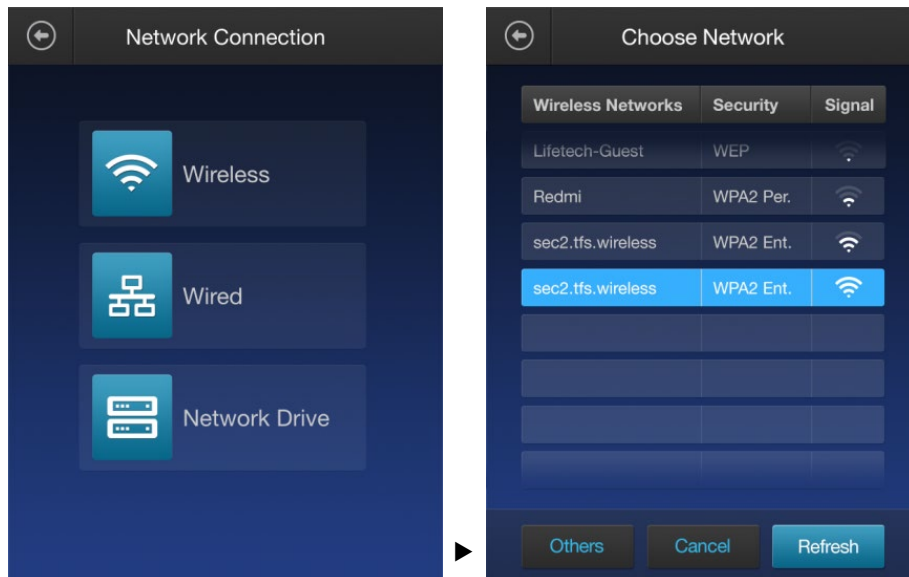
- To connect to a Wi-Fi network, go to page 79.
- To establish a wired connection to a local area network (LAN), go to page 80.
- To map a network drive to save your Qubit™ Flex files, go to page 81.

Connect to a Wi-Fi network

1. Ensure that your USB Wi-Fi dongle is inserted into one of the available USB ports on the instrument (see page 7).
If it is not, insert the Wi-Fi dongle, then restart the instrument by disconnecting and reconnecting the power supply.
2. Press **Settings** ► **Instrument Settings**, then select **Network connection**.



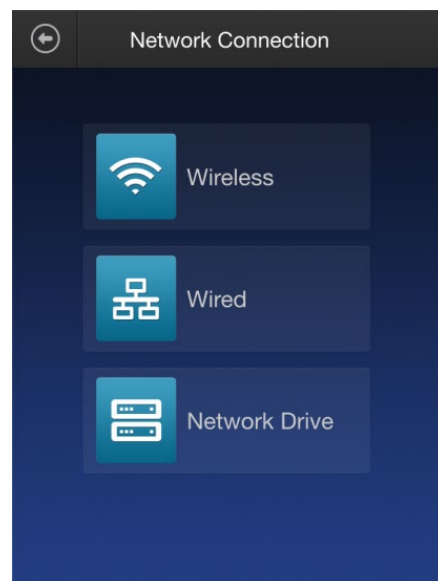
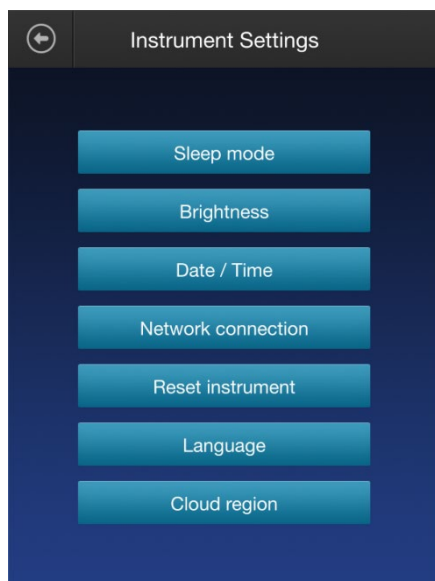
3. On the **Network Connection** screen, press **Wireless**. The instrument searches for available wireless networks within range.



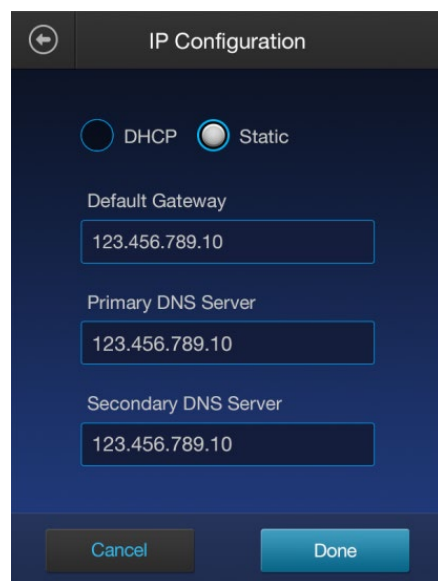
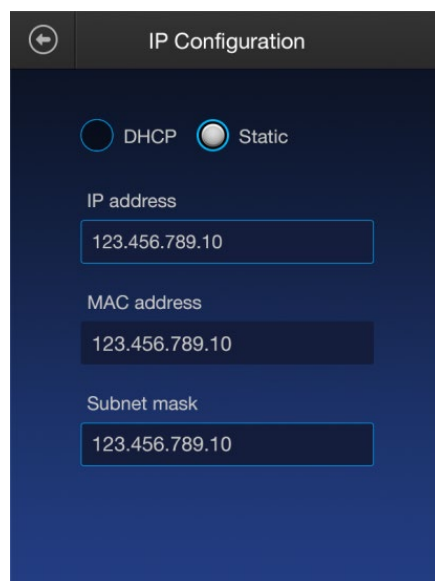
4. On the **Choose Network** screen, press the network you want to join.
5. If required, enter the appropriate security credentials, then press **Join**. After the connection is established, the network is highlighted in blue.

Connect to a local area network (LAN)

1. Ensure that the instrument is connected to an active network jack via the LAN (RJ45) port (page 7) using a standard Category 6 Ethernet cable.
2. On the **Instrument Settings** screen, press **Network connection**, then select **Wired**.



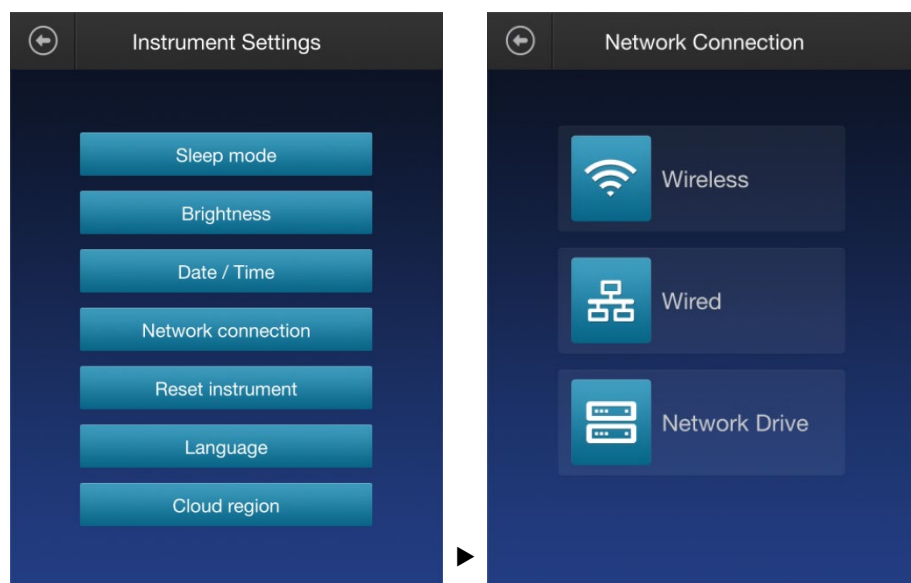
3. On the **IP Configuration** screen, select **DHCP** or **Static**.
4. If you have selected **Static**, enter the static **IP address**, **MAC address**, **Subnet mask**, **Default Gateway address**, and primary and secondary **DNS server addresses** for the LAN port.



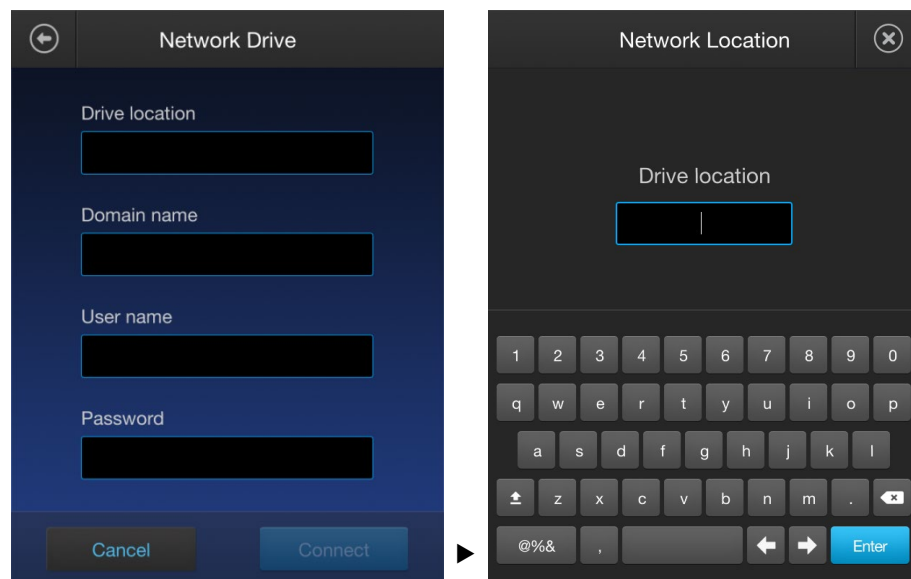
5. Press **Done** to join the local area network.

Map a Network Drive

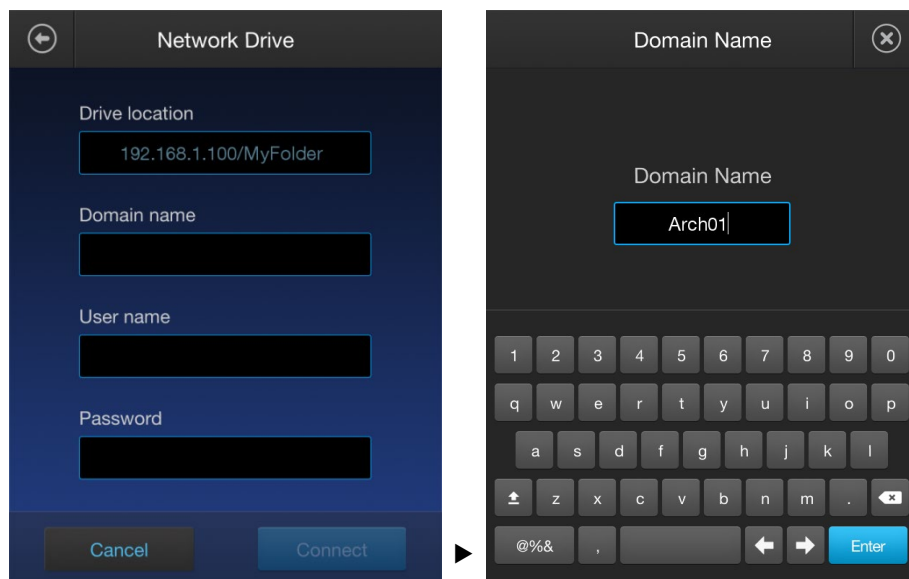
1. Ensure that the instrument is connected to an active network and that you have signed in to your profile (page 24).
2. On the **Instrument Settings** screen, press **Network connection**, then select **Network Drive**.



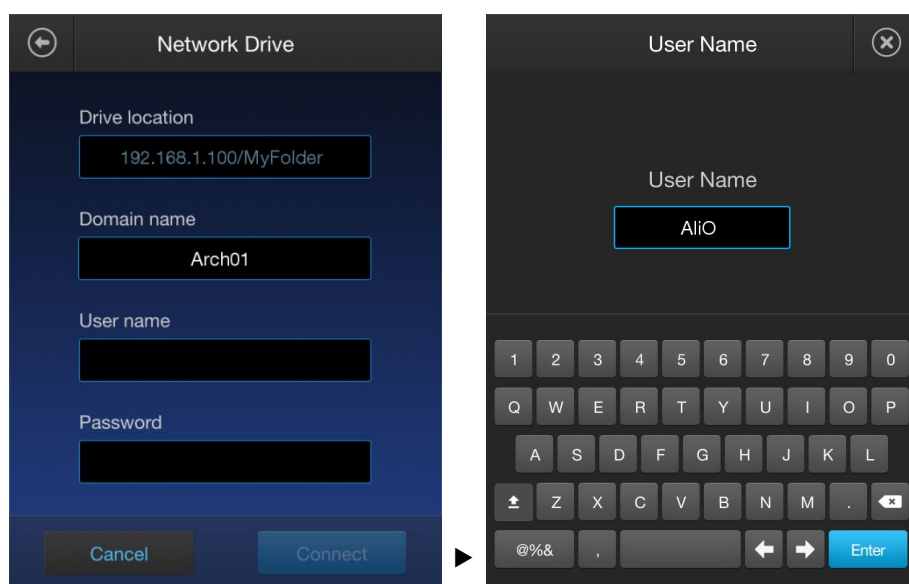
3. On the Network Drive screen, press **Drive location**, enter the location of the drive to save your Qubit™ Flex files, then press **Enter**.



4. Press **Domain name**, enter the domain name where the drive is located, then press **Enter**.



5. Press **User name**, enter your user name for the network drive, then press **Enter**.



6. Press **Password**, enter your password for the network drive, then press **Enter**.

The left screenshot shows the 'Network Drive' configuration screen. It has a title bar with a back arrow and the text 'Network Drive'. Below the title bar are four input fields: 'Drive location' (containing '192.168.1.100/MyFolder'), 'Domain name' (containing 'Arch01'), 'User name' (containing 'AliO'), and 'Password' (empty). At the bottom are two buttons: 'Cancel' and 'Connect'. A right-pointing arrow is positioned between the two screenshots.

The right screenshot shows the 'Password' input screen. It has a title bar with the text 'Password' and a close button. Below the title bar is a 'Password' label and a text input field containing six dots. Below the input field is a numeric keypad with buttons for digits 1-0, letters q-p, a-s, d-f, g-h, j-k, l, and a bottom row with '@%&', a comma, a left arrow, a right arrow, and an 'Enter' button.

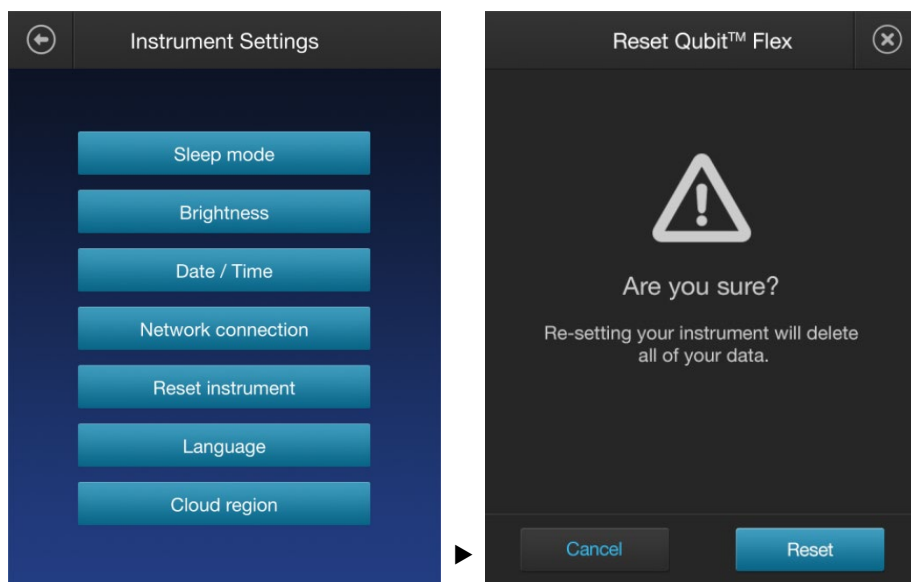
7. When finished entering all the required fields for the Network Drive, press **Connect**.

This screenshot shows the 'Network Drive' configuration screen with all fields filled: 'Drive location' is '192.168.1.100/MyFolder', 'Domain name' is 'Arch01', 'User name' is 'AliO', and 'Password' is masked with six dots. The 'Connect' button at the bottom right is highlighted in blue, indicating it is the next step.

Reset instrument

Reset instrument Reset instrument function returns the Qubit™ Flex Fluorometer to its default factory settings, and **erases all saved data** and **user-defined instrument settings**.

1. On the **Instrument Settings** screen (page 72), press **Reset instrument** to display the Reset Qubit™ Flex screen.



2. To return the instrument to its default factory settings, press **Reset**.
After the reset is complete, all data, user-defined instrument settings, and custom assays are removed, and the instrument displays the Home screen.
Press **Cancel** or **Exit** (ⓧ) to return to the Instrument settings screen without saving the changes.



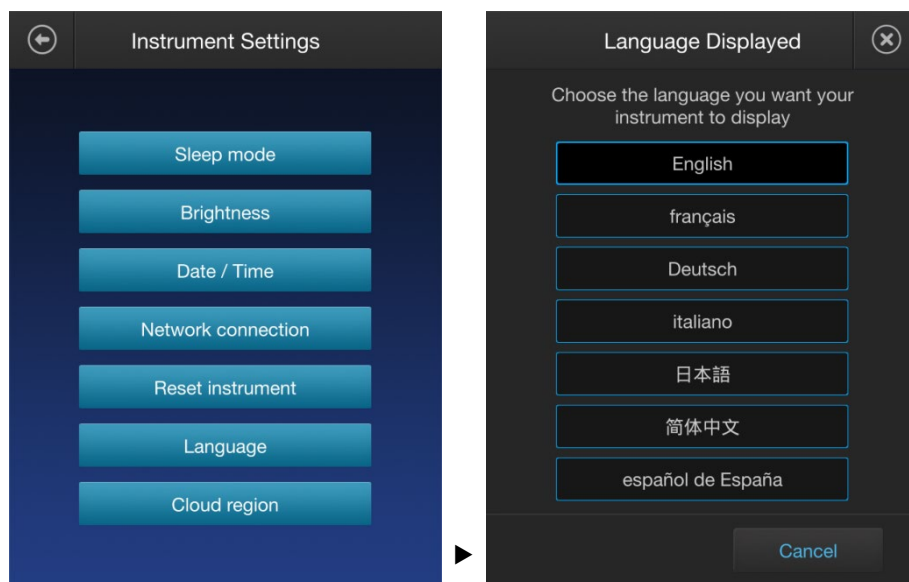
IMPORTANT! The reset function is **not** reversible.

Language

Change the displayed language

You can change the language that the Qubit™ Flex Fluorometer displays to English (default), French, German, Italian, Spanish, simplified Chinese, and Japanese.

1. On the **Instrument Settings** screen (page 72), press **Language** to display the Language screen.



2. Press to select the desired language. Available options are **English, French, German, Italian, Chinese, Japanese, and Spanish**.
3. When prompted, press **Yes** to confirm the change and return to the Instrument settings screen.

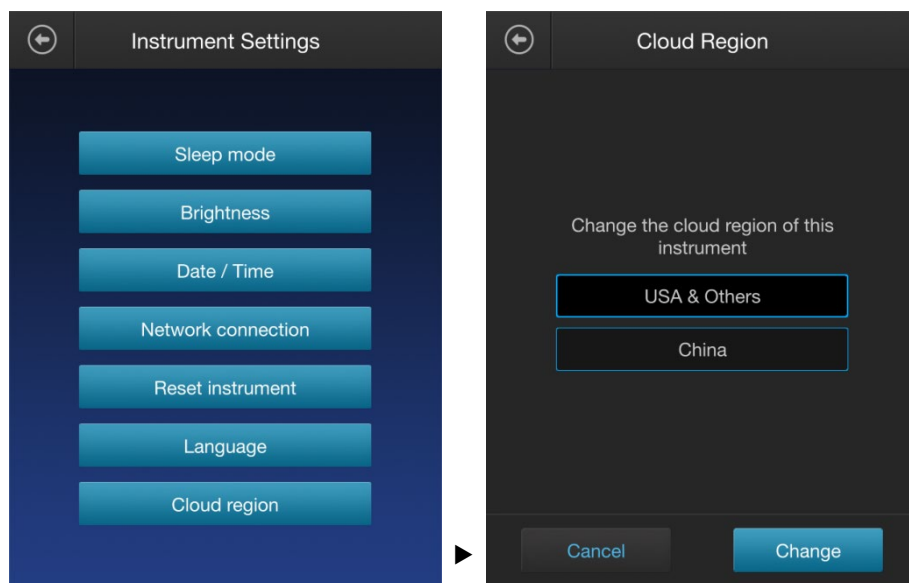
If you do not want to change the language settings, press **Cancel** or **Exit** (✕) to return to the Instrument settings screen without saving the changes.



Cloud region

Change the cloud region

1. On the **Instrument Settings** screen (page 72), press **Cloud region**.



2. Select the cloud region from the available choices, then press **Change**.
3. When prompted, press **Change** to close the warning screen, then press **Change** again to change the cloud region of the instrument. The instrument will restart after changing the cloud region.

If you do not want to change the cloud region, press **Cancel** to return to the previous screen.



Note: If you change the cloud region of the instrument, you must relink all instrument accounts.

6. Instrument maintenance

Maintenance and cleaning

- Maintenance** The Qubit™ Flex Fluorometer does not need regular maintenance. To troubleshoot problems with the instrument, contact Technical Support (page 111).
- **Do not** perform any repairs or service on the Qubit™ Flex Fluorometer to avoid damaging the instrument.
 - **Do not expose the Qubit™ Flex Fluorometer to direct sunlight.**



CAUTION! Never disassemble or service the instrument yourself. Do not remove any covers or parts that require the use of a tool. Unauthorized repairs may damage the instrument or alter its functionality, which may void your warranty. Contact your local distributor to arrange for service.

Clean the Qubit™ Flex Fluorometer

We recommend that you clean the Qubit™ Flex Fluorometer periodically to prevent the buildup of dust and dirt that might reduce its performance and cause contamination.



CAUTION! To avoid electrical shock, always disconnect the power cable before cleaning or decontaminating the instrument.



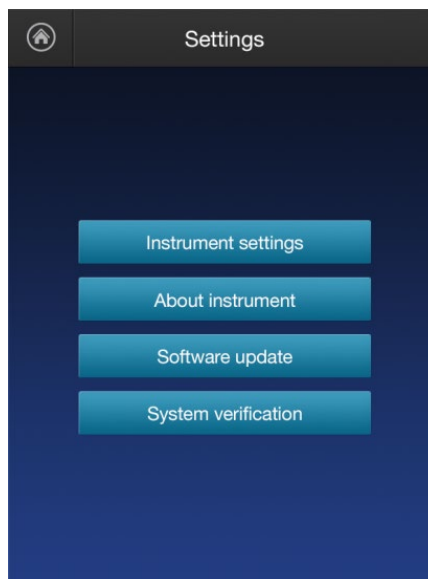
IMPORTANT! Using a cleaning or decontaminating method other than that specified by the manufacturer may result in damage to the instrument.

- Clean the surface of the Qubit™ Flex Fluorometer with a damp cloth.
- To clean the touchscreen, disconnect the power cable, and clean the touchscreen with a soft cloth lightly moistened with LCD (liquid crystal display) cleansing detergent.
- Cleaning the screen with excessive force can damage the touchscreen. Wipe the screen dry immediately.
- Do not use abrasive cleaning solutions or material to prevent the touchscreen from getting scratched.
- To disinfect the instrument, disconnect the power cable from the Qubit™ Flex Fluorometer and clean the instrument, including the touchscreen, with a soft cloth lightly moistened with 70% ethanol, 70% isopropanol, or 10% bleach (0.6% sodium hypochlorite).
- The cloth included with the instrument is not recommended for use with ethanol or isopropanol.
- Ensure that the cleaning solution does not enter the power button, the power inlet, the sample port, or the USB drive ports.
- Never pour or spray any liquids directly on the instrument to avoid electrical shock when the instrument is plugged in.

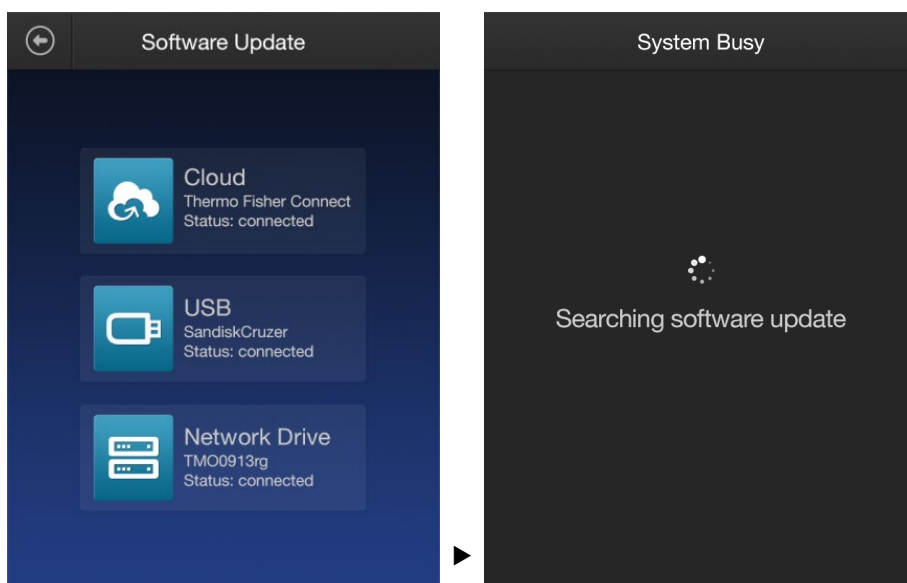
Software updates

- Before you begin**
1. Download the latest software to a USB drive or to your network from thermofisher.com/qubit.
 2. If using a USB drive, insert the USB drive into the instrument.
If using a network drive, ensure that the instrument is connected to the network wirelessly or via an Ethernet cable.

- Update the software**
1. On the Home screen, press **Settings**, then select **Software update**.



2. On the **Software Update** screen, select **Cloud**, **USB**, or **Network Drive**. If you have selected Cloud or Network drive, enter your credentials to sign in.
The instrument searches your Connect™ account, the USB drive, or the network drive for the update.

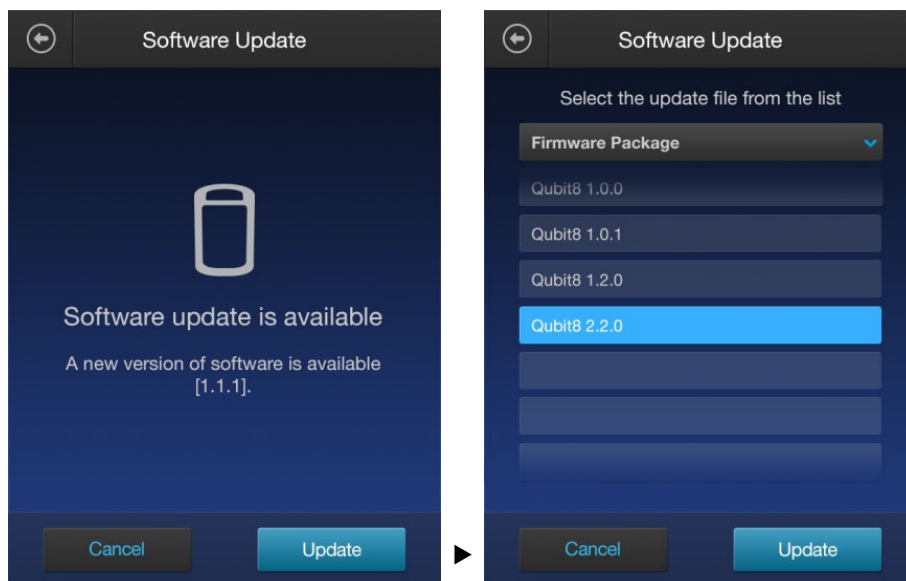




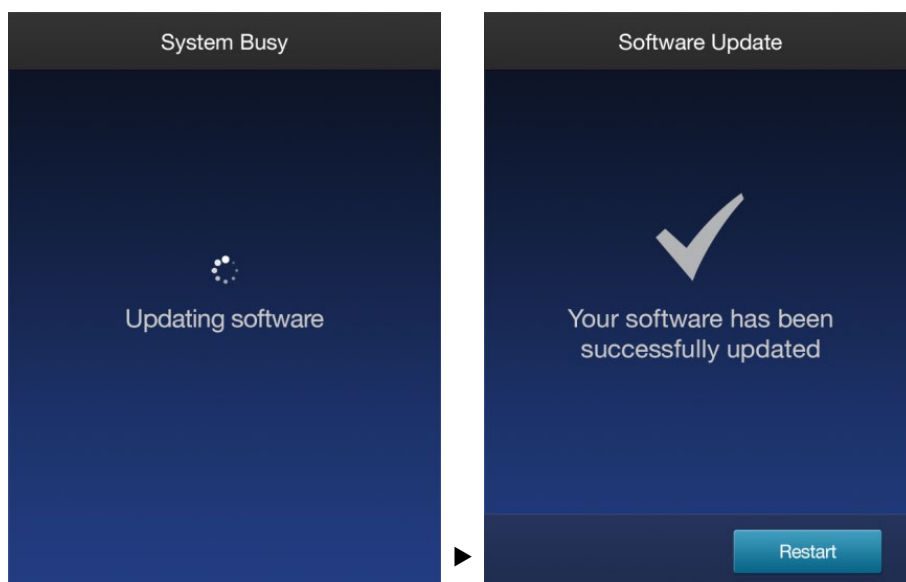
Note: If the USB drive is not inserted into the USB drive port or the instrument does not recognize the USB drive, a warning message is displayed.

To proceed with the software update, insert the correct USB drive into the instrument, then press **Retry**.

3. If a new update is available and the appropriate files are detected, the instrument displays “Software update is available”. Press **Update** to view the available versions of the software.



4. Select the software version you want install on the instrument for the update, then press **Update**.
5. When prompted, press **Restart** to complete the software update.

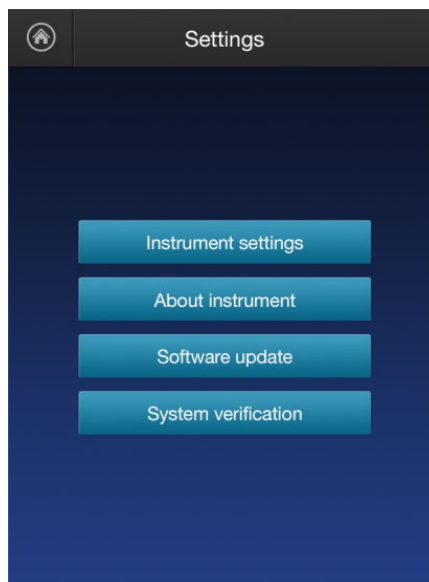


System verification

The system verification checks the internal components of the Qubit™ Flex Fluorometer and requires the use of the Qubit™ Flex System Verification Assay Kit (Cat. No. Q33254). Perform the system verification when a problem with the instrument is suspected. It is not necessary to perform the verification regularly.

Perform System verification test

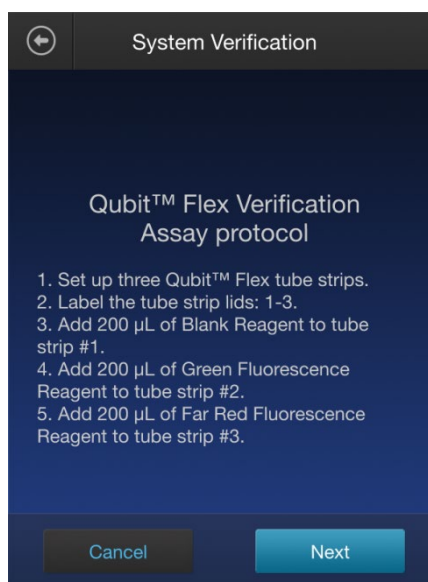
1. On the **Home** screen, press **Settings**, then select **System Verification**.



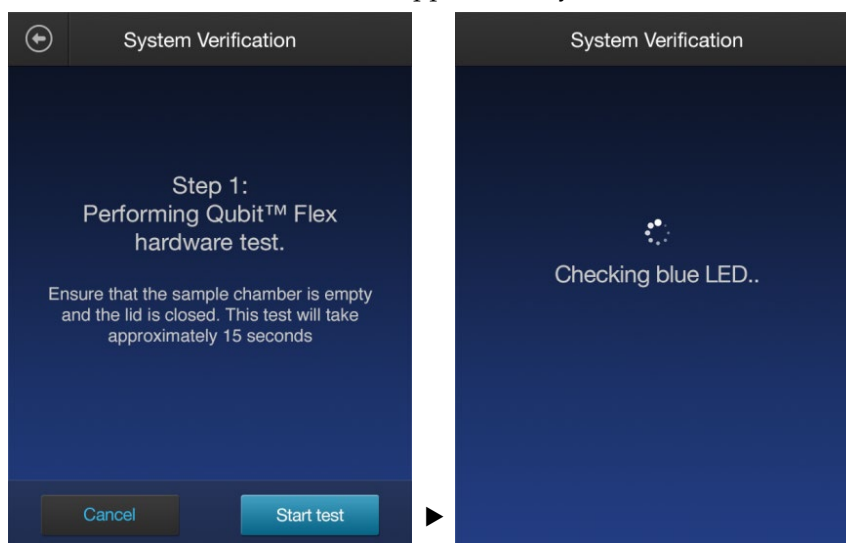
2. On the **System Verification** screen, press **Next**.



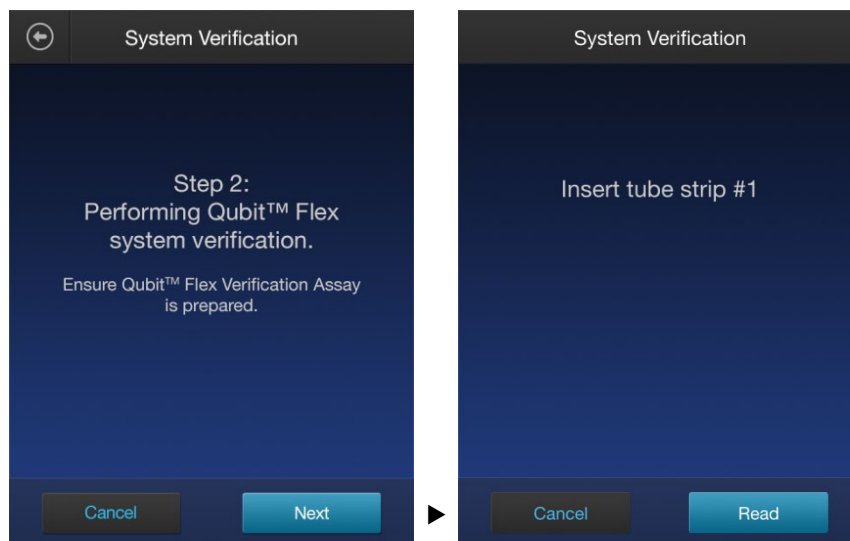
3. When prompted, set up three Qubit™ Flex Tube Strips and label the tube strip lids 1–3.



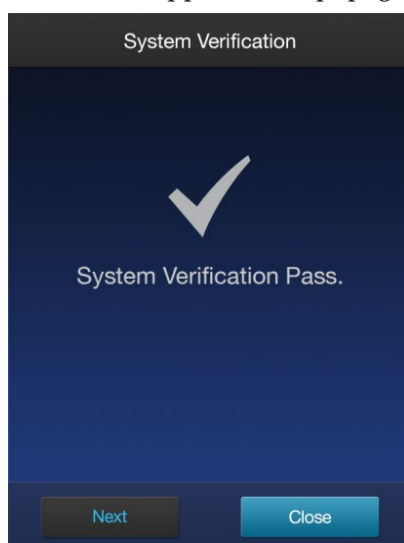
4. Add 200 µL of Blank Reagent to each tube of tube strip #1, 200 µL of Green Fluorescence Reagent to each tube of tube strip #2, and 200 µL of Far Red Fluorescence Reagent to each tube of tube strip #3, then press **Next**.
5. When prompted, ensure that the sample chamber is empty and the lid is closed, then press **Start test** to run the Qubit™ Flex hardware test (Step 1 of System Verification). This test takes approximately 15 seconds.



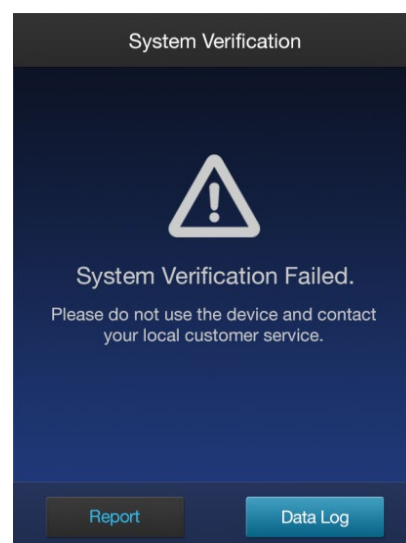
6. When prompted, ensure that the Qubit™ Flex Verification Assay is prepared, then press **Next**.



7. Insert tube strip #1 into the sample chamber, close the lid, then press **Read**.
8. When prompted, read tube strip #2 and tube strip #3 as described for tube strip #1.
9. When the test is complete, the software displays the error status.
- If no errors are found, **System Verification Pass** message appears. Press **Close** to return to the Settings screen or press **Next** to view the System Verification Report (page 93).
 - If errors are found, **Error Reading Reagents** message appears. Verify that the test was run with the lid closed, then press **OK** to re-run the test with the tube strips in the correct order.
 - If the **System Verification Failed** message persists after re-running the tube strips with the lid closed, do **not** use the instrument and contact Technical Support for help (page 111).



System Verification Pass



System Verification Failed

10. Press **Report** to view the *System Verification Report* or press **Data Log** to view and export the available data logs (page 93).

The *System Verification Report* shows the pass/fail status of the instrument components.

The screenshot shows the 'System Verification' screen with a timestamp of 20/09/2019 02:22:00 PM and a 'Saved' status. It contains a table with the following data:

System Verification	Result
Hardware Test	
Blue LED check	PASS
Red LED check	PASS
Photodiode check	PASS
Green Fluorescence Channel	PASS
Far Red Fluorescence Channel	FAIL

At the bottom, there are two buttons: 'Data Log' and 'Close'.

Far Red Fluorescence Channel Fail

The screenshot shows the 'System Verification' screen with a timestamp of 20/09/2019 02:28:33 PM and a 'Saved' status. It contains a table with the following data:

System Verification	Result
Hardware Test	
Blue LED check	PASS
Red LED check	PASS
Photodiode check	PASS
Green Fluorescence Channel	PASS
Far Red Fluorescence Channel	PASS

At the bottom, there are two buttons: 'Data Log' and 'Close'.

All Pass

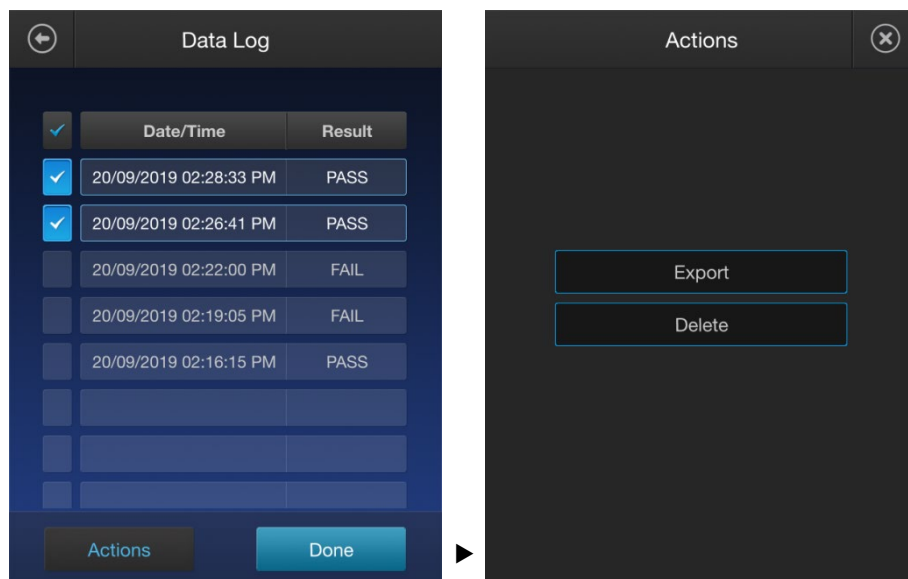
11. Press **Close** to return to the Settings screen or press **Data Log** to view the available data logs.

The screenshot shows the 'Data Log' screen with a back arrow icon at the top left. It contains a table with the following data:

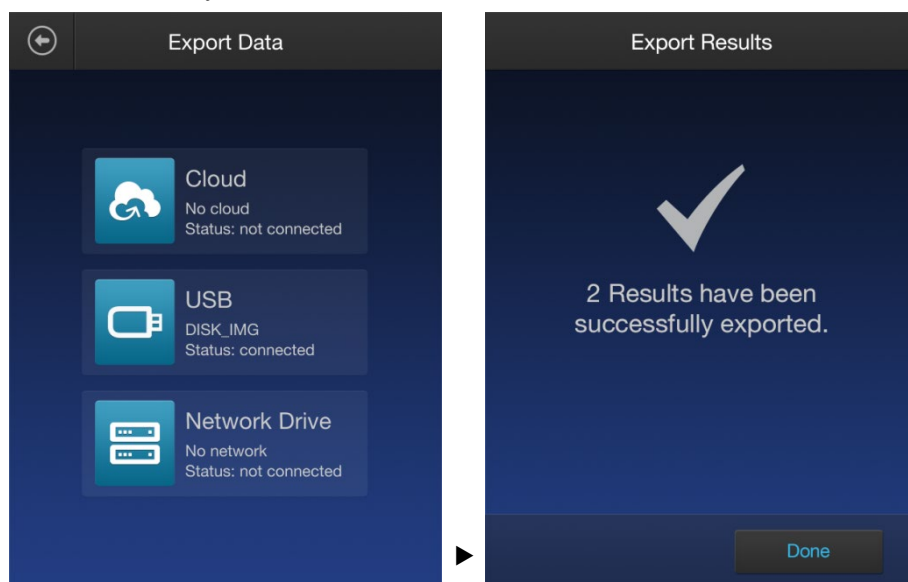
	Date/Time	Result
<input checked="" type="checkbox"/>	20/09/2019 02:28:33 PM	PASS
<input type="checkbox"/>	20/09/2019 02:26:41 PM	PASS
<input type="checkbox"/>	20/09/2019 02:22:00 PM	FAIL
<input type="checkbox"/>	20/09/2019 02:19:05 PM	FAIL
<input type="checkbox"/>	20/09/2019 02:16:15 PM	PASS
<input type="checkbox"/>		
<input type="checkbox"/>		
<input type="checkbox"/>		

At the bottom, there are two buttons: 'Actions' and 'Done'.

12. To export a Data Log as a PDF report, select the desired **Data Log**, press **Actions**, then press **Export**. You can select multiple Data Logs for export.



13. Select **Cloud** (Thermo Fisher Connect™ cloud-based platform), **USB**, or **Network Drive** for the location where you want to save the PDF report of the Verification Assay Test Results.



14. To delete a Data Log, select the desired **Data Log**, press **Actions**, then press **Delete**. You can select multiple Data Logs for deletion.

Replace battery

The Qubit™ Flex Fluorometer contains a 3 V CR2450 battery, which is required to record the export CSV file date and time. When the battery runs out, the system cannot keep the time setting, which indicates the need to replace the battery.

- Replace battery**
1. Disconnect the Qubit™ Flex Fluorometer from the power source.
 2. Remove the four screws (as indicated by the red arrows) on the bottom chassis of the Qubit™ Flex instrument using a Phillips-head screwdriver.



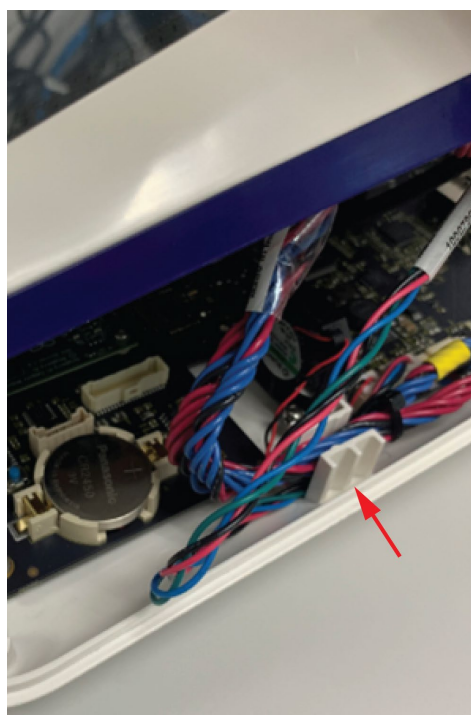
3. Flip the instrument so that the top chassis is facing up.
4. Open the instrument slightly (~ 3 cm) from the bottom right side.



5. Pry the old battery from its housing using a flat-head screwdriver and remove it.



6. Insert the new 3 V CR2450 battery to the battery housing.
7. Arrange two cable assemblies into the groove on the bottom chassis, place the top chassis on the bottom chassis so that the slots for the screws align properly, then tighten the four screws on the bottom chassis using a Phillips-head screwdriver.



Appendix A: Troubleshooting

Troubleshooting

- Handling samples**
- The calibration standards included in the Qubit™ microRNA, Qubit™ RNA HS, and Qubit™ RNA BR Assay Kits are high-quality RNA standards. The integrity and concentration of these standards is critical to the optimal performance of the Qubit™ RNA assays. We highly recommend treating the rRNA standards as you would any other precious RNA. Use appropriate RNase-free handling techniques, including RNase-free gloves, pipette tips, and tubes. Keep the tube lids closed whenever possible; do not press the pipet to the inside wall of the tube when withdrawing a sample. Return the RNA standards to –80°C as soon as possible after use.
 - Ensure that the assay tubes are at room temperature at the time the reading is taken. Do not hold assay tubes in your hand and do not leave assay tubes in the Qubit™ Flex Fluorometer for longer than it takes to read the fluorescence. See “Assay temperature”, page 100.
 - Be careful not to spill sample into the sample chamber. Promptly wipe any spills.
 - The Qubit™ assays are very sensitive and even small amounts of material from a previous sample may result in errors. Use a clean Qubit™ Flex Tube Strip for each reading.
 - The tube **must be clean and dry** on the outside when taking readings. Moisture and condensation on the tube surface can lead to reading errors.
 - Minute bubbles in samples will cause errors in readings. Be sure not to introduce bubbles into samples. Slight tapping on the tube wall or brief centrifugation will often help dissipate bubbles.

- High reading**
- The sample is out of range. Use a sample that is less concentrated or add a smaller volume of sample into the assay to further dilute the sample.
 - For Qubit™ quantification assays, view the Fluorescence vs. Concentration graph in the Results screen to confirm that the values for the samples fall between the values of the standards (page 43).
 - Ensure that the lid is closed while reading standards and samples.
 - Prepare samples and standards according to the instructions in the Qubit™ assay kit you are using.
 - Ensure that the assay is performed entirely at room temperature.



Low reading

- The sample is out of range. Use a sample that is more concentrated or use a lower dilution (for example, 20 μL in 180 μL instead of 10 μL in 190 μL).
- For Qubit™ quantification assays, view the Fluorescence vs. Concentration graph in the Results screen to confirm that the values for the samples fall between the values of the standards (page 43).
- Ensure that you have prepared the Qubit™ working solution correctly (1:200 dilution using the buffer provided in the kit).
- Ensure that you have prepared the standard tubes correctly (10 μL of each standard in 190 μL of Qubit™ working solution).
- Ensure that the standard and sample tubes are filled to 200 μL .
- Protect the Qubit™ reagent and working solutions from light.
- Select the correct Qubit™ Flex Fluorometer assay for the Qubit™ assay you are performing and calibrate the fluorometer correctly. Standards must be used in the correct order.
- Ensure that the assay is performed entirely at room temperature.



Critical Qubit™ Assay considerations

How the Qubit™ Flex Fluorometer calculates concentration

The Qubit™ Flex Fluorometer generates concentration data based on the relationship between the two standards used in calibration (three for the Qubit™ protein assay). The plot below shows the line corresponding to the curve-fitting algorithm (a modified Hill plot) used in the calculation of concentration data for the Qubit™ RNA HS assay. For reference, the positions of the standards and a set of data points from an actual experiment are shown superimposed onto the line. This plot demonstrates that the curve-fitting algorithm gives accurate values for quantification.

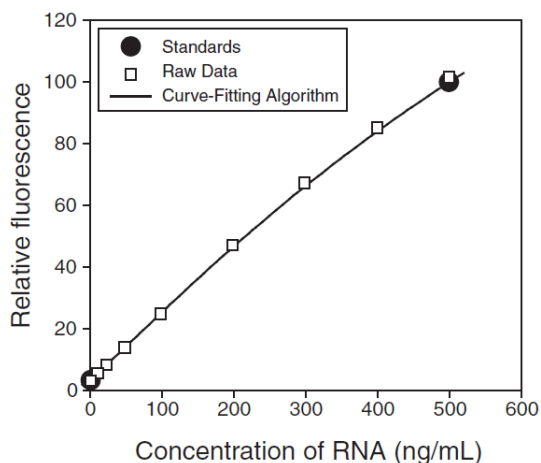


Figure 1. The curve-fitting algorithm used to determine concentration in the Qubit™ RNA HS assay. Data for other Qubit™ quantification assays are generated by similar algorithms.

Incubation time

To allow the Qubit™ assay to reach maximum fluorescence, incubate the tubes for the DNA and RNA assays for 2 minutes after mixing the sample or standard with the working solution. After this incubation period, the fluorescence signal is stable for 3 hours at room temperature for all nucleic acid assays except the Qubit™ ssDNA assay, which is stable for up to 30 minutes.

The Qubit™ protein assay requires 15 minutes of incubation for a stable signal. For greatest accuracy in the protein assay, the incubation time of the samples should be within 10 minutes of the incubation time of the standards.

Photobleaching of Qubit™ reagents

The Qubit™ DNA and protein exhibit high photostability in the Qubit™ Flex Fluorometer, showing <0.3% drop in fluorescence after 9 readings and <2.5% drop in fluorescence after 40 readings. It is important to remember, however, that if the assay tube remains in the Qubit™ Flex Fluorometer for multiple readings, a temporary reduction in fluorescence will be observed as the solution increases in temperature (see Figure 2 in “Assay temperature”, page 100). The RNA assays should only be read once.

Note that the temperature inside the Qubit™ Flex Fluorometer may be as much as 3°C above room temperature after 1 hour. For this reason, if you want to perform multiple readings of a single tube, remove the tube from the instrument and let it equilibrate to room temperature for 30 seconds before taking another reading.

Assay temperature The Qubit™ assays were designed to be performed at room temperature (22–28°C), and temperature fluctuations can influence the accuracy of the assay.

To minimize temperature fluctuations, store all kit reagents at room temperature and insert all assay tubes into the Qubit™ Flex Fluorometer only for as much time as it takes for the instrument to measure the fluorescence, because the Qubit™ Flex Fluorometer can raise the temperature of the assay solution significantly, even over a period of a few minutes. Do not hold the assay tubes in your hand before a measurement, because holding the tubes warms the solution and results in a low reading.

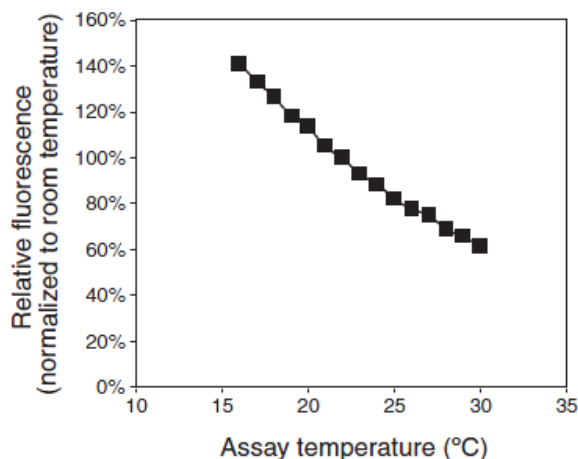


Figure 2. Effect of temperature on the Qubit™ dsDNA BR assay. Qubit™ dsDNA HS, Qubit™ ssDNA, Qubit™ RNA HS, and Qubit™ protein assays show similar sensitivities over the same range.

Qubit™ Flex Fluorometer calibration

For each assay, you have the choice to run standards for a new calibration or to use the values from the previous calibration.

As you first use the instrument, perform a new calibration each time. As you become familiar with the assays, the instrument, your pipetting accuracy, and significant temperature fluctuations within your laboratory, you can determine the level of comfort you have using the calibration data stored from the last time the instrument was calibrated.

Remember also that the fluorescence signal in the tubes containing the standards and the samples is stable for not longer than 3 hours. See Figure 1 in “How the Qubit™ Flex Fluorometer calculates concentration” (page 99) for an example of the calibration curve used to generate the quantification results.

Appendix B: Ordering information

Qubit™ Flex Fluorometer and accessories


The following products can be used with the Qubit™ Flex Fluorometer and are available separately from Thermo Fisher Scientific. For more information, visit [thermofisher.com](https://www.thermofisher.com) or contact Technical Support (page 111).

Product	Quantity	Cat. No.
Qubit™ Flex Fluorometer	1 each	Q33327
Qubit™ Flex Quantitation Starter Kit	1 kit	Q45894
Qubit™ Flex NGS Starter Kit	1 kit	Q45893
Qubit™ Flex USB Flash Drive	1 each	Q46009
Qubit™ Flex Wi-Fi dongle	1 each	A26774
Qubit™ Flex Fluorometer International Power Supply (replacement)	1 each	A36204
Qubit™ Flex Tube Strips	125 strips	Q33252
Qubit™ Flex Reservoir (10 mL)	100 each	Q33253
Qubit™ Flex System Verification Assay Kit	1 kit	Q33254
Qubit™ RNA BR Assay Kit *20–1,000 ng*	100 assays 500 assays	Q10210 Q10211
Qubit™ RNA HS Assay Kit *5–100 ng*	100 assays 500 assays	Q32852 Q32855
Qubit™ ssDNA Assay Kit *1–200 ng*	100 assays	Q10212
Qubit™ dsDNA BR Assay Kit *2–1,000 ng*	100 assays 500 assays	Q32850 Q32853
Qubit™ dsDNA HS Assay Kit *0.2–100 ng*	100 assays 500 assays	Q32851 Q32854
Qubit™ 1X dsDNA HS Assay Kit	100 assays 500 assays	Q33230 Q33231
Qubit™ Protein Assay Kit *0.25–5 µg*	100 assays 500 assays	Q33211 Q33212
Qubit™ microRNA Assay Kit *0.5–100 ng*	100 assays 500 assays	Q32880 Q32881
Qubit™ dsDNA HS Assay - Lambda DNA Standard	5 mL	Q33233



Appendix C: Safety

Symbols on instruments



Electrical symbols The following table describes the electrical symbols that may be displayed.

Symbol	Description
	Indicates a terminal that can receive or supply alternating current or voltage.

Safety symbols The following table describes the safety symbols that may be displayed. Each symbol may appear by itself or in combination with text that explains the relevant hazard (see “Safety labels on instruments”). These safety symbols may also appear next to DANGERS, WARNINGS, and CAUTIONS that occur in the text of this and other product-support documents.



Symbol	Description
	Indicates that you should consult the manual for further information and to proceed with appropriate caution.
	Indicates the presence of an electrical shock hazard and to proceed with appropriate caution.

Environmental symbols The following symbol applies to all Thermo Fisher Scientific electrical and electronic products placed on the European market after August 13, 2005.

Symbol	Description
 	<p>Do not dispose of this product as unsorted municipal waste. Follow local municipal waste ordinances for proper disposal provisions to reduce the environmental impact of waste electrical and electronic equipment (WEEE).</p> <p>European Union customers: Call your Customer Service representative for equipment pick-up and recycling. See thermofisher.com for a list of customer service offices in the European Union.</p>

Safety labels on instruments

The following CAUTION, WARNING, and DANGER statements may be displayed on Thermo Fisher Scientific instruments in combination with the safety symbols described in the preceding section.

Hazard symbol	English	Français
	CAUTION! Hazardous chemicals. Read the Safety Data Sheets (SDSs) before handling.	ATTENTION! Produits chimiques dangereux. Lire les fiches techniques de sûreté de matériels avant toute manipulation de produits.
	CAUTION! Hazardous waste. Refer to SDS(s) and local regulations for handling and disposal.	ATTENTION! Déchets dangereux. Lire les fiches techniques de sûreté de matériels et la réglementation locale associées à la manipulation et l'élimination des déchets.
	DANGER! High voltage.	DANGER! Haute tension.
	WARNING! To reduce the chance of electrical shock, do not remove covers that require tool access. No user-serviceable parts are inside. Refer servicing to Thermo Fisher Scientific qualified service personnel.	AVERTISSEMENT! Pour éviter les risques d'électrocution, ne pas retirer les capots dont l'ouverture nécessite l'utilisation d'outils. L'instrument ne contient aucune pièce réparable par l'utilisateur. Toute intervention doit être effectuée par le personnel de service qualifié venant de chez Thermo Fisher Scientific.

General instrument safety



WARNING! PHYSICAL INJURY HAZARD. Use this product only as specified in this document. Using this instrument in a manner not specified may result in personal injury or damage to the instrument.

Operating the instrument

Ensure that anyone who operates the instrument has:

- Received instructions in both general safety practices for laboratories and specific safety practices for the instrument.
- Read and understood all applicable Safety Data Sheets (SDSs). See “Safety Data Sheets (SDS)”.

Safety precautions

Do not install the instrument in heavy humidity such as a greenhouse or an incubator to avoid a danger of electric shock. If water or other material enters the instrument, the adaptor, or power inlet, disconnect the power cord and contact a service person. For operating environment, refer to “Product specifications” (page 8).

- Do not press the main plug or power cord with wet hands.
- Always ensure that the power supply input voltage matches the voltage available in your location.
- Do not install the instrument on a slant or a place prone to vibrations, which induces the risk of instrument malfunction or damage of the instrument.
- Plug the power cord firmly into the wall outlet and the instrument.
- To avoid potential shock hazard, make sure that the power cord is properly grounded.
- Be sure to position the equipment such that it is easy to disconnect the instrument.
- If the instrument is broken or dropped, disconnect the power cord and contact technical services. Do not disassemble the instrument.
- Use only authorized accessories (adaptor, power cord, and USB drive).
- For operating environment, see “Product specifications” (page 8).
- If the instrument emits smoke, disconnect the power cord from the wall outlet and contact technical services.

Cleaning or decontaminating the instrument



CAUTION! Using cleaning or decontamination methods other than those recommended by the manufacturer may compromise the safety or quality of the instrument.

Removing covers or parts of the instrument



CAUTION! PHYSICAL INJURY HAZARD The instrument is to be serviced only by trained personnel or vendor specified in the user guide. Do not remove any covers or parts that require the use of a tool to obtain access to moving parts.

Chemical safety

Chemical hazard warning



WARNING! CHEMICAL HAZARD. Before handling any chemicals, refer to the Safety Data Sheet (SDS) provided by the manufacturer, and observe all relevant precautions.



WARNING! CHEMICAL STORAGE HAZARD. Never collect or store waste in a glass container because of the risk of breaking or shattering. Reagent and waste bottles can crack and leak. Each waste bottle should be secured in a low-density polyethylene safety container with the cover fastened and the handles locked in the upright position. Wear appropriate eyewear, clothing, and gloves when handling reagent and waste bottles.

General safety guidelines

To minimize the hazards of chemicals:

- Read and understand the Safety Data Sheets (SDSs) provided by the chemical manufacturer before you store, handle, or work with any chemicals or hazardous materials. (See “Safety Data Sheets (SDS)”, page 111)
- Minimize contact with chemicals. Wear appropriate personal protective equipment when handling chemicals (for example, safety glasses, gloves, or protective clothing). For additional safety guidelines, consult the SDS.
- Minimize the inhalation of chemicals. Do not leave chemical containers open. Use only with adequate ventilation (for example, fume hood). For additional safety guidelines, consult the SDS.
- Check regularly for chemical leaks or spills. If a leak or spill occurs, follow the manufacturer’s cleanup procedures as recommended in the SDS.
- Comply with all local, state/provincial, or national laws and regulations related to chemical storage, handling, and disposal.

Chemical waste safety

Chemical waste hazard



CAUTION! HAZARDOUS WASTE. Refer to Safety Data Sheets (SDSs) and local regulations for handling and disposal.

Chemical waste safety guidelines

To minimize the hazards of chemical waste:

- Read and understand the Safety Data Sheets (SDSs) provided by the manufacturers of the chemicals in the waste container before you store, handle, or dispose of chemical waste. (See “Safety Data Sheets (SDS)”, page 111)
- Provide primary and secondary waste containers. (A primary waste container holds the immediate waste. A secondary container contains spills or leaks from the primary container. Both containers must be compatible with the waste material and meet federal, state, and local requirements for container storage.)
- Minimize contact with chemicals. Wear appropriate personal protective equipment when handling chemicals (for example, safety glasses, gloves, or protective clothing). For additional safety guidelines, consult the SDS.
- Minimize the inhalation of chemicals. Do not leave chemical containers open. Use only with adequate ventilation (for example, fume hood). For additional safety guidelines, consult the SDS.
- Handle chemical wastes in a fume hood.
- After emptying the waste container, seal it with the cap provided.
- Dispose of the contents of the waste tray and waste bottle in accordance with good laboratory practices and local, state/provincial, or national environmental and health regulations.

Waste disposal

If potentially hazardous waste is generated when you operate the instrument, you must:

- Characterize (by analysis, if necessary) the waste generated by the particular applications, reagents, and substrates used in your laboratory.
- Ensure the health and safety of all personnel in your laboratory.
- Ensure that the instrument waste is stored, transferred, transported, and disposed of according to all local, state/provincial, and/or national regulations.



IMPORTANT! Radioactive or biohazardous materials may require special handling, and disposal limitations may apply.

Electrical safety



DANGER! ELECTRICAL SHOCK HAZARD. Severe electrical shock can result from operating the Qubit™ Flex Fluorometer without its instrument panels in place. Do not remove instrument panels. High-voltage contacts are exposed when instrument panels are removed from the instrument.

Power



DANGER! ELECTRICAL HAZARD. Grounding circuit continuity is vital for the safe operation of equipment. Never operate equipment with the grounding conductor disconnected.



DANGER! ELECTRICAL HAZARD. Use properly configured and approved line cords for the voltage supply in your facility.



DANGER! ELECTRICAL HAZARD. Plug the system into a properly grounded receptacle with adequate current capacity.

Overvoltage rating

The Qubit™ Flex Fluorometer has an installation (overvoltage) category of II, and is classified as portable equipment.

Biological hazard safety



WARNING! BIOHAZARD. Biological samples such as tissues, body fluids, and blood of humans and other animals have the potential to transmit infectious diseases. Follow all applicable local, state/provincial, and/or national regulations. Wear appropriate protective eyewear, clothing, and gloves. Read and follow the guidelines in these publications.

ATTENTION! BIOHAZARD. Les échantillons biologiques tels que les tissus, les fluides corporels et le sang des humains et d'autres animaux ont la possibilité de transmettre des maladies infectieuses. Suivre tous les règlements municipaux, provinciaux/provincial et / ou nationales en vigueur. Porter des lunettes de protection approprié, des vêtements et des gants.

In the U.S.:

- U.S. Department of Health and Human Services guidelines published in *Biosafety in Microbiological and Biomedical Laboratories* (stock no. 017-040-00547-4; www.cdc.gov/OD/ohs/biosfty/bmbl4/bmbl4toc.htm)
- Occupational Safety and Health Standards, Bloodborne Pathogens (29 CFR§1910.1030; www.access.gpo.gov/nara/cfr/waisidx_01/29cfr1910a_01.html)
- Your company's/institution's Biosafety Program protocols for working with/handling potentially infectious materials.
- Additional information about biohazard guidelines is available at: www.cdc.gov

In the EU:

- Check your local guidelines and legislation on biohazard and biosafety precaution, and the best practices published in the World Health Organisation (WHO) Laboratory Biosafety Manual, third edition www.who.int/csr/resources/publications/biosafety/WHO_CDS_CSR_LYO_2004_11/en/

Safety, Electromagnetic compatibility (EMC), and Environmental standards

This section provides information on:

- U.S. and Canadian safety and EMC standards
- European safety, EMC, and Environmental standards
- Australian EMC standards
- China RoHS Standards

U.S. and Canadian Safety standards



The Qubit™ Flex Fluorometer has been tested to and complies with standard: UL 61010-1/CAN/CSA-C22.2 No. 61010-1, "Safety Requirements for Electrical Equipment for Measurement, Control, and Laboratory Use, Part 1: General Requirements."

UL 61010-2-081/CAN/CSA-C22.2 No. 61010-2-081, "Safety Requirements for Electrical Equipment for Measurement, Control, and Laboratory Use, Part 2-081: Particular Requirements for Automatic and Semi-Automatic Laboratory Equipment for Analysis and Other Purposes."

U.S. EMC standard

This instrument has been tested to and complies with standard 47 CFR FCC Part 15 "Radio Frequency Devices"; Subpart B "Unintentional Radiators".

Note: This equipment has been tested and found to comply with the limits for a Class A digital device, pursuant to part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses, and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation. If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures:

- Reorient or relocate the receiving antenna.
- Increase the separation between the equipment and receiver.
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected.
- Consult the dealer or an experienced radio/TV technician for help.

Canadian EMC standard

This instrument has been tested to and complies with standard ICES-001, "Industrial, Scientific and Medical (ISM) Radio Frequency Generators."

**European Safety,
EMC, and
Environmental
standards**



Safety

This instrument meets European requirements for safety (Low Voltage Directive 2014/35/EU). This instrument has been tested to and complies with standards:

IEC/EN 61010-1:2010, "Safety Requirements for Electrical Equipment for Measurement, Control and Laboratory Use, Part 1: General Requirements."

IEC/EN 61010-2-081:2015, "Safety requirements for electrical equipment for measurement, control, and laboratory use - Part 2-081: Particular requirements for automatic and semi-automatic laboratory equipment for analysis and other purposes."

EMC

This instrument meets European requirements for emission and immunity (EMC Directive 2014/30/EU). This instrument has been tested to and complies with standards:

EN 61326-1:2013 Class A, "Electrical Equipment for Measurement, Control and Laboratory Use – EMC Requirements."

EN 61000-3-2:2014, "Electromagnetic compatibility (EMC) — Part 3 – 2: Limits — Limits for harmonic current emissions (equipment input current ≤16 A per phase)."

EN 61000-3-3:2013, "Electromagnetic compatibility (EMC) — Part 3 – 3: Limits – Limitation of voltage changes, voltage fluctuations and flicker in public low-voltage supply systems, for equipment with rated current ≤ 16 A per phase and not subject to conditional connection (IEC 61000-3-3:2008)."

Restriction of Hazardous Substance (RoHS)

This instrument meets European requirements RoHS Directive 2011/65/EU.

This instrument has been tested to and complies with standard EN 50581:2012, "Technical documentation for the evaluation of electrical and electronic products with respect to restriction of hazardous substances."



Waste of Electrical and Electronic Equipment (WEEE)

This instrument meets European requirement WEEE Directive 2012/19/EU.

**Australian EMC
standards**



This instrument has been tested to and complies with standard AS/NZS CISPR 11, "Limits and Methods Measurement of Electromagnetic Disturbance Characteristics of Industrial, Scientific, and Medical (ISM) Radio frequency Equipment."

**China RoHS
standards**



Environment Friendly Use Period 25 Years

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- Worldwide contact telephone numbers
- Product support, including:
 - Product FAQs
 - Software, patches, and updates
 - Training for many applications and instruments
- Order and web support
- Product documentation, including:
 - User guides, manuals, and protocols
 - Certificates of Analysis
 - Safety Data Sheets (SDSs; also known as MSDSs)

Safety Data Sheets (SDS) Safety Data Sheets (SDSs) are available at thermofisher.com/support.



IMPORTANT! For SDSs for reagents and chemicals from other manufacturers, contact the manufacturer.

Limited product warranty Life Technologies Corporation and/or its affiliate(s) warrant their products as set forth in the Life Technologies' General Terms and Conditions of Sale found on Life Technologies' website at www.thermofisher.com/us/en/home/global/terms-and-conditions.html. If you have any questions, please contact Life Technologies at www.thermofisher.com/support.



IMPORTANT! Wiping the Qubit™ Flex Fluorometer computer (i.e., erasing the hard drive to remove all programs, files, and the operating system) voids the product warranty.
