

Qubit® Protein Assay Kits

For use with the Oubit® 2.0 Fluorometer

Table 1. Contents and storage information

Amount	Concentration	Storage*
300 μL or 1.5 mL	200X concentrate in 1,2-propanediol	Room temperature Desiccate Protect from light
60 mL or 300 mL	Not applicable	Room temperature
1 mL or 5 mL	0 ng/μL in TE buffer with 2 mM sodium azide	
1 mL or 5 mL	200 ng/μL in TE buffer with 2 mM sodium azide	• ≤4°C
1 mL or 5 mL	400 ng/μL in TE buffer with 2 mM sodium azide	
	60 mL or 300 mL 1 mL or 5 mL 1 mL or 5 mL	60 mL or 300 mL Not applicable 1 mL or 5 mL 0 ng/μL in TE buffer with 2 mM sodium azide 1 mL or 5 mL 200 ng/μL in TE buffer with 2 mM sodium azide

Introduction

The Qubit® Protein Assay Kits for use with the Qubit® 2.0 Fluorometer make protein quantitation easy and accurate. The kits provide concentrated assay reagent, dilution buffer, and pre-diluted BSA standards. Simply dilute the reagent using the buffer provided, add your sample (any volume from $1-20 \mu L$ is acceptable), then read the concentration using the Qubit® 2.0 Fluorometer. The assay is accurate for initial sample concentrations from 12.5 μg/mL to 5 mg/mL and exhibits low protein-to-protein variation (Appendix, Figure 1). The assay is performed at room temperature, and the signal is stable for 3 hours. Common contaminants, such as reducing reagents (DTT, β-mercaptoethanol), salts, free nucleotides, amino acids, solvents, or DNA, but not detergents, are well tolerated in the assay; some very slight modifications for the procedure are required for other contaminants (Appendix, Table 2). In addition to the Qubit® Protein Assay Kits described here, we also offer other kits for assaying dsDNA and RNA (Appendix, Table 3).

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Materials required but not provided

- Plastic container (disposable) for mixing the Qubit[®] working solution (step 1.3)
- Qubit* assay tubes (500 tubes, Cat. no. Q32856) or Axygen PCR-05-C tubes (VWR, part no. 10011-830)

Storing the Qubit® **Protein Assay Kits**

The Qubit[®] protein reagent and buffer are designed for room temperature storage. The Qubit[®] protein reagent is supplied in DMSO, which freezes at temperatures lower than room temperature. Store the protein standards at 4°C.

Critical assay parameters

Assay temperature

The Qubit* protein assay for the Qubit* 2.0 Fluorometer delivers optimal performance when all solutions are at room temperature (22-28°C). The Qubit® assays were designed to be performed at room temperature, as temperature fluctuations can influence the accuracy of the assay (Appendix, Figure 2). To minimize temperature fluctuations, store the Qubit® protein reagent and the Qubit® protein buffer at room temperature and insert all assay tubes into the Qubit 2.0 Fluorometer only for as much time as it takes for the instrument to measure the fluorescence; the Qubit® 2.0 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 reading, as this will warm the solution and result in a low reading.

Incubation time

To allow the Qubit* protein assay to reach optimal fluorescence, incubate the tubes for 15 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 greatest accuracy of the protein assay, the incubation time of the samples should be within 10 minutes of the incubation time of the standards.

Photobleaching of the Qubit® reagent

The Qubit reagents exhibit high photostability in the Qubit 2.0 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* 2.0 Fluorometer for multiple readings, a temporary reduction in fluorescence will be observed as the solution increases in temperature (Appendix, Figure 2). Note that the temperature inside the Qubit 2.0 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, you should remove the tube from the instrument and let it equilibrate to room temperature for 30 seconds before taking another reading.

Calibrating the Qubit® 2.0 Fluorometer

For each assay, you have the choice to run 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 should 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 standards and the samples is stable for no longer than 3 hours. See Figure 3 in the Appendix for an example of the calibration curve used to generate the quantitation results.

Handling and disposal

No data are currently available addressing the mutagenicity or toxicity of the Qubit® protein reagent (Component A). This reagent is an organic dye and is provided as a solution in 1,2-propanediol. Treat the Qubit* protein reagent with the same safety precautions as other materials with similar properties and dispose of the dye in accordance with local regulations.

Experimental Protocol

The protocol below assumes that you are preparing standards for calibrating the Qubit 2.0 Fluorometer. If you plan to use the last calibration performed on the instrument, you need fewer tubes (step 1.1) and less working solution (step 1.3). More detailed instructions on the use of the Qubit[®] 2.0 Fluorometer (corresponding to steps 1.9–1.12 and 2.1–2.6) can be found in the user manual accompanying the instrument.

1.1 Set up the required number of 0.5-mL tubes for standards and samples. The Qubit[®] protein assay requires 3 standards.

Note: Use only thin-wall, clear, 0.5-mL optical-grade real-time PCR tubes. Acceptable tubes include Qubit® assay tubes (500 tubes, Cat. no. Q32856) or Axygen™ PCR-05-C tubes (VWR, part no. 10011-830).

1.2 Label the tube lids.

Note: It is important to label the lid of each standard tube correctly as calibration of the Qubit* 2.0 Fluorometer requires that the standards be introduced to the instrument in the right order.

1.3 Make the Qubit* working solution by diluting the Qubit* protein reagent 1:200 in Qubit* protein buffer. Use a clean plastic tube each time you make the Qubit[®] working solution. **Do** not mix the working solution in a glass container.

Note: The final volume in each tube must be 200 μL. Each standard tube requires 190 μL of Qubit[®] working solution, and each sample tube requires anywhere from 180–199 μL. Prepare sufficient Qubit* working solution to accommodate all standards and samples. For example, for 7 samples, prepare enough working solution for the samples and 3 standards: ~200 µL per tube in 10 tubes yields 2 mL of working solution (10 µL of Qubit* reagent plus 1990 µL of Qubit® buffer).

- 1.4 Load 190 μL of Qubit* working solution into each of the tubes used for standards.
- 1.5 Add 10 μL of each Qubit[®] standard to the appropriate tube, then mix by vortexing 2-3 seconds. Be careful not to create bubbles.

Note: Careful pipetting is critical to ensure that exactly 10 μL of each Qubit[®] protein standard is added to 190 µL of Qubit® working solution.

1.6 Load Qubit* working solution into individual assay tubes so that the final volume in each tube after adding sample is 200 μ L.

Note: Your sample can be anywhere from $1-20 \mu L$, therefore, load each assay tube with a volume of Qubit[®] working solution anywhere from 180–199 μL.

1.7 Add each of your samples to the assay tubes containing the correct volume of Qubit* working solution (prepared in step 1.6), then mix by vortexing 2-3 seconds. The final volume in each tube should be 200 µL.

- **1.8** Allow all tubes to incubate at room temperature for 15 minutes.
- 1.9 On the Home Screen of the Qubit® 2.0 Fluorometer, press Protein. The Standards Screen is displayed.

Note: If you have already performed a calibration for the selected assay, the Qubit 2.0 Fluorometer prompts you to choose between reading new standards and using the previous calibration. See Calibrating the Qubit* 2.0 Fluorometer above for calibration guidelines.

1.10 On the Standards Screen, select to run a new calibration or to use the last calibration:

Press **Yes** to run a new calibration, then:

Insert the tube containing Standard #1 in the Qubit* 2.0 Fluorometer, close the lid, then press **Read**. The reading takes approximately 3 seconds. Remove Standard #1.

Insert the tube containing Standard #2 in the Qubit* 2.0 Fluorometer, close the lid, then press Read. Remove Standard #2.

Insert the tube containing Standard #3 in the Qubit* 2.0 Fluorometer, close the lid, then press Read. Remove Standard #3.

OR

Press No to use the last calibration. The Sample Screen is displayed. Insert a sample tube into the Qubit® 2.0 Fluorometer, close the lid, then press **Read**.

After the measurement is completed, the result is displayed on the screen.

Note: The value given by the Qubit® 2.0 Fluorometer at this stage corresponds to the concentration after your sample was diluted into the assay tube. You can record this value and perform the calculation yourself to find out the concentration of your original sample (see Calculating the concentration of your sample below) or the Qubit* 2.0 Fluorometer performs this calculation for you (see *Dilution Calculator* on page 5).

- 1.11 To read the next sample, remove the sample from the Qubit[®] 2.0 Fluorometer, insert the next sample, then press Read Next Sample.
- **1.12** Repeat sample readings until all samples have been read.

Calculating the concentration of your sample

The Qubit[®] 2.0 Fluorometer gives values for the Qubit[®] protein assay in μg/mL. This value corresponds to the concentration after your sample was diluted into the assay tube. To calculate the concentration of your sample, use the following equation:

Concentration of your sample = QF value ×
$$\left(\frac{200}{x}\right)$$

where QF value = the value given by the Qubit[®] 2.0 Fluorometer x = the number of microliters of sample you added to the assay tube

This equation generates a result with the same units as the value given by the Qubit 2.0 Fluorometer (that is, if the Qubit[®] 2.0 Fluorometer gave a concentration in μg/mL, the result of the equation will be in $\mu g/mL$).

Dilution Calculator

- 2.1 After the sample measurement is completed, press Calculate Stock Conc. The Dilution Calculator Screen containing the volume roller wheel is displayed.
- 2.2 Using the volume roller wheel, select the volume of your original sample that you added to the assay tube. When you stop scrolling, the Qubit* 2.0 Fluorometer calculates the original sample concentration based on the measured assay concentration.
- **2.3** To change the units in which the original sample concentration is displayed, press $\mu g/mL$. A pop-up window opens, showing the current unit selection (indicated by a red dash).
- 2.4 Select the unit for your original sample concentration by touching the desired unit in the unit selection pop-up window. To close the unit selection pop-up window, touch anywhere on the screen outside the pop-up.

The Qubit 2.0 Fluorometer automatically converts the units to your selection once the unit selection pop-up window is closed.

Note: The unit button next to your sample concentration reflects the change in the units (for example, if you change the unit to pg/ μ L, the button displays pg/ μ L).

- 2.5 To save the data from your calculation to the Qubit 2.0 Fluorometer, press Save on the Dilution Calculator screen. The last calculated value of your measurement will be saved as a *.csv file and tagged with a time and date stamp.
- 2.6 To exit the Dilution Calculator screen, press any navigator button on the bottom of the screen or Read Next Sample.

Note: When you navigate away from the Dilution Calculator screen, the Qubit[®] 2.0 Fluorometer saves the last values for the sample volume and the units in the Dilution Calculator screen only. Returning to the Dilution Calculator screen displays these last selected values.

Appendix

Protein-to-protein variation of the Qubit® Protein Assay

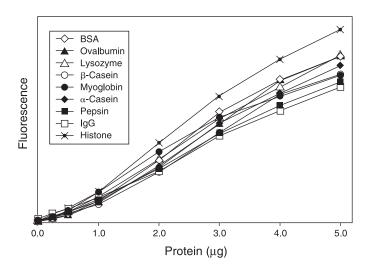


Figure 1. Low protein-to-protein variation in the Qubit® protein assay (Q33211, Q33212). Solutions of the following proteins were prepared, diluted, and assayed in the Qubit® protein assay: bovine serum albumin (BSA), chicken-egg ovalbumin, chicken-egg lysozyme, bovine-milk β -casein, equine myoglobin, bovine-milk α -casein, porcine pepsin, mouse immunoglobulin (IgG), and calf-thymus histone. Fluorescence was measured at 485/590 nm and plotted versus the mass of the protein sample. At 3 µg, the fluorescence variation was 12.4%, or 8.7% excluding the basic histone protein. Background fluorescence has not been subtracted.

Effect of temperature on the **Qubit® Protein Assay**

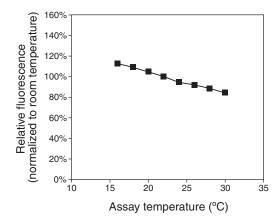


Figure 2. Plot of fluorescence versus temperature for the Qubit® protein assay. The Qubit® assays were designed to be performed at room temperature, as temperature fluctuations can influence the accuracy of the assay.

How the Qubit® 2.0 Fluorometer calculates concentration

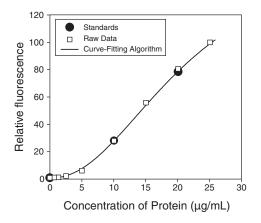


Figure 3. The curve-fitting algorithm used to determine concentration in the Qubit® protein assay. The Qubit® 2.0 Fluorometer generates concentration data based on the relationship between the two standards used in the calibration.This plot shows the line corresponding to the curve-fitting algorithm (a modified Hill plot) used in the calculation of concentration data for the Qubit® protein assay. For reference, the positions of the standards and a set of data points from an actual experiment are shown superimposed onto the line, demonstrating that the curve-fitting algorithm gives accurate values for quantitation.

Contaminants tolerated by the Qubit® Protein Assay

Table 2. Effect of contaminants in the Qubit® protein assay, tested over the range of 1.25 μg/mL to 25 μg/mL*

Contaminant	Final concentration in the assay	Concentration in 20-µL sample	Concentration in 10-µL sample	Result	
Sodium chloride	20 mM	20 mM 200 mM 400 ml		OK†	
Magnesium chloride	2 mM	20 mM	40 mM	ОК	
Potassium chloride‡	200 mM	200 mM	400 mM	ОК	
Calcium chloride‡	2 mM	20 mM	40 mM	ОК	
Ammonium sulfate	5 mM	50 mM	100 mM	OK†	
DTT	1 mM	10 mM	20 mM	OK†	
β-Mercaptoethanol	1 mM	10 mM	20 mM	ОК	
EDTA	1 mM	10 mM	20 mM	ОК	
Sodium azide	1 mM	10 mM	20 mM	ОК	
HEPES, pH 7.4	5 mM	50 mM	100 mM	OK	
Potassium phosphate, pH 7.4	5 mM	50 mM	100 mM	OK	
PBS, pH 7.4	1 mM KPO₄ 15 mM NaCl	10 mM KPO₄ 150 mM NaCl	20 mM KPO ₄ 300 mM NaCl	Protocol modification required***	
Sucrose	50 mM	50 mM 500 mM 1 M		ОК	
Sucrose	100 mM	1 M	2 M	NR	
Glycerol	1%	10%	20%	OK†	
Imidazole	1.25 mM	12.5 mM	25 mM	ОК	
SDS	0.01%	0.1%	0.2%	OK†	
SDS	0.02%	0.2%	0.4%	NR	
Tween 20	0.001%	0.01%	0.02%	NR	
Triton® X-100	0.001%	0.01%	0.02%	NR	
Amino acids§	100 μg/mL	1 mg/mL	2 mg/mL	ОК	
dNTPs**	100 μΜ	1 mM	2 mM	OK†	
DNA	5 μg/mL	50 μg/mL	100 μg/mL	OK†	
DNA	10%††	10%††	10%††	ОК	
DNA	50%††	50%††	50%††	NR	

*BSA standards were assayed in the presence or absence of contaminants at the indicated final concentrations. Equivalent $concentrations \ (approximate) \ in \ 20-\mu L \ or \ 10-\mu L \ sample \ volumes \ are \ also \ listed. \ Results \ are \ given \ as \ OK \ (usually < 10\% \ listed)$ $perturbation) or as NR (not recommended). \\ \dagger An acceptable result, but with some distortion of the standard curve. For a contract of the standard curve is the standard curve$ best results, add the same amount of contaminant to the standard samples. \$A precipitate was observed. \$A mixture of 19 amino acids. **A mixture of dATP, dCTP, dGTP, and dTTP. ††For each data point, the DNA mass was a fixed percentage of the protein mass. ***For accurate results, add the same amount of PBS to standard samples.

Qubit® Assay Kits compatible with the **Qubit® 2.0 Fluorometer**

A number of fluorescence-based quantitation kits are available for use with the Qubit* 2.0 Fluorometer. Table 3 can help you choose a kit based on the target molecule being measured and the number of assays you require.

Table 3. Qubit® Assay Kits for use with the Qubit® 2.0 Fluorometer

Product	Catalog no.	Number of assays*	Target	Notes	
Qubit® dsDNA BR Assay Kit	Q32850	100	dsDNA	 Core range (high confidence): 0.01 μg/mL to 5 μg/mL† Extended range (moderate confidence): 5 μg/mL to 10 μg/mL† Useful for quantitation of genomic and miniprep DNA samples 	
Qubit® dsDNA BR Assay Kit	Q32853	500		Accurate in the presence of RNA, salts, solvents, proteins, and free nucleotides	
Qubit® dsDNA HS Assay Kit	Q32851	100	d-DNA	4-0010	Core range (high confidence): 1 ng/mL to 500 ng/mL† Extended ranges (moderate confidence): 0.5 ng/mL to 1 ng/mL and 500 ng/mL to 600 ng/mL† Usoful for quantitation of RCR products virial DNA and complet for
Qubit® dsDNA HS Assay Kit	Q32854	500	dsDNA	 Useful for quantitation of PCR products, viral DNA, and samples for subcloning Accurate in the presence of RNA, salts, solvents, proteins, and free nucleotides 	
Qubit® ssDNA Assay Kit	Q10212	100	ssDNA	 Core range (high confidence): 5 ng/mL to 1000 ng/mL† Extended ranges (moderate confidence): 1 ng/mL to 5 ng/mL and 1000 ng/mL to 1200 ng/mL† Useful for quantitation of oligos, primers, denatured DNA, PCR products Accurate in the presence of salts, urea, solvents, proteins, ATP, and agarose 	
Qubit® RNA Assay Kit	Q32852	100		Extended ranges (moderate confidence): 20 ng/mL to 500 ng/mL to 1000 ng/mL† Useful for quantitation of samples for microarray RT-F	 Core range (high confidence): 25 ng/mL to 500 ng/mL† Extended ranges (moderate confidence): 20 ng/mL to 25 ng/mL and 500 ng/mL to 1000 ng/mL† Useful for quantitation of samples for microarray, RT-PCR, and Northern
Qubit® RNA Assay Kit	Q32855	500	RNA	blot procedures • Accurate in the presence of DNA, salts, solvents, proteins, and free nucleotides	
Qubit® RNA BR Assay Kit	Q10210	100		 Core range (high confidence): 0.1 μg/mL to 5 μg/mL† Extended ranges (moderate confidence): 0.05 μg/mL to 0.1 μg/mL and 5–6 μg/mL† 	
Qubit® RNA BR Assay Kit	Q10211	500	RNA	 Useful for quantitation of samples for microarray, RT-PCR, and Nor blot procedures Accurate in the presence of DNA, salts, solvents, proteins, and free nucleotides 	
Qubit® Protein Assay Kit	Q33211	100	Protein	Protein	 Core range (high confidence): 1.25 μg/mL to 25 μg/mL† Extended ranges (moderate confidence): 1 μg/mL to 1.25 μg/mL and 25 μg/mL to 26 μg/mL† Little protein-to-protein difference in signal
Qubit® Protein Assay Kit	Q33212	500		 Accurate in the presence of DTT, β-mercaptoethanol, amino acids, and DNA Signal is stable for 3 hours 	

*Based on an assay volume of 200 µL. †Concentration ranges refer to the concentration of sample after dilution in the assay tube.

Product List Current prices may be obtained from our website or from our Customer Service Department.

Cat. no.	Product name	Unit size
Q33211	Qubit® Protein Assay Kit, 100 assays *0.25–5 μg* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q33212	Qubit® Protein Assay Kit, 500 assays *0.25–5 µg* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Related pro	oducts	
Q10210	Qubit® RNA BR Assay Kit, 100 assays *20–1000 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q10211	Qubit® RNA BR Assay Kit, 500 assays *20–1000 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32852	Qubit® RNA Assay Kit, 100 assays *5–100 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32855	Qubit® RNA Assay Kit, 500 assays *5–100 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q10212	Qubit® ssDNA Assay Kit, 100 assays *1–200 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32850	Qubit® dsDNA BR Assay Kit, 100 assays *2–1000 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32853	Qubit® dsDNA BR Assay Kit 500 assays *2–1000 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32851	Qubit® dsDNA HS Assay Kit, 100 assays *0.2–100 ng* *for use with the Qubit® 2.0 Fluorometer	1 kit
Q32854	Qubit® dsDNA HS Assay Kit, 500 assays *0.2–100 ng* *for use with the Qubit® 2.0 Fluorometer*	1 kit
Q32856	Qubit® assay tubes *set of 500*	1 set
Q32866	Qubit® 2.0 Fluorometer	each
Q32867	Qubit® 2.0 Fluorometer USB	each
Q32868	Qubit® 2.0 Fluorometer International Power Cord (replacement)	each

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Additional international offices are listed at www.lifetechnologies.com

These high-quality reagents and materials must be used by, or directly under the supervision of, a technically qualified individual experienced in handling potentially hazardous chemicals. Read the Safety Data Sheet provided for each product; other regulatory considerations may apply.

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