



User Manual

Disclaimer: Products are intended for research use only

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SHENTEK

Residual BHK DNA Quantitation Kit

User Guide

Version: A/0
For Research Use Only
Product No.: 1101116
Reagents for 100 Reactions

Biofargo, Inc.

(IMPORTANT: Please read this document carefully before experiment.)

1. Product information

■ Product description

SHENTEK® Residual BHK DNA Quantitation Kit is used to quantitate residual BHK host cell DNA in different stages of biopharmaceutical products, from in-process samples to final products. This kit utilizes fluorescent quantitative PCR technique to perform a rapid, specific, and reliable quantitation assay at the femtogram (fg) level. The kit provides BHK DNA Control. For extraction information, please refer to the SHENTEK® Residual Host Cell DNA Sample Preparation Kit User Guide (Product No. 1104191).

■ Kit contents and storage

WARNING: Please read the Material Safety Data Sheets (MSDSs) and follow the handling instructions. Wear appropriate protective eyewear, mask, clothing, and gloves.

Table 1. Kit components and storage

Reagent	Part No.	Quantity	Storage
BHK DNA Control	NNA029	50 μ L \times 1 tube	-20°C
qPCR Reaction Buffer	NNB001	850 μ L \times 2 tubes	-20°C, protect from light
BHK Primer&Probe MIX	NNC076	300 μ L \times 1 tube	-20°C, protect from light
DNA Dilution Buffer (DDB)	NND001	1.5 mL \times 3 tubes	-20°C

The kit components can be stored at appropriate conditions for up to 24 months.

Please check the expiration date on the labels.

■ Applied instruments, including but not limited to the following

- SHENTEK-96S Real-Time PCR System
- 7500 Real-Time PCR System
- LightCycler 480 II Real-Time PCR System

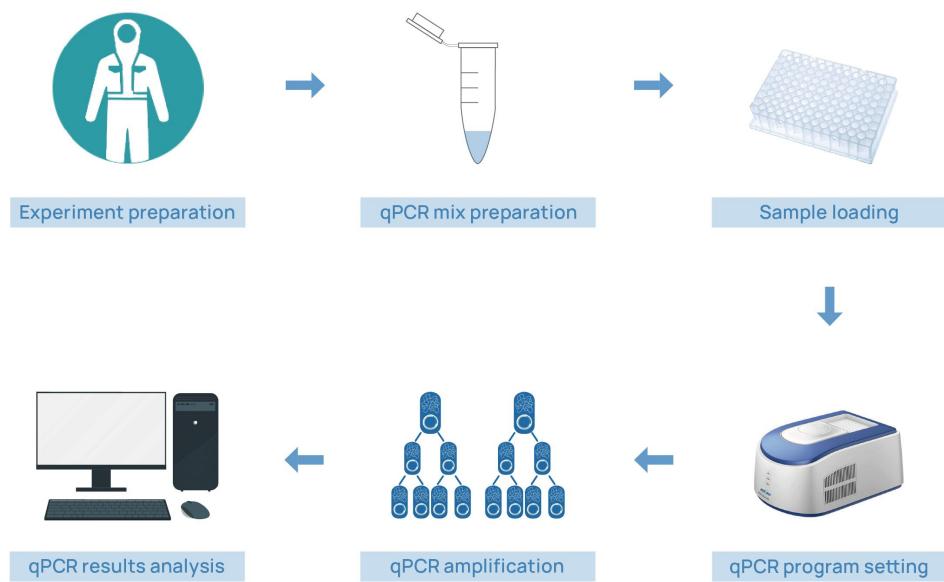
■ Required materials not included in the kit

- Nonstick, DNase-free & Low Retention Microfuge Tubes, 1.5 mL
- Nonstick, Low Retention Tips: 1000 μ L, 100 μ L and 10 μ L
- 96-well qPCR plates with sealing film or PCR 8-strip tubes with caps

■ Related equipment

- Real-Time PCR System
- Vortex mixer
- Desktop microcentrifuge
- Micropipettes: 1000 μ L, 100 μ L and 10 μ L

■ Workflow



2. Methods

■ Experiment preparation

1. Wear appropriate protective eyewear, mask, clothing and gloves.
2. Irradiate the tabletop, micropipettes and tubes with UV for 30 minutes, and

disinfect with 75% ethanol.

3. Thaw the kit completely at 2-8°C or melt on ice, vortex and spin briefly.

■ DNA Control serial dilutions for the standard curve

Please check the concentration labeled on the tube containing BHK DNA Control prior to dilution.

1. Thaw BHK DNA Control and DNA Dilution Buffer completely at 2-8°C or melt on ice. Vortex to mix well and quickly spin down the reagents for 3-5 seconds in microcentrifuge, and repeat 3 times.
2. Label six nonstick 1.5 mL microfuge tubes: ST0, ST1, ST2, ST3, ST4 and ST5.
3. Dilute the BHK DNA Control to 3000 pg/µL with DDB in the ST0 tube. Vortex to mix well and quickly spin down the reagents for 3-5 seconds in microcentrifuge, and repeat 3 times to mix thoroughly.
4. Add 90 µL DDB to each tube of ST1, ST2, ST3, ST4 and ST5.
5. Perform the serial dilutions according to Table 2:

Table 2. Dilution for BHK DNA Control

Serial dilution tube	Dilution	Conc. (pg/µL)
ST0	Dilute the DNA Control with DDB	3000
ST1	10 µL ST0 + 90 µL DDB	300
ST2	10 µL ST1 + 90 µL DDB	30
ST3	10 µL ST2 + 90 µL DDB	3
ST4	10 µL ST3 + 90 µL DDB	0.3
ST5	10 µL ST4 + 90 µL DDB	0.03

- *The remaining unused DDB need to be stored at 2-8°C. If the solution is cloudy or contains precipitates, heat at 37°C until it clear.*
- *At least five concentration of standard curve should be included. To select appropriate sample dilutions, we recommend to perform method validation before sample testing*

■ Sample preparation

➤ Test Sample Preparation

Take 100 μ L of each test sample and add to a new 1.5 mL microfuge tube.

➤ Extraction Reference Control (ERC) samples Preparation

According to the BHK DNA spike concentration in ERC samples (Take the sample containing 30 pg of BHK DNA as example), specific preparation procedure is as follows:

- (1) Take 100 μ L of test sample to a new 1.5 mL microfuge tube.
- (2) Add 10 μ L of ST3 solution and mix thoroughly, label as ERC sample.

➤ Negative Control Sample (NCS) Preparation

Add 100 μ L of DDB to a new 1.5 mL microfuge tube, and label as NCS.

NCS and samples should be prepared in same way for DNA extraction.

■ qPCR MIX preparation

1. Determine the number of reaction wells based on your selected standard curve, with the number of test samples and control samples. Generally, triplicates are tested for each sample.

Number of reaction wells = (5 standard points on the standard curve + 1 NTC + 1 NCS + test samples) \times 3

2. Prepare qPCR Mix according to the number of reaction wells in Table 3.

Table 3. qPCR MIX Preparation

Reagents	Volume/reaction	Volume for 30 reaction (includes 10% overage)
qPCR Reaction Buffer	17 μ L	561 μ L
BHK Primer&Probe MIX	3 μ L	99 μ L
Total volume	20 μ L	660 μ L

3. Mix thoroughly and place on ice, aliquot 20 μ L/well into 96-well qPCR plate or PCR 8-strip tubes.

■ qPCR Reaction MIX preparation

1. Prepare qPCR Reaction MIX according to Table 4 and 96-well plate layout as shown in Table 5.

Table 4. qPCR Reaction MIX Preparation

Tubes	Standard curve	NTC	NCS	Test sample
qPCR MIX	20 μ L	20 μ L	20 μ L	20 μ L
Samples	10 μ L ST1 - ST5	10 μ L DDB	10 μ L purified NCS	10 μ L purified test sample
Total Volume	30 μ L	30 μ L	30 μ L	30 μ L

Table 5. Example of 96-well Plate layout

ST5	ST5	ST5		S1	S1	S1	S1 ERC	S1 ERC	S1 ERC		NCS	A
ST4	ST4	ST4		S2	S2	S2	S2 ERC	S2 ERC	S2 ERC		NCS	B
ST3	ST3	ST3		S3	S3	S3	S3 ERC	S3 ERC	S3 ERC		NCS	C
ST2	ST2	ST2		S4	S4	S4	S4 ERC	S4 ERC	S4 ERC			D
ST1	ST1	ST1		S5	S5	S5	S5 ERC	S5 ERC	S5 ERC		NTC	E
											NTC	F
											NTC	G
												H
1	2	3	4	5	6	7	8	9	10	11	12	

- This example represents the assay for a standard curve with 5 concentration gradients (ST1-ST5), 1 NTC, 1 NCS, 5 test samples (S1-S5) and 5 ERC samples (S1 ERC- S5 ERC), with 3 replicates for each sample.
- In specific testing, the plate layout for sample loading can be adjusted based on the sample quantity. Please refer to the example shown in Table 5.

2. Seal the 96-well plate with sealing film. Mix well in microplate shaker, then spin down the reagents for 10 seconds in microcentrifuge and place it on the qPCR instrument.

■ qPCR program setting

NOTE: The following instructions apply only to the ABI7500 instrument with SDS v1.4. If you use a different instrument or software, refer to the applicable instrument or software documentation.

1. Create a new document, then in the Assay drop-down list, select Standard Curve (**Absolute Quantitation**).
2. In the Run Mode drop-down list, select **Standard 7500**, then click **Next**.
3. Click **New Detector**:
 - a. Enter BHK-DNA in the Name field.
 - b. Select **FAM** in the Reporter Dye drop-down list and select **(none)** in the Quencher Dye drop-down list, then click **OK**.
 - c. Select a color for the detector, then click **Add** to add the detectors to the document.
4. Select **ROX** as the passive reference dye, then Click **Next**.
5. Select the applicable set of wells for the samples, then select BHK-DNA detector for each well.
6. Select Finish, and then set thermal-cycling conditions:
 - a. Set the thermal cycling reaction volume to 30 μ L.
 - b. Set the temperature and time as following (Table 6):

Table 6. qPCR running temperature and time

Step	Temp.	Time(mm:sec)	Cycles
Activation	95°C	10:00	1
Denaturation	95°C	00:15	40
Annealing/extension	60°C*	1:00	

*Instrument will read the fluorescence signal during this step

7. Save the document, then click **Start** to start the real-time qPCR run.

■ Results analysis

1. Select **Set up** tab, then set tasks for each sample type by clicking on the Task

Column drop-down list:

- a. NTC: target DNA detector task = **NTC**
- b. NCS, test samples, and ERC wells: target DNA detector task = **Unknown**

2. Set up the standard curve as shown in table 7:

Table 7. Settings for Standard curve

Tube label	Task	Quantity (pg/µL)
ST1	Standard	300
ST2	Standard	30
ST3	Standard	3
ST4	Standard	0.3
ST5	Standard	0.03

- 3. Select the **Results** tab, then select Amplification Plot.
- 4. In the Data drop-down list, select **Delta Rn vs Cycle**.
- 5. In the Analysis Settings window, enter the following settings:
 - a. Select **Manual Ct**.
 - b. In the Threshold field, BHK-DNA enter 0.02.
 - c. Select **Automatic Baseline**.
- 6. Click the button  in the toolbar, then wait the plate analyzing.
- 7. Select the **Result** tab> >**Standard curve** tab, then verify the Slope, Intercept and R² values.
- 8. Select the **Report** tab, then achieve the mean quantity and standard deviation for each sample.
- 9. Select **File > > Export > > Results**. In the Save as type drop-down list, select **Results Export Files**, then click **Save**.
- 10. In the Report panel of Results, the 'Mean Quantity' column shows the detection values of NTC, NCS, test sample, and ERC sample, in pg/µL.
- 11. The recovery rate of ERC samples is calculated based on the value of test samples and the ERC samples. The recovery rates should be between 50% and 150%.

12. The Ct value of NCS should be larger than the mean Ct value of the lowest concentration in the standard curve. If the proven limit of quantification (LOQ) concentration is lower than the lowest concentration of the standard curve, the Ct value of NCS should be larger than the Ct value of LOQ.
13. The Ct value of NTC should be no less than 35.00 cycles or undetermined, or set standards based on laboratory validation results.

Note: The parameter settings of the result analysis should be configured on the specific model and the software version, and generally can also be automatically interpreted by the instrument.

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Support & Contact

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