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User Manual

Disclaimer: Products are intended for research use only

SHENTEK Biofargo

www.biofargo.com
804-529-2296
contact@biofargo.com

SHENTEK

Residual CHO DNA Quantitation Kit (2G)

User Guide

Version: A/1
For Research Use Only
Product No.: 1101100-1
Reagents for 100 Reactions

Biofargo, Inc.

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

1. Product information

■ Product description

SHENTEK® Residual CHO DNA Quantitation Kit (2G) is used to quantitate host-cell line residual DNA from CHO cell line, which is used for production of biopharmaceutical products. This kit uses duplex real-time PCR technology to performs rapid, specific, and reliable quantitation assay at the fg level. IPC(Internal Positive Control) is included in the CHO Primer&Probe MIX to evaluate the performance of each PCR reaction. 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, clothing and gloves.

Table 1. Kit components and storage

Reagent	Part No.	Quantity	Storage
CHO DNA Control	NNA001	50 µL × 1 tube	-20℃
qPCR Master MIX	NNB023	850 µL × 2 tubes	-20℃, protect from light
DNA Dilution Buffer (DDB)	NND001	1.5 mL × 3 tubes	-20℃
CHO Primer&Probe MIX (Incl IPC)	NNC114	500 µL × 1 tube	-20℃, protect from light

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
- CFX96 Real-Time PCR System

- Lightcycler 480 Real-Time PCR System

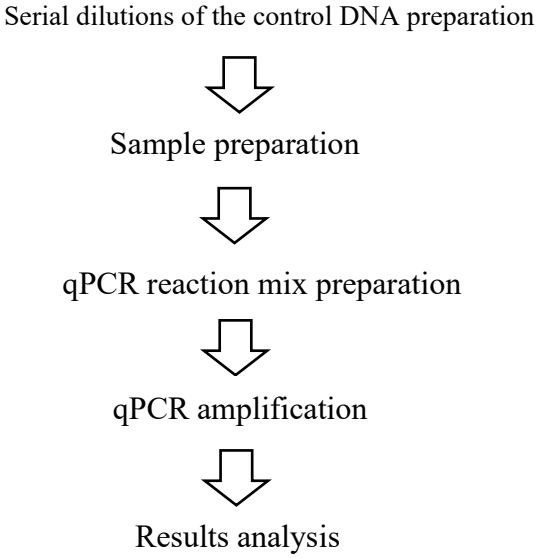
■ Required materials not included in the kit

- Nonstick, DNase-free & Low Retention Microfuge Tubes, 1.5 mL
- Nonstick, Low Retention Tips, 1000 µL, 100 µL, 10 µL
- 96-well qPCR plates or PCR 8-strip tubes

■ Related equipment

- Real-Time PCR System
- Vortex mixer
- Microplate shaker
- Pipettes, 1000 µL, 100 µL, 10 µL

■ Workflow



2. Methods

■ Experiment preparation

1. Wear appropriate protective eyewear, mask, clothing and gloves.
2. Irradiate the tabletop, pipettes 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 on the label of the tube containing the CHO DNA Control prior to dilution.

1. Thaw CHO 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 seven nonstick 1.5 mL microfuge tubes: ST0, ST1, ST2, ST3, ST4, ST5, ST6.
3. Dilute the CHO 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 it thoroughly.
4. Add 90 μL DDB to each tube: ST1, ST2, ST3, ST4, ST5, ST6.
5. Perform the serial dilutions:

Table 2. Dilution for CHO 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
ST6	10 μL ST0 +90 μL DDB	0.003

- *The remaining unused DDB need to be stored at 2-8℃. If the solution is cloudy*

or contains precipitates, heat at 37℃ until it clears.

- *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 µL of the test sample and add it to a new 1.5 mL centrifuge tube.

➢ Extraction Reference Control (ERC) samples Preparation

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

(1) Take 100 µL of the test sample to a new 1.5 mL centrifuge tube.

(2) Add 10 µL of ST3 solution and mix thoroughly, label it as the ERC sample.

➢ Negative Control Sample (NCS) Preparation

Add 100 µL of DDB to a new 1.5 mL centrifuge tube, and label it as NCS.

■ qPCR MIX preparation

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

Number of reaction wells = (6 standard points on the standard curve + 1 NTC + 1 NCS + test samples)×3

2. Prepare qPCR MIX according to the number of reaction wells.

Table 3. qPCR MIX Preparation

Reagents	Volume/reaction	Volume for 30 reaction (includes 10% overage)
qPCR Master MIX	15 µL	495 µL
CHO Primer&Probe MIX (Incl IPC)	5 µL	165 µL
Total volume	20 µL	660 µL

3. After thoroughly mixing qPCR MIX, follow 20 μL each tube is divided into PCR 8-strip tubes or 96-well qPCR plate.

■ 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

Standard curve	20 μL qPCR MIX + 10 μL ST1/ST2/ST3/ST4/ ST5/ST6
NTC	20 μL qPCR MIX + 10 μL DDB
NCS	20 μL qPCR MIX + 10 μL purified NCS
Test sample	20 μL qPCR MIX + 10 μL purified test sample
Test sample ERC	20 μL qPCR MIX + 10 μL purified ERC sample

Table 5. Example of 96-well Plate layout

NTC		S1	S1	S1	S1 ERC	S1 ERC	S1 ERC		ST6	ST6	ST6	A
NTC		S2	S2	S2	S2 ERC	S2 ERC	S2 ERC		ST5	ST5	ST5	B
NTC		S3	S3	S3	S3 ERC	S3 ERC	S3 ERC		ST4	ST4	ST4	C
		S4	S4	S4	S4 ERC	S4 ERC	S4 ERC		ST3	ST3	ST3	D
NCS		S5	S5	S5	S5 ERC	S5 ERC	S5 ERC		ST2	ST2	ST2	E
NCS									ST1	ST1	ST1	F
NCS												G
												H
1	2	3	4	5	6	7	8	9	10	11	12	

- *This example represents the assay for a standard curve with 6 concentration gradients (ST1 to ST6), 1 NTC, 1 NCS, 5 test samples (S1 to S5), 5 ERC samples (S1 ERC to S5 ERC), and 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 it well in microplate shaker, then spin down the reagents for 10 seconds in centrifuge and place it in 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 CHO-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 **Create Another**.
4. Click **New Detector**:
   a. Enter IPC in the Name field.
   b. Select **VIC** 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 **OK**.
   d. Select the detectors, then click **Add** to add the detectors to the document.
5. Select **ROX** as the passive reference dye, then Click **Next**.
6. Select the applicable set of wells for the samples, then select CHO-DNA detector and IPC detector for each well.
7. Select Finish, and then set thermal-cycling conditions:
   a. Set the thermal cycling reaction volume to 30 μL.
   b. Set the temperature and the time as following:

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

*Instrument will read the fluorescence signal during this step.

8. 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 the following table:

Table 6. 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
ST6	Standard	0.003

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, CHO-DNA enter 0.05 and IPC enter 0.1.
   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 R^2 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 can read the detection values of NTC, NCS, test sample, and ERC sample, in pg/reaction.
11. The recovery rate of ERC samples should be calculated based on the test results of the 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, and it shows normal amplification curve in the VIC signal channel.
13. The Ct value of NTC should be ≥ 35.00, or set specific standards based on the laboratory's own validation results, and it shows normal amplification curve in the VIC signal channel.

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

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Support & Contact

SHENTEK

Biofargo, Inc.
www.biofargo.com
Address: 1716 E Parham Rd Richmond, Va, 23228, USA
E-mail: contact@biofargo.com
Phone: 804-529-2296