

Product Information

ExcelTaq™ series

2X Fast Q-PCR Master Mix (SYBR, ROX)

TQ1210 200 RXN

2X Fast Q-PCR Master Mix (SYBR, ROX)	1 ml x 2
--------------------------------------	----------

TQ1211 500 RXN

2X Fast Q-PCR Master Mix (SYBR, ROX)	1 ml x 5
--------------------------------------	----------

Storage

Aliquot to avoid multiple freeze-thaw cycles

Protect from light

-20°C for 12 months

Features

- High sensitivity and signal intensity
- Compatible with fast PCR program
- With smart blue contrast dye as a visual aid for reaction setup
- Low background
- With ROX reference dye

Description

The ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX) is a ready-to-use reagent with all the essential components for quantitative real-time PCR (qPCR) except primers and templates. The master mix features high sensitivity (Fig. 1) and signal intensity as well as low background and better compatibility with fast PCR program.

The ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX) contains hot-start *Taq* polymerase in an optimized buffer with dsDNA specific SYBR green fluorescent dye. This master mix allows sensitive, precise amplification, real-time tracking of the amplification process, and simultaneous quantification for targeted DNA molecules.

With inert smart blue contrast dye, the ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX) is ready-to-use and greatly reduces pipetting errors, while largely improving the reproducibility of the process. The ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX) also includes ROX reference dye for normalizing the fluorescent reporter signal in real-time quantitative PCR.

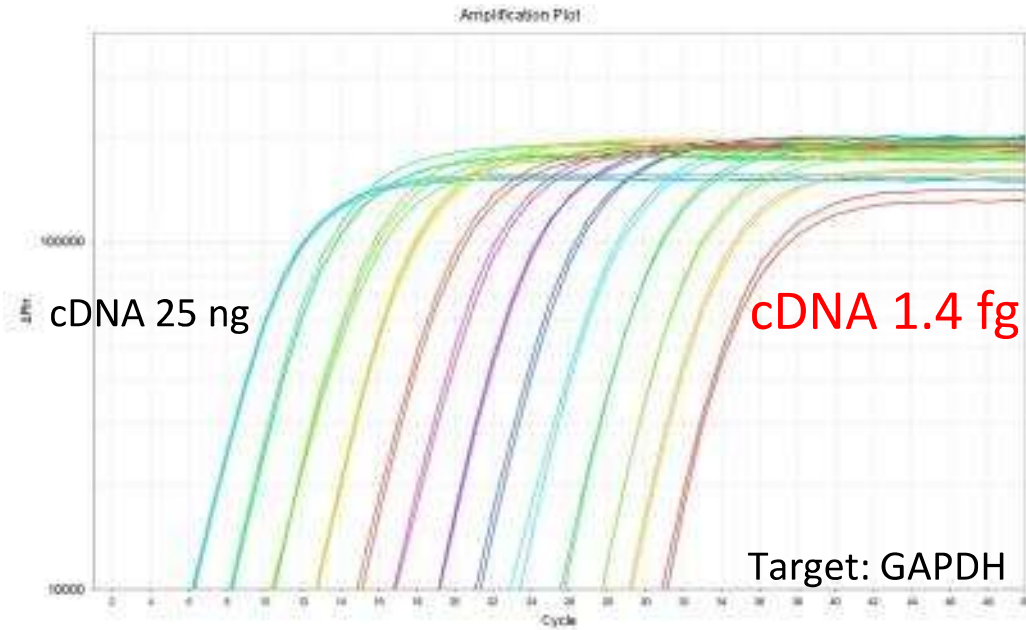


Fig. 1. The amplification plot of real-time PCR with cDNA template ranged from 25 ng to 1.4 fg in quantity, analyzed by using TQ1210 ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX) for qPCR amplification.

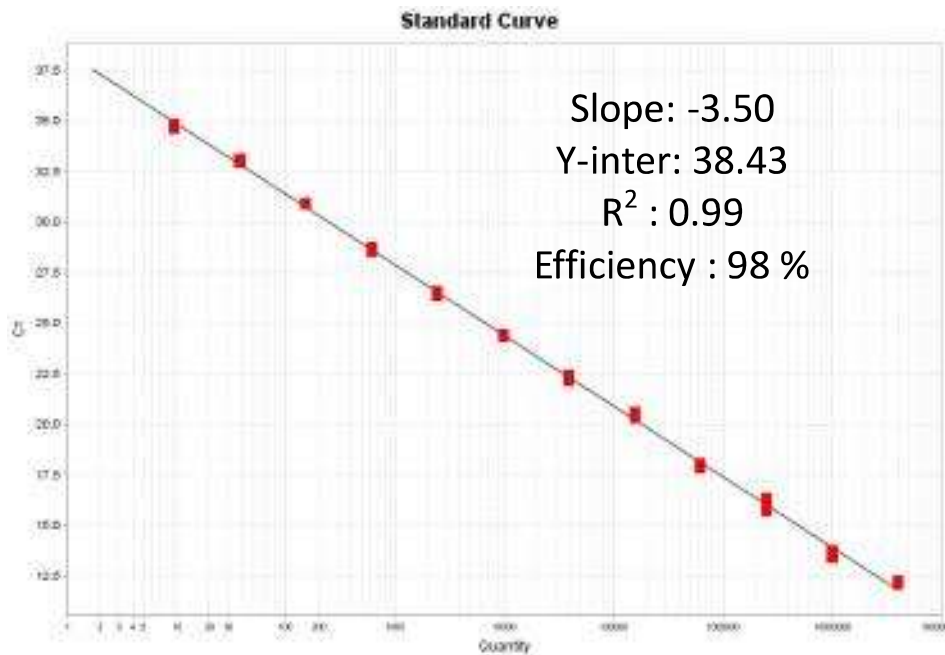


Fig. 2. The standard curve of TQ1210 ExcelTaq™ 2X Fast Q-PCR Master Mix (SYBR, ROX).

Application

- Quantitative real-time PCR
- Quantitative two-step real-time PCR

Instrument compatibility

- Applied Biosystems system:
 - 7000, 7300, 7700,
 - 7900, 7900HT, 7900HT FAST
 - StepOne™ / StepOnePlus™

The Product is also compatible with several instruments that do not require the use of ROX as following.

- BioRad system:
 - CFX96 / CFX384
 - Chromo 4™ Real-Time Detector
 - Opticon™ / Opticon™ 2
- Cepheid system:
 - Smart Cyclor®
- Eppendorf system:
 - Mastercycler® ep realplex
- Roche system:
 - Roche LightCycler® 480 / Nano
- QIAGEN system:
 - Rotor-Gene™ Q

Recommended primer design

- Amplicon size: 80-150 bp
- T_m value: around 60°C (calculated with Primer3 software)
- Primer length: 17-25 mer
- Sequence:
 - 45-55% of GC content is recommended.
 - Avoid regional high GC or AT content
 - Avoid palindrome sequence
 - Sequence with G or C at the 3' end is recommended.
- Specificity of primers should be confirmed through a BLAST search.

Recommended reaction mixture set up for qPCR

Template	2 µl*
Forward primer	50 – 400 nM**
Reverse primer	50 – 400 nM**
2X Fast Q-PCR Master Mix (SYBR, ROX)	10 µl
ddH ₂ O	to 20 µl
Total volume	20 µl

*Final template concentration varies depending on the copy number of target in the template solution. The recommended amount of template is: **100 fg -100 ng of cDNA, 80 pg -50 ng of gDNA or 10²-10⁸ molecules of plasmid.**

**The PCR primer concentration for an optimal qPCR reaction may vary according to primers' properties and template condition.

Recommended qPCR program

The Fast program is recommended for 2X Fast Q-PCR Master mix (SYBR, ROX). If the result is not good, please optimize the PCR program or try the standard program.

Fast program for qPCR

Steps	Temp.	Time	Cycles
Template denature and enzyme activation	95°C	20 sec	1
Denature	95°C	3 sec	40
Annealing /Extension	60°C	30 sec	
Melting curve analysis	Refer to instrument manual		

Standard program for qPCR

Steps	Temp.	Time	Cycles
Template denature and enzyme activation	95°C	2 min	1
Denature	95°C	15 sec	40
Annealing/Extension	60°C	60 sec	
Melting curve analysis	Refer to instrument manual		

For more information, please refer to our website:
www.smobio.com

Other Information

SMOBIO Technology, Inc. claims all warranties with respect to this document, expressed or implied, including but not limited to those of merchantability or fitness for a particular purpose. In no event shall SMOBIO Technology, Inc. be liable, whether in contract, tort, warranty, or under any statute or any other basis for special, incidental, indirect, punitive, multiple or consequential damages in connection with or arising from this document, including but not limited to the use thereof.

Caution: Not intended for human or animal diagnostic or therapeutic uses.

Related Products

CK1000	Champion E. coli Transformation Kit
DM2300	ExcelBand 100 bp+3K DNA Ladder, 500 µl
DL5000	FluoroDye DNA Fluorescent Loading Dye (Green, 6×), 1 ml
NS1000	FluoroVue Nucleic Acid Gel Stain (10,000X), 500 µl
PM2510	ExcelBand Enhanced 3-color Regular Range Protein Marker, 250 µl × 2
RP1000	ExcelRT Reverse Transcriptase, 20,000 U
RP1100	ExcelRT One-step RT-PCR Kit, 50 RXN
RI1000	RNAok RNase Inhibitor, 2000 U
TF1000	SMO-HiFi DNA Polymerase, 100 U × 1
TP1000	ExcelTaq Taq DNA Polymerase, 500 U × 1
TP1200	ExcelTaq 5X PCR Master Dye Mix, 200 RXN
TQ1110	ExcelTaq 2X Q-PCR Master Mix (SYBR, ROX), 200 RXN
TQ2110	ExcelTaq 2X Q-PCR Master Mix (TaqMan, ROX), 200 RXN

The latest version of the manual can be downloaded from www.smobio.com/shop.

For research use only

2024 ver. 2.3.1