Datasheet

Features:

- optimized realtime PCR Mastermix using probe based detection (e.g. FRET, Molecular Beacons or TaqMan)
- The Master mix contains dUTP instead of dTTP
- The qPCR / RT-PCR Mastermix DLP2 is ready-to-use and is optimized for high specificity and sensitivity because of optimized reaction buffer
- easy to us because ready-to-use Master Mix for block based PCR Cycler
- The Master Mix can be used with ROX as reference dye (1x concentrated)

Applications:

- Detection and quantification of DNA and cDNA targets
- Profiling gene expression
- Microbial detection
- Viral load determination

Description:

The Master Mix contains all reagents required for qPCR (except template and primer) in a premixed 2x concentrated ready-to-use solution. The high specificity and sensitivity of the mix is achieved by an optimized hot-start polymerase. Its activity is blocked at ambient temperature and switched on automatically at the onset of the initial denaturation. The thermal activation prevents the extension of non-specifically annealed primers and primer-dimer formations at low temperatures during PCR setup. The mix contains dUTP instead of dTTP and allows an UNG (Uracil-N-Glycosylase) treatment at the onset of thermal cycling to prevent carry-over contaminations of DNA from previous PCR reactions.

Concentration: The Master mix is 2x concentrated

List of components: Hot-Start Polymerase for qPCR, dATP, dCTP, dGTP, dUTP, optimized reaction buffer with KCl and MgCl2, Uracil-Glycosylase (UDG), stabilizers and enhancers, PCR-grade water

Transportation: with blue ice

Storage: at 4°C for 3 months, at -20°C for more than 12 months

Components	Volume per reaction	final conc.
2X qPCR Master mix DLP1	25 µl	1x
Up-stream primer (10 µM stock)	1,5 µl (range: 0,5-2.5 µl)	300 nM
Down-stream primer (10µM stock)	1,5 µl (range: 0.5-2,5 µl	300 nM
Template DNA	5 µl (0.1-15 ng/ml plasmid DNA) (1-10 µg/ml genomic DNA)	< 500ng DNA
Sterile dest. Water (included)	up to 50 µl total reaction volume	

by Bioswisstec AG
Ebnatstrasse 65
8200 Schaffhausen

visit seraglob.com
info@seraglob.com

Tel +41 52 620 33 44
Fax +41 52 620 33 45
Version 1

- vortex all solutions carefully before using and before PCR
- may you add the enzyme mix after Template DNA
- an individual optimization of annealing temperature may be necessary for new combinations of primers and Template DNA

General Thermo-Cycler protocol for qPCR / RTD-PCR Master mix:

Step	Time	Temperature
UNG treatment	2 min	50°C
Initial denaturation	1-3 min	95°C
30-40 Cycles: Denaturation Annealing Extension	 15-30 sec 30-65 sec 30 sec (per 500bp)	 95°C 55-65°C 72-75°C

Note: an individual optimization of annealing temperature may be necessary for new combinations of primers and Template DNA

Order Information

Prod. No.	Description	Quantity
S9200	qPCR/real-time-PCR Master Mix DLP2	100 rcs (2,5 ml)
S9200L	qPCR/real-time-PCR Master Mix DLP2	500 rcs (12,5 ml)
S9200XL	qPCR/real-time-PCR Master Mix DLP2	1000 rcs (25 ml)