

MegaMix Emerald 2X qPCR Mastermix, with UNG Separate ROX

P. Code	Size in 200 Rxn	Size in 1000 Rxn	Component	Description	Lot Number	Expiry
2MMEU-1	2 x 1 mL	10 x 1 mL	2X MegaMix Emerald qPCR Mastermix with UNG	Hot Start Taq, 0.2 mM dUTPs, 3 mM MgCl ₂ thermolabile UNG and intercalating dye in optimised buffer (final concentrations).		
ROX-0.1	1 x 100 µL	5 x 100 µL	25 µM ROX Reference Dye	Passive ROX reference dye		

Applications

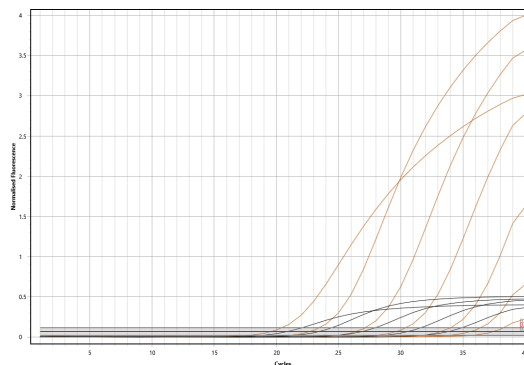
- Quantification of any DNA template (gDNA, cDNA, viral DNA), low copy number samples.
- Genotyping assays.
- Gene expression analysis.
- Pathogen detection.
- qPCR assays using fluorescence of intercalating dye.
- High resolution melting.

Key Features

- Specificity—Hot Start Taq DNA Polymerase in optimized buffer eliminates non-specific amplification and the formation of primer dimers.
- Sensitivity—Single copy detection.
- Eliminates carry over contamination—incorporation of thermolabile UNG and dUTP prevent amplicon contamination from previous runs without degrading generated amplicons.
- Versatile— compatible with standard and fast cycling conditions, GC/AT rich templates.
- Reproducibility and convenience—ready to use 2X format.
- Inhibitor Resistance—suitable for direct to PCR with products such as microLYSIS-Plus and microLYSIS-RNA.
- Third generation intercalating dye— microGREEN dye does not inhibit PCR, even at high concentrations.

Product Description

Containing all the components needed to perform qPCR swiftly and reliably. The 2X mix contains chemically modified Hot Start Taq DNA polymerase, intercalating dye, dUTP and UNG in enhancing buffer optimised for amplifying low copy DNA targets. The third generation intercalating fluorescent dye binds to double stranded DNA, making MegaMix Emerald the perfect choice for qPCR, Melt Curve Analysis and High Resolution Melting (HRM). The Hot Start Taq polymerase is chemically inactivated until heating to 95°C, providing excellent sensitivity and specificity; eliminating the formation of non-specific amplification and primer-dimers. The presence of UNG and dUTP's eliminates carryover contamination, the thermolabile UNG is active at room temperature before being completely and irreversibly inactivated when heated to 95°C, this means your PCR product is suitable for downstream processing.



MegaMix Emerald qPCR with UNG (orange) exhibits earlier Cq values and superior sensitivity vs competitor A (black), when amplifying the RNase P gene from a 6X, 1 in 10 serial dilution of 1 µg human DNA. BMS MIC.

Protocol

This product is to be used as follows.

Thaw all reagents completely and mix well before use.

Prepare a master mix as described in the table below. This reaction can be scaled according to the quantity of reactions required.

Mix gently, avoiding bubbles, centrifuge if necessary.

Include a no template control and positive control as required.

Components	Volume
2X MegaMix Emerald qPCR Mastermix with UNG	10 µL
25 µM ROX Reference Dye (Optional)	Low 0.04 µL / High 0.4 µL
Primer mix	x µL
Template	y µL
Just Water (Molecular grade water)	make up to 20 µL

The ROX concentration required will depend on the qPCR instrument in use.

Incubate the reactions at room temperature for 2-5 minutes prior to cycling to allow the UNG to eliminate carryover contamination.

Product Handling

Storage

To ensure the quality of the product until the expiry date keep at the recommended storage temperature and limit exposure to light.

Contamination Control

To prevent erroneous results ensure work environment is free of contamination by cleaning your workstation and equipment daily with a DNA decontaminant, wear gloves, use sterile tubes and filter pipette tips.

Thermocycling

Transfer the reactions to the thermal cycler and set as follows:

Annealing temperature (60°C) may require optimisation depending on the specific primers in use.

Cycles	Temperature	Time
1	95°C	2 min
40	95°C	3 sec
	60°C	20 sec

The run time can be shortened by optimising the steps of the thermocycling profile.

The included dye, microGREEN, has an absorption wavelength of 487 nm and a excitation wavelength of 511 nm, therefore acquisition can be performed in the FAM/SYBR channel of any compatible thermal cycler.

Note: Low ROX instruments require a ROX final concentration of 50 nM and high ROX instruments require a ROX final concentration of 500 nM.

For research use only

Simple | Effective | Efficient