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Produktinformation

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- Diagnostik & molekulare Diagnostik
- Laborgeräte & Service

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Lieferung & Zahlungsart

siehe unsere Liefer- und Versandbedingungen

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Product Information

Forget-Me-Not™ EvaGreen® qPCR Master Mix (Low ROX or High ROX)

Catalog Numbers

Low ROX: 31045-1mL, 31045-5mL, 31045-20mL
High ROX: 31046-1mL 31046-5mL, 31046-20mL

For low ROX instruments:

Component	31045-1mL 100 x 20 uL reactions	31045-5mL 500 x 20 uL reactions	31045-20mL 2000 x 20 uL reactions
2X Forget-Me-Not™ EvaGreen® qPCR Master Mix (Low ROX)	1 X 1 mL	5 X 1 mL	2 X 10 mL

For high ROX instruments:

Component	31046-1mL 100 x 20 uL reactions	31046-5mL 500 x 20 uL reactions	31046-20mL 2000 x 20 uL reactions
2X Forget-Me-Not™ EvaGreen® qPCR Master Mix (High ROX)	1 X 1 mL	5 X 1 mL	2 X 10 mL

Storage and Handling

Forget-Me-Not™ EvaGreen® qPCR Master Mix is shipped on blue ice and should be stored at -20°C upon arrival. Store protected from light. When stored as recommended the product is stable for at least 1 year from the date of receipt. Before use, thaw at room temperature and mix well by gentle vortexing. After thawing, the master mix should be kept on ice before use. It can be refrozen for storage, or kept at 4°C for 1 week without loss of activity.

EvaGreen® Dye

Forget-Me-Not™ features EvaGreen® dye, a unique DNA-binding dye with features ideal for both qPCR and high resolution melting (HRM) analysis. EvaGreen® dye binds to dsDNA via a novel "release-on-demand" mechanism, which permits the use of a relatively high dye concentration in qPCR without PCR inhibition (Ref 1). The absorption and fluorescence emission spectra of DNA-bound EvaGreen® dye are very similar to those of SYBR® Green I or FAM (Figure 1).

λabs/λem = 500/530 nm (DNA bound); λabs = 471 nm (without DNA)

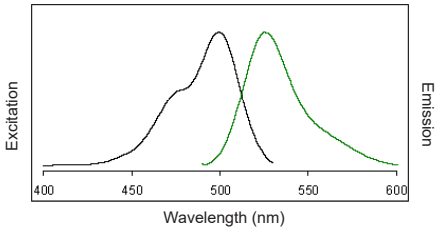

Figure 1. Excitation (left) and emission (right) spectra of EvaGreen® dye bound to dsDNA in PBS. Also see Ref. 1.

EvaGreen® dye is safer than SYBR® Green I. DNA-binding dyes are inherently dangerous due to their potential to cause mutation, but EvaGreen® dye cannot cross cell membranes, thus preventing it from coming in contact with genomic DNA in live cells. Independent labs have confirmed that EvaGreen dye is nonmutagenic, noncytotoxic and safe to aquatic life for direct disposal down the drain. Visit Biotium's website for a full EvaGreen® dye safety report.

Product Description

Forget-Me-Not™ qPCR Master Mix is a 2X hot-start EvaGreen® based master mix for use in real-time PCR applications and DNA melt curve analysis. The master mix has been formulated for fast cycling PCR parameters but can be used with regular cycling protocols.

The master mix contains EvaGreen® dye, Cheetah™ HotStart Taq DNA Polymerase, dNTPs, ROX, and a low concentration of an inert blue dye, which allows the user to visually distinguish wells containing reaction mix from empty wells, and can thereby reduce pipetting errors, saving time and reagents.

Cheetah™ HotStart Taq DNA Polymerase is Biotium's proprietary chemically-modified hot-start Taq that is completely inactive at room temperature. Cheetah™ Taq is fully activated in 2 minutes with high activity recovery, making it particularly suitable for fast PCR.

For certain instruments ROX is necessary for accurate Ct determination from well to well. Refer to Table 1 for the recommended ROX concentration (high or low) for your instrument. ROX may add noise to melt curve analysis, which could be mistaken for real peaks. Thus, in case of unexpected melt-curve peaks, un-check "ROX" in the "Passive Reference Dye" box in the software so that data is not collected from the ROX fluorescence channel, then re-analyze the data. We also offer Forget-Me-Not™ EvaGreen® qPCR Master Mix with ROX supplied separately, and Forget-Me-Not™ Probe qPCR Master Mixes (see related products).

Reaction Setup

Reaction component	Amount required per 20 uL reaction	Final concentration
2X Forget-Me-Not™ qPCR Master Mix	10 uL	1X
Primers	x uL each	0.1-0.5 uM each
Template DNA	x uL	See note [a]
H_2O	Add to 20 uL	

[a] Template concentration: The optimal amount of template DNA varies by application. Recommended amounts of genomic DNA template per reaction typically range from 50 pg to 50 ng per reaction. For two-step RT-PCR: the A_{260} measurement of a reverse transcription reaction does not accurately quantify cDNA. Add undiluted or diluted cDNA from a RT reaction (generated from < 1 µg RNA), but the RT reaction volume must not exceed 10% of the final PCR volume.

General Considerations

1) Primer design and amplicon length: For optimal results, use appropriate software to design primers with melting temperatures (Tm's) of approximately 60°C that amplify products of 60-200 bp. For longer amplicons, extension times may need to be extended.

2) EvaGreen® dye can be used for high resolution melting (HRM) analysis. Follow the qPCR system's instructions for data collection and analysis.

3) Gel electrophoresis analysis of PCR product: After PCR with EvaGreen® dye, PCR products need not be stained with another DNA gel stain. Simply add DNA loading buffer to your PCR reaction solution, load on a gel, and conduct electrophoresis as usual. Gel visualization can be carried out using a 254 nm UV box, or a blue LED imager using a SYBR® Green filter. Alternatively, the gel may be imaged using a 488 nm laser-based gel scanner.

4) Roche LightCycler® users: If using glass capillaries for reactions, add BSA to the PCR reactions at ~250 ng/uL final concentration. BSA is not necessary if plastic capillary tubes are used.

Cycling Protocols

Choice of cycling protocol depends on your instrument capability and on the nature of your amplicon. If your instrument does not support fast cycling, use the parameters recommended in your instrument manual.

A. Two-step fast cycling protocol

This cycling protocol should be applicable to most amplifications where the primer Tm's are designed to be 60°C.

Cycling Step	Temperature	Holding Time	Number of cycles
Enzyme activation	95°C	2 min	1
Denaturation	95°C	2-5 sec	40
Annealing / extension / data acquisition	60°C	20-30 sec	
Dissociation / melt curve	Set up as per instrument guidelines		

B. Three-step fast cycling protocol

Use this protocol when optimal primer annealing and extension temperatures are desired.

Cycling Step	Temperature	Holding Time	Number of cycles
Enzyme activation	95°C	2 min	1
Denaturation	95°C	2-5 sec	40
Annealing	55-65°C	10 sec	
Extension / data acquisition	72°C	10-20 sec	
Dissociation / melt curve	Set up as per instrument guidelines		

Table 1. Instrument Compatibility

Reference Dye	PCR Instrument
Low ROX (~50 nM)	Applied Biosystems®: 7500, 7500 Fast, ViiA™7, QuantStudio™ instruments Stratagene (Agilent): MX4000P, MX3000P, MX3005P
High ROX (~500 nM)	Applied Biosystems®: 5700, 7000, 7300, 7700, 7900, 7900HT, 7900HT Fast, StepOne™, StepOnePlus™
No ROX required [a]	BioRad: iCycler™, MyiQ™, MiQ™ 2, iQ™ 5, CFX-96 Touch™, CFX-384 Touch™ and Connect™, Chromo4™, MiniOpticon™ Qiagen: Rotor-Gene® Q, Rotor-Gene® 3000 & 6000 Eppendorf: Mastercycler® Realplex Illumina: Eco™ RealTime PCR System Cepheid: SmartCyler® Roche: LightCycler® 480, LightCycler® 2.0
Fluorescein [b]	BioRad: iCycler™, MyiQ™, MiQ™ 2, iQ™ 5

[a] Instruments that do not require ROX reference dye are generally compatible with qPCR master mixes containing ROX (check with the manufacturer before use). We also sell Forget-Me-Not™ EvaGreen® qPCR Master Mix kits without ROX (cat# 31041), and with a separate tube of ROX (cat# 31042) that can be used to make your own low or high ROX master mixes.

[b] Bio-Rad's iCycler™, MyiQ™, MiQ™, and iQ™ users do not need to add fluorescein to the PCR reaction as EvaGreen® dye has a slight background fluorescence that provides adequate and stable baseline level fluorescence. For these instruments we recommend using Forget-Me-Not™ EvaGreen® qPCR Master Mix without ROX (cat# 31041).

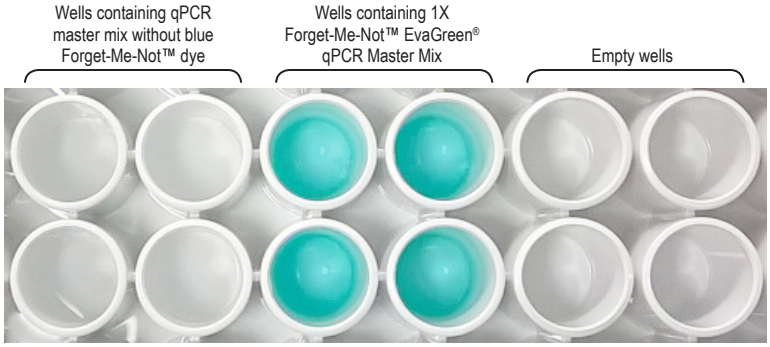

Figure 2: Forget-Me-Not™ EvaGreen® qPCR Master Mix contains a low concentration of an inert blue dye, which allows the user to visually distinguish wells containing reaction mix from empty wells, and can thereby reduce pipetting errors, saving time and reagents.

Related Products

Catalog number	Product
31043	Forget-Me-Not™ Universal Probe qPCR Master Mix
31041	Forget-Me-Not™ EvaGreen® qPCR Master Mix (no ROX)
31042	Forget-Me-Not™ EvaGreen® qPCR Master Mix (separate ROX)
31000	EvaGreen® Dye, 20X in water
29050	Cheetah™ HotStart Taq DNA Polymerase
29054	HotStart Polymerase Modification Kit
29051	EvaEZ™ Fluorometric Polymerase Activity Assay Kit
41003	GelRed® Nucleic Acid Gel Stain, 10,000X in water
41005	GelGreen® Nucleic Acid Gel Stain, 10,000X in water
31042	Ready-to-Use 100 bp DNA Ladder
31022	Ready-to-Use 1 kb DNA Ladder
41024-4L	Water, Ultrapure Molecular Biology Grade
41006	TBE Buffer, 5X
E90003	Gel-Bright™ LED Gel Illuminator

References

1) Mao, et al. Characterization of EvaGreen Dye and the implication of its physico-chemical properties for qPCR applications. BMC Biotechnology 7, 76-91 (2007).

Please visit our website at www.biotium.com for information on our life science research products, including environmentally friendly EvaGreen® qPCR master mixes, fluorescent CF® dye antibody conjugates and reactive dyes, apoptosis reagents, fluorescent probes, and kits for cell biology research.