

Product Information

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

For low ROX instruments:

| Component | 31045-1mL 100 x 20 uL reactions | 31045-5mL 500 x 20 uL reactions | 31045-10mL 1000 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 | 1 x 10 mL | 2 x 10 mL |

For high ROX instruments:

| Component | 31046-1mL 100 x 20 uL reactions | 31046-5mL 500 x 20 uL reactions | 31046-10mL 1000 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 | 1 x 10 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.

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.

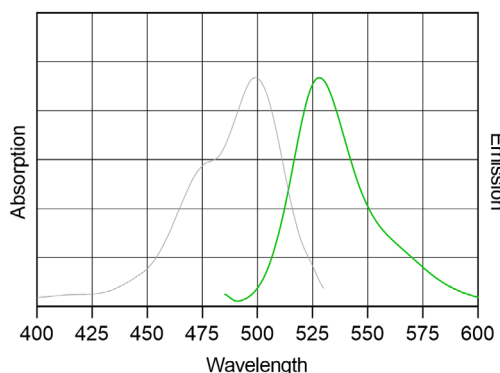


Figure 1. Absorption (left) and emission (right) spectra of EvaGreen® Dye bound to dsDNA in PBS. Also see Ref. 1.

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 (Figure 1) are very similar to those of SYBR® Green I or FAM with Ex/Em at 500/530 nm with DNA.

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 non-mutagenic, non-cytotoxic and safe to aquatic life for direct disposal down the drain. Visit Biotium’s website for a [full EvaGreen® Dye safety report](#).

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.

References

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

General Considerations

- Primer design and amplicon length: For optimal results, use appropriate software to design primers with melting temperatures (T_m) of approximately 60°C that amplify products of 60-200 bp. For longer amplicons, extension times may need to be extended.
- EvaGreen® Dye can be used for high resolution melting (HRM) analysis. Follow the qPCR system's instructions for data collection and analysis.
- Template concentration: The optimal amount of template DNA varies by application. We recommend 50 pg to 50 ng genomic DNA 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 ug RNA), but the RT reaction volume must not exceed 10% of the final PCR volume.
- Gel electrophoresis analysis of PCR products: 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.
- 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).
- 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. No. 31041), and with a separate tube of ROX (Cat. No. 31042) that can be used to make your own low or high ROX master mixes.
- 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.

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 | BioRad: iCycler™, MyiQ™, MiQ™ 2, iQ™ 5, CFX Opus, 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: SmartCycler Roche: LightCycler® 480, LightCycler® 2.0 |
| Fluorescein ^[a] | BioRad: iCycler™, MyiQ™, MiQ™ 2, iQ™ 5 |

^[a] 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. No. 31041).

Protocols for Use

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 ₂ O | Add to 20 uL | |

^[a] See General Considerations for template concentration recommendations.

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 T_m of the primers is designed to be 60°C.

| Cycling Step | Temp | 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 | Temp | 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 | | |

Related Products

| Cat. No. | 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) |
| 31078 | N-Flux™ 5X Digital PCR Master Mix |
| 31000 | EvaGreen® Dye, 20X in Water |
| 31077 | EvaGreen® Plus Dye, 20X in Water |
| 29050 | Cheetah™ HotStart Taq DNA Polymerase |
| 31079 | EvaRuby™ Dye, 20X in Water |
| 29087-50uL | VeriFluor™ Far-Red Passive Reference Dye, 400X in Water |
| 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 |
| 31085 | 100 bp DNA Ladder, Ready-to-Load |
| 31084 | 1 kb DNA Ladder, Ready-to-Load |
| 41024-4L | Water, Ultrapure Molecular Biology Grade |
| 41006 | TBE Buffer, 5X |
| E90005 | Gel-Bright™ Laser Diode Gel Illuminator |

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.

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