

TOOLS 5X EvaGreen qPCR Mix without ROX Cat. no.: TF-5EVNR250 Storage: -20°C for up to 1 year Product Size: 1 mL

Introduction

The TOOLS $5 \times$ EvaGreen qPCR Mix without ROX is optimized for real-time quantitative polymerase chain reaction (PCR) assays. The ready-to-load mix includes TOOLS HotTaq DNA polymerase, ultrapure dNTPs, MgCl₂, and EvaGreen® dye. Only water, template, and primers need to be added. TOOLS HotTaq DNA polymerase is activated by a 15-min incubation step at 95°C. This prevents extension of nonspecifically annealed primers and primer-dimers formed at low temperatures during qPCR.

EvaGreen® Dye: EvaGreen® is a DNA-binding dye for qPCR that, compared with SYBR® Green I, has similar spectra but much lower PCR inhibition. It is extremely stable and has been shown to be nonmutagenic and noncytotoxic. EvaGreen® is compatible with all common real-time PCR cyclers—simply select the standard settings for SYBR® Green or FAM

EvaGreen[®] is a registered trademark and licensed for sale by Biotium, Inc. SYBR[®] is a registered trademark of Molecular Probes, Inc.

Composition TOOLS HotTaq DNA polymerase 5× qPCR buffer E 12.5 mM MgCl₂ (1× PCR solution–2.5 mM MgCl₂) dNTPs, including dTTP to improve reaction sensitivity and efficiency compared with dUTP EvaGreen® dye

Cycler compatibility

The TOOLS 5× EvaGreen qPCR Mix without ROX can be used with all major qPCR cyclers that do not require a reference dye, including Bio-Rad iQTM 5, OpticonTM, and OpticonTM 2; Chromo 4TM, MiniOpticon, CFX96, and CFX384; Corbett Rotor-GeneTM 3000 and Rotor-GeneTM 6000; and Roche LightCycler® 480 and Techne QuanticaTM.

TOOLS 5X EVAGREEN QPCR MIX WITHOUT ROX

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Component	Volume	Concentration		
TOOLS 5X EvaGreen qPCR Mix	4 μL	1x		
Primer Forward (10 pmol/µl)	0.16–0.5 μL	80–250 nM		
Primer Reverse (10 pmol/µl)	0.16–0.5 μL	80–250 nM		
DNA template	1–5 µL	1–50 ng/µL		
H ₂ O PCR grade	up to 20 µL			
Total	20 µL			

Recommended qPCR reaction mix

Recommended qPCR cycles

Cycle step	Temp.	Time	Cycles
Initial denaturation	95°C	15 min	1
Denaturation	95°C	15 s	40
Annealing	60°-65°C	20 s	
Elongation	72°C	20 s	

Important: To activate the polymerase, incubate it at 95°C for 15 min at the beginning of the qPCR cycle.

The product is for research only and not for diagnostic and clinical use.

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