



# igScript™ Probe-Based qPCR 2x Master Mix

# Manual

Catalog #	4233	4235	4237
Package Size	500 reactions	1,000 reactions	2,500 reactions
Volume	20μΙ		



# Important!

# -20°C Storage Required

- \* Immediately inspect packages
- \* Freeze upon receipt

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# igScript™ Probe-Based qPCR 2x Master Mix

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# **Description:**

igScript<sup>™</sup> Probe-Based qPCR 2x Master Mix contains igScript<sup>™</sup> *Taq* DNA polymerase, MgCl<sub>2</sub>, dNTPs, stabilizers, enhancers and low ROX reference dye with standard buffer providing improved qPCR efficiency, wider dynamic range, superior sensitivity and specificity. igScript<sup>™</sup> qPCR 2x Master Mix is a ready-to-use cocktail containing all components except primers, probe and template, for the amplification and detection of DNA in qPCR. This 2x master mix requires minimal handling during reaction setup and offer consistent and robust qPCR reactions. *Taq* DNA Polymerase is a thermostable DNA polymerase that possesses a 5′ $\rightarrow$ 3′ polymerase (1, 2) and a 5′ $\rightarrow$ 3′ exonuclease activity (3, 4). The amplification step features a high quality *Taq* DNA Polymerase which offers robust, reliable and better amplification.

## **Product Components and Storage:**

- igScript™ Probe-Based qPCR 2x Master Mix
- -20 °C

# **Applications:**

- Gene expression data validation
- Multiplexing
- Mutation detection
- Pathogen and viral detection
- Genetically modified organisms (GMO) characterization and Genetic profiling

#### **Benefits:**

- Enhanced efficiency, specificity, and sensitivity
- Compatible with all real-time PCR instruments
- Superior gene expression results under various cycling conditions.



#### **Protocol:**

- 1. Place kit components and DNA samples on ice.
- 2. Mix and then centrifuge briefly to collect contents at the bottom of the tube.
- 3. Prepare a master mix for each reaction and control plus 10% extra to allow for pipetting error, according to the following table:

PCR Reaction Set Up:			
Template DNA	x µl		
Forward primer (5 µM)	1.0 µl		
Reverse primer (5 µM)	1.0 µl		
Probe (5 μM)	0.5 µl		
igScript™ qPCR 2x Master Mix	10 µl		
H <sub>2</sub> O up to	20.0 µl		

- 4. Mix the reaction mixture thoroughly.
- 5. Program the thermal cycler according to the manufacturer's instructions.
- 6. A typical PCR cycling program is outlined in the following table:

PCR Cycling Conditions						
Steps	Temperature	Time	Cycles			
Initial denaturation	95°C	3 min	1			
Denaturation	95°C	5 sec				
Annealing/ Extension*	55-60°C	30 sec	40			
Melting curve analysis	According to instrument guidelines					

- 7. Place the PCR tubes in the thermal cycler and start the cycling program.
- 8. Analyze the data according to manufacturer protocol.

**Note:** For 3 step cycling protocols, anneal at optimal annealing temperature for 30 sec followed by the minimum time required for data acquisition at 72 °C according to instrument guidelines.



#### **Related Products:**

- igScript™ Probe Based One Step RT-qPCR Kit (Cat.#4243, 4245, 4247)
- igScript™ One Step RT-PCR Kit (Cat.# 4211)
- igScript™ One Step RT-qPCR Kit (Cat.# 4214)
- igScript™ First Strand cDNA Synthesis Kit (Cat.# 4312)
- igScript<sup>™</sup> Reverse Transcriptase (Cat.# 3344)
- ig® SYBR Green qPCR 2x Master Mix (Cat.# 3354)

### **Ordering Information:**

- Order online within the USA. Place orders on www.intactgenomics.com using our secure Shopping Cart.
- Order by email, phone, or fax.

Email: sales@intactgenomics.com

Phone: (314) 942-3655 | Toll-free: 855-835-7172 | Fax: (314) 942-3656

• Order via our distributors.

#### **References:**

- 1. Chien, A., Edgar, D. B. and Trela, J. M. (1976). J. Bact. 127, 1550-1557.
- 2. Lawyer, F. C. et al. (1993). PCR Methods and Appl. 2, 275-287.
- 3. Longley, M. J., Bennett, S. E. and Mosbaugh D. W. (1990). Nucleic Acids Res. 18, 7317-7322.
- 4. Lyamichev, V., Brow, M. A. and Dahlberg, J. E. (1993). Science. 260, 778-783.

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Our hours are Monday - Friday, 8AM to 5PM, U.S. Central Standard Time.

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