

## RealQ Fast 2x Master Mix Green

Cat. No.: A463499  
40 reactions (25 µl)

MADE IN DENMARK

|                   | RealQ Fast<br>2x Master Mix<br>Green | ROX<br>Internal Reference Dye<br>200 µM |
|-------------------|--------------------------------------|---|
| <b>ID No.</b>     | <b>5030300-1250</b>                  | <b>5700300-0050</b>                     |
| <b>Cap colour</b> | <b>Amber</b>                         | <b>Amber</b>                            |
| <b>Content</b>    | 1 x 1.25 ml                          | 1 x 0.05 ml                             |

### Contents

RealQ Fast 2x Master Mix Green is a ready-to-use, optimized 2x master mix for dye-based real-time PCR. ROX is included for PCR real-time instruments requiring an internal reference dye.

- RealQ Fast 2x Master Mix Green: optimized buffer system, 2Taq Hot Start DNA Polymerase, dNTPs, and fluorescent dye.
- 200 µM ROX internal reference dye.

### Compatibility

All real-time PCR instruments with a FAM/SYBR filter (addition of ROX may be needed, see Table 1).

### Recommended Storage and Stability

| Temperature      | Duration                         |
|------------------|----------------------------------|
| Room temperature | Up to 3 days                     |
| 4° C             | Up to 3 months                   |
| -20° C           | Long term.<br>See expiry on tube |

Multiple freeze-thaw cycles should be avoided. Solutions containing fluorescent green DNA dye should be protected from light whenever possible.

### ROX Reference Dye

ROX is used as passive reference dye in qPCR. The addition of ROX enables fluorescence normalization which helps to achieve a higher level of precision. The required concentration of ROX depends on the real-time PCR instrument – see table below.

**Table 1:** Recommended ROX concentrations for various real-time PCR instruments

| Final ROX concentration | Real-time PCR instrument   |
|-------------------------|--|
| 0 nM                    | <b>Bio Molecular:</b> Mic qPCR Cycler<br><b>Bio-Rad:</b> iCyclerR iQ, iQ5 and MyiQ™, OpticonR, CFX 96, CFX 384<br><b>Roche:</b> LightCyclerR 480, LightCycler 96, LightCyclerR 2.0, iCycler iQ System<br><b>Corbett:</b> Rotor-Gene™ 3000, 6000, Rotor-Gene Q<br><b>Eppendorf:</b> MasterCycler™ ep <i>realplex</i><br><b>Cepheid:</b> Smart Cycler<br><b>MyGo:</b> Mini and Pro |
| 30 nM                   | <b>Applied Biosystems:</b> ABI 7500 and ABI 7500 Fast, ABI ViiA7<br><b>Agilent:</b> Mx3000™, Mx3005P™, and Mx4000™, Mx4000R, AriaMx  |
| 300 nM                  | <b>Applied Biosystems:</b> ABI 5700, 7000 PRISM, 7300, 7700, 7900, 7900HT and 7900HT Fast, StepOne™, StepOnePlus™  |

### Protocol

Allow master mix to reach room temperature. Ensure sufficient mixing of the master mix (i.e. by vortexing) prior to reaction assembly.

- Combine master mix, primers, template DNA and water according to the following table.

**Table 2:** Recommended reaction setup

| Component                  | Vol./reaction       | Final concentration*   |
|----------------------------|---------------------|--|
| RealQ Fast 2x Master Mix   | 12.5 µl             | 1x   |
| Primer A (10 µM)           | 0.5 µl (0.2 – 1 µl) | 0.2 µM (0.1 – 0.5 µM)**  |
| Primer B (10 µM)           | 0.5 µl (0.2 – 1 µl) | 0.2 µM (0.1 – 0.5 µM)**  |
| 200 µM ROX                 | X µl***             | 30 nM – Low ROX (Optional)<br>300 nM – High ROX (Optional)   |
| PCR-grade H <sub>2</sub> O | X µl                | -  |
| Template DNA               | X µl                | genomic DNA: 20 ng (1 – 100 ng)<br>plasmid DNA: 0.5 ng (0.1 – 1 ng)<br>bacterial DNA: 5 ng (1 – 10 ng) |
| <b>Total volume****</b>    | 25 µl               | -  |

\* Suggested starting conditions, optimisation range in parenthesis.

\*\* Optimization of primer and probe concentrations is highly recommended.

\*\*\* Prepare fresh 1:10 or 1:100 dilutions of 200 µM ROX with PCR-grade H<sub>2</sub>O to increase transferred volume.

\*\*\*\* Reaction volumes < 10 µl is not recommended. Smaller reaction volumes decrease signal intensity.

- Mix the components thoroughly, then centrifuge to collect liquid at the bottom of the tube.
- Transfer the appropriate volume to an optical plate or strip compatible with the chosen real-time PCR instrument. Seal the plate / strip.
- Set up PCR program using one of the following settings:

**Table 3:** Standard 2-step cycling conditions.

| STANDARD | Step                         | Temperature | Duration | Cycle |
|----------|------------------------------|-------------|----------|-------|
|          | Activate enzyme <sup>a</sup> | 95°C        | 2 min    | 40    |
|          | Denature                     | 95°C        | 15 sec   |       |
|          | Anneal/extend <sup>b</sup>   | 60°C        | 60 sec   |       |

**Table 4:** Fast 2-step cycling conditions.

| FAST | Step                         | Temperature | Duration | Cycle |
|------|------------------------------|-------------|----------|-------|
|      | Activate enzyme <sup>a</sup> | 95°C        | 2 min    | 40    |
|      | Denature                     | 95°C        | 5 sec    |       |
|      | Anneal/extend <sup>b</sup>   | 60°C        | 30 sec   |       |

**Table 5:** Super-fast 2-step cycling conditions. Sensitivity may be affected by fast cycling conditions.

| SUPER-FAST | Step                         | Temperature | Duration | Cycle |
|------------|------------------------------|-------------|----------|-------|
|            | Activate enzyme <sup>a</sup> | 95°C        | 2 min    | 40    |
|            | Denature                     | 95°C        | 5 sec    |       |
|            | Anneal/extend <sup>b</sup>   | 60°C        | 15 sec   |       |

<sup>a</sup> Can be reduced to 30 sec for non-complex templates.

<sup>b</sup> Choose an appropriate annealing temperature for the primer set used.

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Other product sizes, combinations and customized solutions are available. Please look at [www.ampliqon.com](http://www.ampliqon.com) or ask for our complete product list for PCR Enzymes. For customized solutions please contact us.

Made in Denmark