

# Data Sheet

## FIREPol<sup>®</sup> Master Mix with 12.5 mM MgCl<sub>2</sub>, 5x

Cat. No.	Pack Size	20 µl rxn
04-11-00S25	0.1 ml	25
04-11-00125	1 ml	250
04-11-00125-5	5 x 1 ml	1250
04-11-00125-10	10 x 1 ml	2500

For *in vitro* use only

### Description:

FIREPol<sup>®</sup> Master Mix is a 5x-concentrated ready-to-use solution containing all reagents required for PCR (except template, primers and water).

### Applications:

- Suited for a wide range of PCR assays
- TA cloning

### Mix Composition:

- FIREPol<sup>®</sup> DNA polymerase
- 5x Reaction Buffer B  
*0.4 M Tris-HCl, 0.1 M (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>, 0.1% w/v Tween-20*
- 12.5 mM MgCl<sub>2</sub>  
*1 x PCR solution – 2.5 mM MgCl<sub>2</sub>*
- 1 mM dNTPs of each  
*1 x PCR solution – 200 µM dATP, 200 µM dCTP, 200 µM dGTP and 200 µM dTTP*

### Shipping and Storage conditions:

Routine storage: -18°C to -28°C

Shipping and temporary storage for up to 1 month at room temperature or storage for up to 6 months at 2-8°C has no detrimental effects on the quality of the product.

Manufactured by Solis BioDyne in compliance with the ISO 9001 and ISO 13485 certified Quality Management System.

**FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES.**

### Recommendations:

We recommend using FIREPol<sup>®</sup> Master Mix in any PCR application that will be visualized by agarose gel electrophoresis and ethidium bromide staining.

In order to prevent contamination, we recommend you to setup the reaction under laminar or in PCR box.

### Recommended PCR reaction mix:

Component	Volume	Final conc.
FIREPol <sup>®</sup> Master Mix (5x)	4 µl	1x
Forward primer (10 µM)	0.2–0.6 µl	0.1–0.3 µM
Reverse primer (10 µM)	0.2–0.6 µl	0.1–0.3 µM
DNA template	variable	variable <sup>1</sup>
H <sub>2</sub> O	Up to 20 µl	

<sup>1</sup>Conc. of cDNA 0.01 pg/µl -0.1 ng/µl; gDNA 0.1 ng/µl – 10 ng/µl

### Recommended PCR cycles:

Operation	Temp.	Time	Cycles
<b>Initial denaturation<sup>2</sup></b>	<b>95°C</b>	<b>3–5 min</b>	1
Denaturation	95°C	15–30 s	25–30
Annealing <sup>3</sup>	54–66°C	30–60 s	
Elongation <sup>4</sup>	72°C	40 s–4 min	
Final elongation	72°C	5–10 min	

<sup>2</sup> Complex templates, such as gDNA, require longer time to denature (5 min). With low complexity templates (i.e. lambda, plasmid DNA), initial denaturation time can be reduced to 3 min.

<sup>3</sup> The annealing temperature (T<sub>a</sub>) depends on the melting temperature (T<sub>m</sub>) of the primers. A T<sub>a</sub> that is about 2 to 5°C lower than the T<sub>m</sub> of the primers is generally suitable. Performing temperature gradient is recommended.

<sup>4</sup> Elongation time depends on the length of the fragment to be amplified. A time of 1 min/kb is recommended.

**Safety warnings and precautions:**

This product and its components should be handled only by persons trained in laboratory techniques. It is advisable to wear suitable protective clothing, such as laboratory overalls, gloves and safety glasses. Care should be taken to avoid contact with skin or eyes. In case of contact with skin or eyes, wash immediately with water. Refer to Safety Data Sheet for more information.

**Technical support:**

Contact your sales representative for any questions or send an email to [support@solisbiodyne.com](mailto:support@solisbiodyne.com)

Online chat is available at [www.solisbiodyne.com](http://www.solisbiodyne.com)

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Solis BioDyne

Teaduspargi 9, 50411 Tartu, Estonia, tel: +372 740 9960, fax: +372 740 2079, e-mail: [info@solisbiodyne.com](mailto:info@solisbiodyne.com), [www.solisbiodyne.com](http://www.solisbiodyne.com)