

PCRMax™

# Lyophilised 2X qPCR Master Mix

Instructions for use of Lyophilised Master Mix

The logo for PCRmax features the text "PCRmax" in a blue, sans-serif font. The "max" portion is smaller and positioned to the right of "PCR". To the right of the text is a decorative graphic consisting of three parallel, curved lines that sweep upwards and to the right, resembling a stylized wave or a graph line.

PCRmax

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# Introduction

Our Lyophilised 2X qPCR Master Mix is optimised for use in real-time PCR. The product is stable at ambient temperatures for at least 18 months and can be conveniently shipped and stored at room temperature. The master mix contains a thermo-stable TAQ Polymerase as well as buffer, dNTPs, MgCl<sub>2</sub> and stabilisers at concentrations optimised for the enzyme. The master mix has been freeze-dried to produce a room temperature stable preparation.

The kit includes the lyophilised master mix, resuspension buffer and a tube of ROX dye which can be added as required when the master mix is to be used on hardware platforms that use ROX as a passive reference dye.

The resuspended solution requires only the addition of your cDNA, primers and probe to be PCR ready.

## Kit contents

- 3 x Lyophilised Master Mix (50 reactions per glass ampule)
- 1 x Lyophilised ROX (BROWN)
- 4 x Resuspension buffer (BLUE)

## Kit storage

The Lyophilised 2X qPCR Master Mix should be stored at ambient temperature on arrival. The kit is stable for at least 18 months at ambient temperature. Once resuspended in the provided buffer the kit should be stored at -20°C. Repeated freeze/thawing will not compromise the performance of the product. Under these conditions reagents are stable for six months from date of resuspension.

## Suitable sample material

All kinds of sample material suited for PCR amplification can be used. Please ensure the samples are suitable in terms of purity, concentration and DNA integrity. Always run at least one negative control with the samples. To prepare a negative-control, replace the test sample with RNase/DNase free water.

## Licensing agreement and limitations of use

PCR is covered by several patents owned by Hoffman-Roche Inc and Hoffman-LaRoche, Ltd. Purchase of this kit does not include or provide licence with respect to any patents owned by Hoffman-La Roche or others.

# Resuspension protocol

## 1. For each glass ampule resuspend lyophilised Master Mix in 525µl of resuspension buffer

Do not replace the resuspension buffer with water or any other buffer.

The master mix is then ready to use as a 2X qPCR master mix

## 2. Add ROX if required

ROX is required for platforms that use ROX as a passive reference guide. Use table 1 below to see if ROX addition is required for your hardware platform. If ROX is required then follow the instructions below.

- Resuspend the lyophilised ROX (**BROWN**) in the correct volume of resuspension buffer (**BLUE**) according to table 1 below.
- Add resuspended ROX to each ampule at the correct level.

**Table 1. ROX addition**

Real-time PCR platform	ROX resuspension volume	ROX addition per ampule
Applied Biosystems 7700, 7000, and 7900, 7300 StepOne, StepOnePLUS and ViiA7 platforms, Roche capillary Lightcyclers.	100µl	20µl
All Stratagene platforms	200µl	15µl
Applied Biosystems 7500 platform	700µl	10µl
All Other machines	NOT REQUIRED	NOT REQUIRED

## qPCR detection protocol

- **When using PCRmax qPCR test kits.**

For each 20µl qPCR reaction add the following to each reaction tube

Components	1 Reaction
Lyophilised 2X qPCR Master Mix	10 µl
Primer/probe mix	1 µl
Template (25ng)	5 µl
RNase/DNase free water	4 µl
<b>Final volume</b>	<b>20 µl</b>

- **Suggested use with user supplied primers and probe.**

For each 20µl qPCR reaction add the following to each reaction tube

Components	1 Reaction
Lyophilised 2X qPCR Master Mix	10 µl
Primers (6pmols Forward and Reverse*)	x µl
Probe (3pmols)	x µl
Template (25ng)	x µl
RNase/DNase free water (up to Final volume)	x µl
<b>Final volume</b>	<b>20 µl</b>

\*6pmols of primer gives a working concentration of 300nM in a 20µl reaction

# qPCR Amplification protocol

- For use with PCRmax qPCR test kits.

	Step	Time	Temp
	Enzyme Activation	2 min	95°C
Cycling x50	Denaturation	10 s	95°C
	<b>DATA COLLECTION*</b>	60 s	60°C

\*Fluorogenic data should be collected during this step through the FAM channel.