



Sep 22, 2019

Q5 PCR DNA Amplification (Protocol for Q5® High-Fidelity 2X Master Mix)

Alba Balletbó¹¹Wageningen University

1

Works for me

dx.doi.org/10.17504/protocols.io.7cwhixe



Alba Balletbó



ABSTRACT

This protocol is for PCR with Q5® High-Fidelity 2X Master Mix

GUIDELINES

We recommend assembling all reaction components on ice and quickly transferring the reactions to a thermocycler preheated to the denaturation temperature (98°C). All components should be mixed prior to use.

MATERIALS

NAME	CATALOG #	VENDOR
Q5 High-Fidelity 2X Master Mix - 500 rxns	M0492L	New England Biolabs
Q5 High-Fidelity 2X Master Mix - 100 rxns	M0492S	New England Biolabs

MATERIALS TEXT

Reagent	25 µl Reaction	50 µl Reaction	Final concentration
Q5 High-Fidelity 2X Master Mix	12.5 µl	25 µl	1X
Forward Primer (10 µM)	1.25 µl	2.5 µl	0.5 µM
Reverse Primer (10 µM)	1.25 µl	2.5 µl	0.5 µM
Template DNA	variable	variable	< 1,000 ng
Nuclease-Free Water	to 25 µl	to 50 µl	

Notes: Gently mix the reaction. Collect all liquid to the bottom of the tube by a quick spin if necessary. Overlay the sample with mineral oil if using a PCR machine without a heated lid.

SAFETY WARNINGS

PCR reagents are classified as non-hazardous. Follow the specified handling and disposal considerations included in the safety data sheets provided by the manufacturer.

BEFORE STARTING

Please note that protocols with Q5 High-Fidelity DNA Polymerase may differ from protocols with other polymerases. Conditions recommended below should be used for optimal performance.

- 1 Add all components in a 250 µL tube making up to a 25 or 50 µl reaction. If performing various PCR with different templates, a Master Mix is recommended to be done.

When doing a Master Mix always add Q5 enzyme last, then vortex the solution briefly and centrifugate before use.

'Pro-Tip': Use DMSO 3% or 5X GC enhancer as PCR additives when amplifying particularly difficult or high GC amplicons.

- 2 Gently mix the PCR reactions and transfer the tubes to a thermocycler. Thermocycling conditions for a routine PCR:

Step	Temperature	Time
Initial Denaturation	98°C	30 seconds
25–35 Cycles	98°C	5–10 seconds
*50–72°C	10–30 seconds	
72°C	20–30 seconds/kb	
Final Extension	72°C	2 minutes
Hold	4–10°C	

*The use of the NEB_{Tm} Calculator is highly recommended.



This is an open access protocol distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited