

Sep 20, 2019

 PCR

DOI

[dx.doi.org/10.17504/protocols.io.7hyhj7w](https://dx.doi.org/10.17504/protocols.io.7hyhj7w)

宏亮 董<sup>1</sup>

<sup>1</sup>Northeast Forest University

2019 iGEM NEFU\_China

Tech. support email: [shengyianwork@gmail.com](mailto:shengyianwork@gmail.com)



宏亮 董

OPEN  ACCESS



DOI: [dx.doi.org/10.17504/protocols.io.7hyhj7w](https://dx.doi.org/10.17504/protocols.io.7hyhj7w)

**Protocol Citation:** 宏亮 董 2019. PCR. [protocols.io https://dx.doi.org/10.17504/protocols.io.7hyhj7w](https://dx.doi.org/10.17504/protocols.io.7hyhj7w)

**License:** This is an open access protocol distributed under the terms of the [Creative Commons Attribution License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

**Protocol status:** Working

We use this protocol and it's working

**Created:** September 20, 2019

**Last Modified:** September 20, 2019

**Protocol Integer ID:** 27928

**Keywords:** PCR

## Materials

2×high Taq Master Mix

ddH<sub>2</sub>O

Template

F/R Primer

Bio-rad S1000TM Thermo Cycler.

## Safety warnings

- ! Please wear gloves for the experiment, don't try to touch the lid after PCR program initiation.

## Before start

Synthesize primers in advance.

## Before the start

- 1 Set up a small box with ice, put DNA and 2×high Taq Master Mix into it before going into the Bio-rad S1000TM thermocycler cycler. Set up a small box with ice, put DNA and 2×high Taq Master Mix into it before going into the Bio-rad S1000TM Thermo Cyler.
- 2 Add the following reagent to a PCR tube.(50 µl).

2×high Taq Master Mix (Enzyme)	25 µl
Template	2 µl
Forward Primer (10 µM)	2 µl
Reverse Primer (10 µM)	2 µl
ddH <sub>2</sub> O	19 µl

- 3 Program the thermocycler as follows:

Temper ature	Time
98°C	5 min
98°C	15 s
Tm-4°C	15 s
72°C	(5-10s) × (1 kb)
72°C	5 min
4°C	∞