

2X Pfu Master Mix

Product No : PLMM03
Quantity : 100 reactions



Lot :
Expiry Date :
Supplied with : 4 x 625µl **2X Pfu Master Mix***
3ml of Nuclease-free Water

Store at -20°C
*2X Pfu Master Mix consists of Pfu DNA Polymerase,
(2X ViBuffer Pfu, 0.4mM dNTPs and enhancers

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Description :

2X Pfu Master Mix is an optimized ready-to-use 2X concentrated DNA amplification mixture containing Pfu DNA Polymerase, reaction buffer, dNTPs and enhancers. It contains all the components required for routine DNA amplification except template and primers. Pfu DNA Polymerase is an extremely thermostable proof-reading DNA polymerase. It exhibits the 3' to 5' proofreading activity, resulting in over 10-fold higher fidelity than possible with Taq DNA Polymerases.

Features:

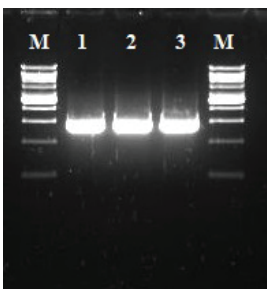
- Saves time and reduces contamination due to reduced number of pipetting steps.
- Recommended for use in high-fidelity amplification, amplification of GC-rich sequences or problematic secondary structures, primer extension reactions at elevated temperatures and cloning of blunt-ended amplification products.
- Stable at 4°C for 6 months, allowing immediate reaction setup without the time-consuming thawing of reagent.
- Suitable for all routine DNA amplification applications.

Storage and Stability:

- Stable at -20°C for 18 months or at 4°C for 6 months if properly stored.
- Stable for 20 freeze-thaw cycles. To avoid frequent freeze-thaw, keeping small aliquot at -20°C is recommended.
- For daily use, keeping an aliquot at 4°C is recommended.

Quality Control:

All preparations are assayed for contaminating endonuclease, exonuclease, and non-specific DNase activities. Functionally tested in DNA amplification.



Amplification of 1.5kb DNA fragment from pTZ DNA using 2X Pfu Master Mix in a 50µl reaction mixture.

- Lane M : VC 1kb DNA Ladder.
Lane 1 : DNA amplification product generated with 1u of Pfu DNA polymerase.
Lane 2 : DNA amplification product generated with 2X Pfu Master Mix (store at -20°C).
Lane 3 : DNA amplification product generated with 2X Pfu Master Mix (after 20 freeze-thaw cycles).

1% TBE agarose gel.

RECOMMENDED PROTOCOL FOR 2X Pfu Master Mix:
Gently mix all solutions after thawing. Spin down briefly and keep on ice. Add the following components in a 0.2ml thin walled PCR tube on ice.
For 50µl reaction volume:

Reagent:	Volume	Final Concentration
2X Pfu Master Mix	25µl	*1X
Primers (Fwd / Rev)	Variable	0.1 - 1 µM each
DNA Template	Variable	0.02 - 5µg
Water, nuclease-free	Adjust final volume to 50µl	

* 1 unit Pfu DNA Polymerase, 1X ViBuffer Pfu, 0.2mM dNTPs and enhancers.
Note : Smaller reaction volume may be achieved provided that the same final concentration of each reaction component is maintained.

CYCLING CONDITIONS (100bp-5kb)	
Denaturation	94°C for 5 minutes
Denaturation	94°C for 30 seconds
Annealing	50 - 68°C for 1 minutes
Extension / 1kb	72°C for 2 minutes
Final Extension	72°C for 7 minutes

} 25 - 35 cycles

This protocol may change depending on the template DNA and primers used.

Product Use Limitation

This product is for research purpose an *in vitro* use only