

#### **Data sheet**

This data sheet provided for information only

# 5X MaSTaqPLUS-2025

#### **RESEARCH USE ONLY**

## Ready-to-use MasterMix for DNA amplification

Cat.No.	Pack	Conc.	
MPL5-100	100 rnx	5 X	
MPL5-500	500 rnx	5 X	

#### **Stability:**

**5X MasTaq**<sup>PLUS</sup> stable for 24 months at -20°C, **or** for 6 months at +4°C storage without freezing.

#### **CONTENT:**

1X: SmarTaq Polymerase
0.2mM of dATP,dCTP,dGTP+
0.6mM dUTP
2.0 mM MgCL<sub>2</sub>
Reaction Buffer Components
Stabilizer/enhancer

#### **DESCRIPTION**

**5X MasTaq**PLUS is a ready-to-use premix of all components for amplification of target DNA, contains **stabilizer/enhancer**, which improves thermostabilization of enzyme during PCR amplification and storage.

**5X MasTaq**<sup>PLUS</sup> contains antibodies-blocked Taq polymerase (**SmarTaq**), which is not active at ambient temperature (during PCR set-up) and activated automatically during the first PCR cycle at the temperature >70°C.

**5X MasTaq**PLUS contains **dUTP** instead of dTTP, what makes this mix very useful for "anticontamination" PCR , allowing to prevent false-positive amplification results, because of possible contaminations by PCR amplicons.

To prevent contaminations **5X MasTaq**<sup>PLUS</sup> should be used in combination with pre-treatment of PCR reaction by UDG (Uracil DNA glycosilase) to destruct of UTP containing amplicons to avoid possible contaminations in the next PCR reaction.

It's no need for prolonged heating for activation of enzyme for PCR.

**5X MasTaq**<sup>PLUS</sup> is optimized for PCR with complex, **low-copy number DNA templates, multiplex PCR, "real-time" PCR** allows to improve specificity of your PCR by titration of MgCl<sub>2</sub> (not included) concentration.

You can use an appropriate volume of **5X MasTaq**<sup>PLUS</sup> for amplification reaction, depending on total final reaction volume.

Just place it into the tube/plate adds primers and template of choice, mix all components and run PCR.

#### **Recommended PCR assay**

50μl PCR assay		Final Conc.
10µl	5X MasTaq <sup>PLUS</sup> -2025	1X
$0.2\text{-}1\mu\text{M}$	each Primer	
Variable*	<b>DNA Template</b>	
To 50μl	PCR Grade Water	

# \*- depending on DNA template initial concentration **APPLICATIONS:**

- -"anti-contamination" PCR
- Routine PCR
- Real-Time PCR
- Low-copy PCR (SmarTaq Polymerase)
- Multiplex PCR

#### **SHIPPING CONDITIONS:**

Should be shipped at ambient Store 5) temperature. For land distance shipments storage). preferably in **Blue Ice** 

#### **STORAGE CONDITIONS:**

Store **5X MasTaq**<sup>PLUS</sup> at  $-20^{\circ}$ C (for long term storage).

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# General Protocol for amplification with 5X MasTaqPLUS-2025

## Add and mix the following components:

Component	50μL reaction	25μL reaction	Final concentration
PCR Grade Water	Up to 50 μL	Up to 25 μL	
5X MasTaq <sup>PLUS</sup> -2025	10 μL	5 μL	1X
Primers			0.2-0.5 μM each
UDG, 1U/μl	1μΙ	0 <b>,</b> 5μΙ	
Template DNA	optionally	optionally	10-50ng

In some cases we recommends to optimize Mg concentration in the range 2.0-3.0mM We recommend to use  $25\mu l$  reaction for the PCR with **5X MasTaq**<sup>PLUS</sup>

**Cycling Protocol:** 

Cycling i rotocon						
Cycle step	3-step a	Cycles				
	T°C	Time				
<b>UDG</b> treatment	47-50°C	2 min	1			
Initial	95°C	95°C 2 min				
Denaturation						
Denaturation	95°C	10-30 S				
Annealing	55-68*	10-30 S 25-3				
Extension	72°C	30-60 S/Kb**				
Final extension	72°C	5-10 min	1			
	4°C	hold				

<sup>\*</sup>Optimal Tm for the primer pair recommended as Tm of the lower primer, for the standard oligos <20nt.

<sup>\*\*</sup>For non-complex DNA templates (plasmid DNA, phage DNA, BAC clone) extension time could be reduced up to 15-30 sec/Kb.

For complex DNA templates (human DNA) strongly recommended to apply Extension time as 60 sec/Kb