

# Apex Hot Start 2.0X Master Mix Blue

with Apex Buffer II

Cat #: 42-149

2.0X Master Mix Kit (1.5mM Final MgCl<sub>2</sub> Conc.)

Unit size: 1000

Contents: 20 x 1.25ml

Storage: -20°C.

Reagent for in vitro laboratory use only

### **General Description**

Apex Hot Start Master Mix BLUE is a ready-to-use 2.0X master mix. Simply add primers, template and water to successfully carry out primer extensions and other molecular biology applications.

Apex Hot Start DNA Polymerase, NH4+ buffer system, dNTPs and magnesium chloride are present in Apex Hot Start Master Mix with Apex Buffer II. Each reaction requires 25  $\mu$ L of the 2.0X reaction mix. Simply add primers, template and water to a total reaction volume of 50  $\mu$ L.

Apex Hot Start DNA Polymerase is a modified form of Apex Taq DNA Polymerase, which is activated by heat treatment. A chemical moiety is attached to the enzyme at the active site, which renders the enzyme inactive at room temperature. Thus, during setup and the first ramp of thermal cycling, the enzyme is not active and misprimed primers are not extended. The result is higher specificity and greater yields when compared to standard DNA polymerases.

Apex Hot Start Master Mix BLUE offers several advantages: Direct gel loading, no need to use separate loading dyes for electrophoresis and subsequent visualization, and the chance of contaminating component stocks is eliminated. Reduction of reagent handling steps leads to better reproducibility. Standard tests can be set up with the confidence that results will be consistent every time.

# Composition of 2.0 X Apex Hot Start Master Mix BLUE w/ buffer II

- Tris-HCl, pH 8.7, Balanced KCl/(NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>,
  3 mM MgCl<sub>2</sub>, 0.2% Tween 20®.
- 0.4 mM dNTPs
- Apex Hot Start DNA Polymerase
- Inert Blue Dye
- Stabilizer

#### **Protocol**

This protocol serves as a guideline for primer extensions. Optimal reaction conditions such as incubation times, temperatures, and amount of template DNA may vary and must be individually determined.

#### **Notes:**

- Set up reaction mixtures in an area separate from that used for DNA preparation or product analysis.
- The table below shows the reaction set up for a final volume of 50  $\mu$ L.
- Important: Mix the solutions completely before use to avoid localized concentrations of salts.
- 1. Set up each reaction as follows:

Component	Vol./Reaction	Final Conc.
Apex Hot Start Master Mix Blue with Apex Buffer II	25 μL	1X
Primer A	Variable	0.1–1.0 μΜ
Primer B	Variable	0.1–1.0 μΜ
PCR Grade Water	Variable	
Template DNA	Variable	Variable
TOTAL volume	50 μL	

- 2. Mix gently by pipetting the solution up and down a few times.
- 3. Program the thermal cycler according to the manufacturer's instructions.
- 4. Each program must start with an initial heat activation step at 95°C for 15 minutes.

For maximum yield and specificity, temperatures and cycling times should be optimized for each new template target or primer pair.



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A typical thermal cycling program is shown below:

<b>95°C</b> for 15 min.		Activate Apex Hot Start Polymerase
30-40 cyc	les:	
95°C	30 sec.	Denature template
45-65°C	30 sec.	Anneal primer
72°C	1-5 min.	Elongation
<b>72°C</b> for 5 min.		Elongation

5. Place the tubes in the thermal cycler and start the reaction.

## **Related Products**

Taq Polymerase kits (500 units)	Cat#	
With 10X Standard and Ammonium Reaction Buffer	42-800B1	
With 10X Combination Buffer	42-800B3	
Glycerol Free	42-800B4	
Hot Start DNA Polymerase (500 units)	Cat#	
With 10X Ammonium and Combination Reaction Buffer	42-106	
High Fidelity - Proof reading (500 units)	Cat#	
Hi-Fi PR™ Taq 2.5 U/μl	42-110	
All polymerases are also available in kits, Mg <sup>2+</sup> free buffers and 50 mM MgCl <sub>2</sub> .		

Master Mixes (500 reactions)	Cat#
2X Taq DNA Polymerase Master Mix, 1.5 mM MgCl <sub>2</sub>	42-132
2X Taq RED Master Mix, 1.5 mM MgCl <sub>2</sub>	42-138
2X Hot Start Master Mix Buffer I, 1.5 mM MgCl <sub>2</sub>	42-198

The shown master mixes are ammonium based. Also available with balanced ammonium and potassium based buffers.

Real-time PCR (400 reactions)	Cat#
qPCR 2X Master Mix for Probe, without ROX <sup>TM</sup>	42-116P
qPCR 2X Master Mix for Probe, low ROX <sup>™</sup>	42-118P
qPCR 2X Master Mix for Probe, high ROX <sup>™</sup>	42-120P
qPCR 2X GREEN Master Mix, without ROX <sup>™</sup>	42-
4PCK 2A GREEN Master Mix, without ROA	116PG
qPCR 2X GREEN Master Mix, low ROX <sup>™</sup>	42-
	118PG
qPCR 2X GREEN Master Mix, high ROX <sup>™</sup>	42-
——————————————————————————————————————	120PG
Ultrapure dNTPs	Cat#
dNTP set, 100 mM each:	42-410
250 μl of each dA, dC, dG and dT	42-410
dNTP Set, 100 mM each:	42-403
1 ml of each dA, dC, dG and dT	72 403
dNTP Mix 40 mM (1 x 500 μl):	42-411
10 mM each dA, dC, dG, dT	72 711
dNTP Mix 100 mM (2 x 1 ml):	42-405
25 mM each dA, dC, dG, dT	72 703
dNTP Mix 10 mM (10 x 1 ml):	42-406
2.5 mM each dA, dC, dG, dT	12 400
Other concentrations and Single dNTPs are available.	
DNA Laddors	Cat#

DNA Ladders	Cat#
Apex 100 bp-Low DNA Ladder, 250 applications	19-109
Apex 1 kb DNA Ladder, 333 applications	19-115
Apex 200 bp DNA Ladder, 200 applications	19-111
Apex ECON Mini DNA Ladder, 100 applications	19-130
Apex ECON Low DNA Ladder, 100 applications	19-131
Apex ECON PCR Ladder, 100 applications	19-132
Accessory reagents	Cat#
50 mM MgCl_ 3 x 1 5 ml	42-303

Nuclease-Free Water, PCR Grade, 6 x 5 ml 42-710

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