



# 2x HotStart PCR Master Mix with Loading Dye

| Article      | Content                     |
|--------------|-----------------------------|
| SL-9712-smp  | 1 ml, 100 rxn × 20 µl       |
| SL-9712-5ML  | 5 x 1 ml, 500 rxn × 20 µl   |
| SL-9712-10ML | 10 x 1 ml, 1000 rxn × 20 µl |
| SL-9712-20ML | 20 x 1 ml, 2000 rxn × 20 µl |



Long-Term Storage at -20°C in the dark

Short-Term Storage at 4°C in the dark

#### DESCRIPTION

Our **primaAMP 2x HotStart PCR Master Mix** includes a recombinant, thermostable Taq from Thermus aquaticus. It possesses a 5' to 3' polymerase activity as well as a 5'-flap endonuclease activity.

The **primaAMP** Taq DNA polymerase can be used to amplify DNA fragments up to a length of 5 kb. Moreover, it generates A (adenine) overhangs at the 3' end, which can be used for TA-cloning.

The sophisticated buffer system contains potassium chloride as well as ammonium sulfate and allows the amplification of difficult templates (e.g GC-rich).

A **red-colored loading dye is included** in the Master Mix which allows for direct loading onto agarose gels after PCR.



#### **DID YOU KNOW?**

- primaAMP is also available as a stand-alone polymerase.
- For a Master Mix without HotStart polymerase, please order SL-9612.



## 🛞 primaAMP

# **Recommended Reaction Mixture per Well**

## 

## **BEFORE YOU START**

- After thawing, please invert the Master Mix tube 6-8 times.
- **Do not vortex** the Master Mix to avoid damaging of the enzyme.

| Component                     | 20 µl<br>Reaction       | 10 μl<br>Reaction       | Final<br>Concentration       |
|-------------------------------|-------------------------|-------------------------|------------------------------|
| 2x primaAMP<br>PCR Master Mix | 10 µl                   | 5 µl                    | 1x                           |
| Forward Primer                | variable<br>(e.g. 2 µl) | variable<br>(e.g. 1 µl) | 100 - 400 nM                 |
| Reverse Primer                | variable<br>(e.g. 2 µl) | variable<br>(e.g. 1 µl) | 100 - 400 nM                 |
| Template DNA                  | variable                | variable                | 0.01 - 10 ng<br>per reaction |
| Sterile Water                 | adjust to 20 µl         | adjust to 10 µl         |                              |





# **Suggested Cycling Conditions**

| Step                 | Time                                       | Temperature                             |  |  |
|----------------------|--|---|--|--|
| Initial Denaturation | 3 minutes                                  | 92°C - 95°C                             |  |  |
| 25 - 35 cycles       |  |   |  |  |
| Denaturation         | 5 - 10 seconds                             | 92°C - 95°C                             |  |  |
| Annealing            | 5 - 10 seconds                             | 55°C - 68°C<br><b>depends on primer</b> |  |  |
| Extension            | 5 - 30 seconds per 1 kb<br>amplicon length | 72°C                                    |  |  |



### NOTE

- The optimal annealing temperature is usually 2°C 5°C below the primer melting temperature.
- Recommended elongation time is 5 30 seconds per 1 kb of amplicon length. For more complicated templates, we suggest 45 seconds for elongation.
- For maximum yield and specificity, we recommend to optimize annealing temperatures, annealing time, extension time, and the number of cycles should be optimized and the number of cycles for each template and primer pair.







## **Further Information**

For more information, please visit our website: www.steinbrenner.de



In der Au 17 | 69257 Wiesenbach

J +49 (0) 6223 / 96 73 00
Mail@steinbrenner.de

## **Further Products**

Products that may also be interesting to you

😢 primaDIRECT 🔤

- Direct-PCR without DNA extraction
- From sample to PCR in 15 minutes
- For cell culture, tissue, plants, mouse tails/ear, meat

S primaPROOF HIGH-FIDELITY

- High-fidelity / proofreading PCR
- For NGS and cloning