

# 10X STANDARD BUFFER 1,5MM MGCL<sub>2</sub>

#### Cat. No.: 257757

-	10x Standard Buffer, 1,5mM MgCl₂
ID No.	CL1.500-0009
Cap color	Clear
Content	3 x 1.5 ml

#### Features and General Description

10x Standard Buffer are usually supplied in 10x formulations with 15 mM MgCl $_2$  included but are also available as Mg $^{2+}$  free buffer, detergent free buffer as well as Mg $^{2+}$  and detergent free buffer.

#### **Standard Buffer**

Standard Buffer is the traditional potassium ( $K^+$ ) buffer. Standard Buffer promotes high specificity and careful optimization of primer annealing temperatures and  $Mg^{2+}$  concentrations may be required.

#### Magnesium

 ${\rm Mg}^{2+}$  is required for polymerase activity. Low  ${\rm Mg}^{2+}$  concentrations increase the fidelity but with too low  ${\rm Mg}^{2+}$  concentrations the polymerase will not work. The  ${\rm Mg}^{2+}$  concentration available in the reaction is dependent on several parameters e.g. the presence of chelators or the dNTP concentration. Therefore, the  ${\rm Mg}^{2+}$  concentration should be optimized.

#### Tween

Non-ionic detergents are used to prevent the polymerase to stick to the walls of the tube, to stabilize the polymerase and increase yield. However, these agents might increase non-specific amplification or interfere with downstream reactions. Tween can be used to neutralize SDS contaminations in the DNA template.

#### **Recommended Storage and Stability**

Long term storage at -20  $^{\circ}\text{C}.$  Product expiry at -20  $^{\circ}\text{C}$  is stated on the label.

Option: Store at +4 °C for up to 6 months.

#### **Quality Control**

Each lot of buffer is functionally tested in PCR.

#### Kit Components

### 10x Standard Buffer

Tris-HCl pH 8.5, KCl, 15 mM MgCl<sub>2</sub>, 1% Tween<sup>®</sup> 20.

## Determining the optimal buffer system for your application

ClearLine offers several PCR buffers to allow the customer to choose the optimal buffer system for a specific amplification process.

For your specific application the optimal reaction condition can be determined by comparing PCR reactions containing the different buffers.

The final concentration of the buffer in the reaction should be 1x.

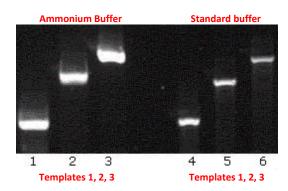


Figure 1: Amplification of three different cDNA templates using Ammonium Buffer versus Standard Buffer.

Other product sizes, combinations and customized solutions are available. Please look at www.dutscher.com or ask for our complete product list for PCR Enzymes. For customized solutions please contact us.

Made in Europe

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