

SuperHot Taq DNA Polymerase

Taq DNA Polymerase for HotStart PCR. Product No. A5231

Description

SuperHot Taq DNA Polymerase is the optimized mixture of Taq DNA Polymerase and anti-Taq DNA polymerase monoclonal antibodies. Antibodies block polymerase activity during the set-up of the PCR reactions at ambient temperature (20 - 22°C). The inhibition of Taq DNA polymerase is completely reversed when the temperature is above 70°C. The PCR products obtained with SuperHot Taq are free from unspecific products and from primer-dimers.

Unit definition: One unit is defined as the amount of enzyme required to incorporate 10 nmoles of dNTP into an acid-insoluble DNA fraction in 30 minutes at 72°C.

Supplied in Storage buffer: 10 mM Tris · HCl (pH 7.0), 50 mM KCl, 0.1 mM EDTA, and 50 % glycerol.

Supplied with 3 Reaction buffers (10X):

1 Tube	'incomplete'	160 mM (NH ₄) ₂ SO ₄ , 670 mM Tris · HCl (pH 8.8), 0.1 % Tween [®] 20.
1 Tube	'complete'	160 mM (NH ₄) ₂ SO ₄ , 670 mM Tris \cdot HCl (pH 8.8), 0.1 % Tween [®] 20, 25 mM MgCl ₂
1 Tube	'complete II KCl'	500 mM KCl, 100 mM Tris \cdot HCl (pH 8.8), 0.1 % Tween [®] 20, 15 mM MgCl ₂
1 Tube	$MgCl_2$ (100 mM)	

Recommended Reaction buffer (1X):	16 mM (NH ₄) ₂ SO ₄ , 67 mM Tris \cdot HCl (pH 8.8); 1.5 - 7 mM MgCl ₂ , 0.01% Tween $^{\circledast}$ 20
Recommended PCR conditions:	Use PCR conditions optimized for <i>Taq</i> DNA polymerase. In the case of low amount of DNA template, additional cycles may be used.
Applications:	Complex genomic or cDNA templates, low copy number targets, large numbers of thermal cycles, multiplex PCR
Storage Conditions:	-20°C
Concentration:	5000 units/ml

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