

Datasheet for ABIN5519492

## TransFast® Taq DNA Polymerase (with 2.5 mM dNTPs)

### Overview

Quantity:	500 units
Application:	Polymerase Chain Reaction (PCR)

### Product Details

Purpose:	TransFast® Taq DNA Polymerase is an engineered version of Taq DNA Polymerase.
Brand:	TransFast®
Specificity:	The enzyme consists of a single polypeptide with a molecular weight of approximately 94 kDa. TransFast® Taq DNA Polymerase has 5'-3' DNA polymerase activity and 5'-3' exonuclease activity. It lacks 3'-5' exonuclease activity.
Characteristics:	<ul style="list-style-type: none"> <li>- Extension rate is about 6 kb/min.</li> <li>- Template-independent "A" can be generated at the 3' end of the PCR product. PCR products can be directly cloned into pEASY®-T vectors.</li> <li>- Amplification of genomic DNA fragment up to 4 kb.</li> </ul>
Components:	DNA Polymerase, 10X Taq Buffer, 2.5 mM dNTPs, 6X DNA Loading Buffer
Unit Definition:	One unit of TransFast® Taq DNA Polymerase incorporates 10 nmol of deoxyribonucleotide into acid-precipitable material in 30 minutes at 74°C.

### Application Details

Application Notes:	Routine PCR, High throughput PCR, Colony PCR
Comment:	TransFast® Taq DNA Polymerase has passed the following quality control assays: functional absence of double- and single-strand endonuclease activity, >99% homogeneous measured by SDS-PAGE. Each batch of TransFast® Taq DNA Polymerase has been assayed for amplification efficiency to amplify p53 gene from 10 ng of human genomic DNA.
Restrictions:	For Research Use only

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## Handling

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Buffer:	Storage Buffer: 20 mM Tris-HCl ( pH 8.0), 0.1 mM EDTA, 1 mM DTT, 100 mM KCl, 50 % glycerol, stabilizers. 10xTransFast® Taq Buffer (with Mg <sup>2+</sup> ): 200 mM Tris-HCl ( pH 8.4), 100 mM KCl, 100 mM (NH <sub>4</sub> ) <sub>2</sub> SO <sub>4</sub> , 20 mM MgSO <sub>4</sub> , others
Storage:	-20 °C
Storage Comment:	at -20°C for two years
Expiry Date:	24 months

## Publications

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Product cited in:	Johnson, Drugan, Miller, Evans: "38" in: , Vol. 1363, Issue Nucleic acids research, pp. 28-39, (1991)
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