

Datasheet for ABIN5519583

TransTaq®-T DNA Polymerase (with 2.5 mM dNTPs)

Overview

Quantity:	250 units
Application:	Polymerase Chain Reaction (PCR)

Product Details

Purpose:	TransTaq®-T DNA Polymerase is a mixture of EasyTaq® DNA Polymerase with a proofreading 3'-5' exonuclease.
Brand:	TransTaq®
Specificity:	TransTaq®-T DNA Polymerase is a mixture of EasyTaq® DNA Polymerase with a proofreading 3'-5' exonuclease. The fidelity is equal to EasyPfu DNA Polymerase. The yield is equal to that from EasyTaq® DNA Polymerase. It is more suitable for high fidelity TA cloning.
Characteristics:	<ul style="list-style-type: none"> - TransTaq®-T DNA Polymerase offers 18-fold fidelity as compared to EasyTaq® DNA Polymerase. - Extension rate is about 1-2 kb/min. - Template-independent "A" can be generated at the 3' end of the PCR product. PCR products can be directly cloned into pEASY®-T vectors. - Amplification of genomic DNA fragment up to 8 kb.
Components:	DNA Polymerase, 10X Taq Buffer, 2.5 mM dNTPs, 6X DNA Loading Buffer
Unit Definition:	One unit of TransTaq®-T DNA Polymerase incorporates 10 nmol of deoxyribonucleotide into acid-precipitable material in 30 minutes at 74°C.

Application Details

Application Notes:	Complex templates, TA Cloning
Comment:	TransTaq®-T 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 TransTaq®-T DNA Polymerase has been assayed for amplification

Order at www.genomics-online.com

USA & Canada: +1 877 302 8632 | support@antibodies-online.com

Application Details

efficiency to amplify p53 gene from 10 ng of human genomic DNA.

Restrictions: For Research Use only

Handling

Buffer: Storage Buffer: 20 mM Tris-HCl (pH 8.0), 0.1 mM EDTA, 1 mM DTT, 100 mM KCl, 50 % glycerol, stabilizers
10xTransTaq@-T Buffer: 200 mM Tris-HCl (pH 9.0), 100 mM KCl, 100 mM (NH₄)₂SO₄, 20 mM MgSO₄, others

Storage: -20 °C

Storage Comment: at -20°C for two years

Expiry Date: 24 months

Publications

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