

Lot 0006247171

Page 1 of 2

Catalog Number 600684

Product Name Paq5000 DNA Polymerase

Expiration Date 2016-08-30

Quantity 5000 U

Certified By Dana Faver

Quality Controlled By Tricia Molina

Shipping Conditions Shipped on dry ice.

Storage Conditions Store at -20°C upon receipt.

Concentration

5 U/µ1

Description

Paq5000 DNA Polymerase* is a new alternative to *Taq* DNA polymerase that provides higher PCR yields. In addition, Paq5000 DNA Polymerase is optimized for fast cycling conditions to save time and increase throughput. The enzyme is a recombinant DNA-dependent DNA polymerase isolated from a Pyrococcus species. Paq5000 DNA Polymerase is ideal for endpoint PCR for up to 6 kb genomic targets. It is not recommended for high fidelity cloning or 5' nuclease assays.

Agilent Technologies

10x Reaction Buffer

Use only the provided 10^{\times} Paq5000 Reaction Buffer. Changing the reaction buffer may adversely affect enzyme performance. The 10^{\times} buffer provides a final 1^{\times} Mg²⁺ concentration of 2 mM.

Unit Activity

One unit of activity is the amount of enzyme required to incorporate 10 nanomoles of [3H]dTTP in 30 minutes at 72°C.

Test Conditions

Paq5000 DNA polymerase was tested by polymerase chain reaction (PCR) assays using multiple primer-template systems, with targets sizes ranging from 0.9–6.0 kb. The resulting amplification products were visualized on an agarose gel to verify the production of discrete products of the expected size.

Contamination Test Conditions

Paq5000 DNA polymerase is tested to confirm the absence of detectable nonspecific nuclease and DNA contamination.

PCR Protocol

To a nuclease-free microcentrifuge tube, add the following components in order:

x μl of dH_2O for a final reaction volume of 50 μl

5 μl of 10× Paq5000 Reaction Buffer

 $0.4~\mu l$ of 100~mM dNTP mix (25 mM each dNTP)

1 μl of 10 μM forward primer

 $1~\mu l$ of $10~\mu M$ reverse primer

100 ng genomic DNA or 5-30 ng vector DNA

 $0.5~\mu l$ of Paq5000 DNA Polymerase (5 U/ μl)

Mix the reaction gently, then place the reaction in a thermocycler and run the following PCR amplification program:

Cycles	95°C	Duration 2 minutes
30	95°C Primer T _m –5°C 72°C	20 seconds 20 seconds 30 seconds/kb
1	72°C	5 minutes

Endnotes

This product is provided under an agreement between Bio-Rad Laboratories and Agilent Technologies, and the manufacture, use, sale or import of this product is subject to EP Pat. No. 1 283 875 B1, owned by Bio-Rad Laboratories, Inc. Purchase of this product conveys to the buyer the non-transferable right to use the purchased amount of the product and components of the product in PCR (but not real-time PCR) in the Research Field including all Applied Research Fields (including but not limited to forensics, animal testing, and food testing).

Purchase of this product conveys to the purchaser the non-transferable right under these patents to use the product for research use only.

*U.S. Patent Nos. 6,734,293; 6,444,428; 6,183,997; 5,489,523.

Limited Product Warranty

This warranty limits our liability to replacement of this product. No other warranties of any kind, express or implied, including, without limitation, implied warranties of merchantability or fitness for a particular purpose, are provided by Agilent. Agilent shall have no liability for any direct, indirect, consequential, or incidental damages arising out of the use, the results of use, or the inability to use this product.

For research use only. Not for use in diagnostic procedures. This certificate is a declaration of analysis at the time of manufacture.