Certificate of Analysis



Advantage® Genomic LA Polymerase Mix

Catalog No. Amount Lot Number

639152 100 rxns Specified on product label.

Description

The Advantage Genomic LA Polymerase Mix is composed of a full-length Taq DNA polymerase, a small amount of proofreading enzyme, and a hot-start antibody that inhibits polymerase activity at ambient temperature. The Advantage Genomic LA Polymerase Mix allows synthesis of products up to 30 kb for complex templates such as genomic DNA, and up to 48 kb for less complex templates such as λ DNA. This polymerase also has a 6.5X higher fidelity than wild-type Taq DNA polymerase due to the presence of the proofreading 3'-to-5' exonuclease. The robust enzyme/buffer system requires less optimization and produces higher yields than other "long and accurate polymerases." Sufficient polymerase mix and buffer are provided for 100 reactions (25 μ l each).

Package Contents

- 25 μl Advantage Genomic LA Polymerase Mix (5 units/μl)
- 500 μl 10X Advantage Genomic LA Buffer (containing 25 mM MgCl₂)

Storage Conditions

• Store at -20°C.

Expiration Date

• Specified on product label.

Shipping Conditions

Dry ice

Product Documents

Documents for our products are available for download at <u>takarabio.com/manuals</u> The following documents apply to this product:

Advantage Genomic LA Polymerase Mix Protocol-At-A-Glance

Quality Control Data

Raw Material Quality Control

Purity

No nicking activity, endonuclease activity, or exonuclease activity was detected after incubation of 0.6 mg of supercoiled pBR322 DNA, 0.65 mg of λ DNA, or 0.6 mg of λ -Hind III digest with 10 units of this enzyme for 1 hr at 74°C.

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Unit definition

One unit is the amount of enzyme that will incorporate 10 nmol of dNTP into acid-insoluble products in 30 min at 74°C using activated salmon sperm DNA as template.

Reaction mixture for unit definition

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25 mM TAPS (pH 9.3 at 25°C)
50 mM KCl
2 mM MgCl<sub>2</sub>
1 mM 2-mercaptoethanol
200 mM Each dATP, dGTP, dTTP
100 mM [α-<sup>32</sup>P]-dCTP
0.25 mg/ml Activated salmon sperm DNA
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Functional Quality Control

PCR performance

The Advantage Genomic LA Polymerase Mix was tested for PCR performance using a λ DNA control template and primers. Reactions (50 μ l) were assembled and performed using a standard PCR protocol. When the PCR products were examined by electrophoresis on a 1.0% agarose/EtBr gel, a major band of 35 kb was observed.

To ensure optimum PCR performance, the Advantage Genomic LA Polymerase Mix was also tested using human genomic DNA as template. Reactions were assembled and performed using a standard PCR protocol. When the PCR products were examined by electrophoresis on a 1.0% agarose/EtBr gel, a major band of 17.5 kb was observed.

Hot-start antibody

Hot-start antibody inhibition of the Advantage Genomic LA enzyme was confirmed to be greater than 80% following a 10 min incubation of the reaction at 55°C.

It is certified that this product meets the above specifications, as reviewed and approved by the Quality Department.

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NOTICE TO PURCHASER:

Our products are to be used for **Research Use Only**. They may not be used for any other purpose, including, but not limited to, use in humans, therapeutic or diagnostic use, or commercial use of any kind. Our products may not be transferred to third parties, resold, modified for resale, or used to manufacture commercial products or to provide a service to third parties without our prior written approval.

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Takara Bio USA, Inc.

2560 Orchard Parkway, San Jose, CA 95131, USA U.S. Technical Support: technical_support@takarabio.com

 United States/Canada
 Asia Pacific
 Europe
 Japan

 800.662.2566
 +1.650.919.7300
 +33.(0)1.3904.6880
 +81.(0)77.565.6999

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