

Titanium® Taq PCR Kit

Catalog Nos.	Amount	Lot Number
639210	100 rxn	Specified on product label.
639211	30 rxn	Specified on product label.

Description

Titanium *Taq* is a mixture consisting of a 5' exonuclease-deficient *Taq* polymerase, and TaqStart® Antibody, a monoclonal antibody which inhibits *Taq* at ambient temperatures. TaqStart Antibody provides automatic hot start PCR. Enough enzyme and buffer are supplied for 30 PCR reactions of 50 μl each. An aliquot of calf thymus DNA is provided as a control template for amplifying a 407-bp fragment of the BPTI gene using the Control Primer Mix.

Package Contents

<u>639210</u>	<u>639211</u>	
(100 rxns)	(30 rxns)	
100 μ1	30 µl	50X Titanium Taq DNA Polymerase
600 µl	200 μ1	10X Titanium Taq PCR Buffer
120 μ1	40 μ1	50X dNTP Mix (10 mM each)
100 μ1	30 µl	Control DNA Template (100 ng/µl)
100 μ1	30 μ1	Control Primer Mix (10 µM each)
4 x 1.25 ml	2 x 1.25 ml	PCR-Grade Water

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:

- Titanium Taq PCR Kits User Manual
- Titanium Taq PCR Kit Protocol-At-A-Glance

Titanium Taq PCR Kit

Quality Control Data

Raw Material Quality Control

Purified Titanium *Taq* DNA Polymerase was tested for enzymatic activity and PCR performance. Endonuclease, exonuclease, and DNA contamination assays were also performed.

PCR Performance

Titanium Taq was serially diluted and each serial dilution used in a separate PCR reaction with λ genomic DNA as a template. Optimal dilution per reaction was determined as the amount of enzyme required to amplify >20 ng/µl of a 3.5 kb λ fragment with minimal background.

Functional Quality Control

Amplification from a cDNA template

Titanium Taq was tested in a 50 μ l PCR reaction using 5 μ l of Marathon®-Ready Human Placenta cDNA (Cat. No. 639311) as a template and primers specific for a 1.3 kb fragment from the transferrin receptor gene (0.2 μ M each). Conditions were set at:

5 μ l of the PCR product was run on a 1.2% TAE/agarose gel to confirm the presence of a 1.3 kb band with minimal background. PCR product concentration was measured by fluorometry. The yield was determined to be >5 ng/ μ l.

Amplification from a genomic DNA template

Titanium Taq was tested in a 50 μ l PCR reaction using 100 ng of calf thymus genomic DNA as a template and primers specific for a 407 bp fragment of the BPTI gene (0.4 μ M each). Conditions were set at:

5 μ l of PCR product was electrophoresed on a 1.2% TAE/agarose gel to confirm the presence of a 407 bp band with minimal background. PCR product concentration was measured by fluorometry. The yield was determined to be \geq 15 ng/ μ l.

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639210 & 639211

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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