

## Titanium® DNA Amplification Kit

<b>Catalog No.</b> 639240	<b>Amount</b> 300 rxns	<b>Lot Number</b> Specified on product label.
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### Description

Titanium *Taq* is a mixture consisting of a 5' exonuclease-deficient *Taq* polymerase and TaqStart® Antibody, a monoclonal antibody which inhibits Titanium *Taq* at ambient temperatures. TaqStart Antibody provides automatic hot start PCR. An optimized buffer mix (yielding a final concentration of 3.5 mM MgCl<sub>2</sub>) and a purified mixture of dNTPs (2.5 mM each) is also included. The RNase-free GC-Melt Reagent (5M) improves the specificity and yield of PCR reactions, especially when using templates with a high GC-content or complex secondary structure. This kit contains sufficient reagents for Titanium reactions of 100 µl each. The Titanium DNA Amplification Kit is designed to be used with Affymetrix DNA Mapping products (see Table 1).

### Package Contents

- 0.67 ml 50X Titanium *Taq* DNA Polymerase Mix
- 3.4 ml 10X Titanium *Taq* PCR Buffer
- 4.7 ml dNTP Mixture (2.5 mM each)
- 6.7 ml GC-Melt Reagent (5.0 M)

### Storage Conditions

- Store at -20°C

**NOTE:** At times, precipitate may be observed in the GC-Melt. This precipitate does not affect the performance of the kit. The precipitate can be dissolved rapidly by mixing at room temperature or warming at 37°C for a few minutes.

### Expiration Date

- Specified on product label.

### Shipping Conditions

- Dry ice

### Quality Control Data

#### Raw Material Quality Control

Purified N-terminal deletion mutant *Taq* polymerase was tested for enzymatic activity and PCR performance. Endonuclease, exonuclease, and DNA contamination assays were also performed.

#### PCR Performance

N-terminal deletion mutant *Taq* polymerase was serially diluted (0.6–0.2 µg/µl) and each serial dilution used in a separate PCR reaction with λ genomic DNA as a template. The optimal protein concentration per reaction was determined as the amount of protein (in µg) required to amplify >20 ng/µl of a 3.5 kb λ fragment with minimal background.

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## Functional Quality Control

### Amplification from a cDNA template

Titanium *Taq* DNA polymerase and 10X Titanium Buffer were tested in a 50  $\mu$ l PCR reaction using 5  $\mu$ l of Marathon®-Ready Human Placenta cDNA (Cat. No. 639311) and control primers specific for a 1.3 kb fragment from the transferrin receptor (TFRC) gene (0.2  $\mu$ M each). Conditions were set at:

95°C	1 min
30 cycles:	
98°C	15 sec
68°C	1.5 min

5  $\mu$ l of the PCR product was electrophoresed 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  $\geq 5$  ng/ $\mu$ l.

### Amplification from a genomic DNA template

Titanium *Taq* DNA polymerase and 10X Titanium Buffer were tested in a 50  $\mu$ l PCR reaction using 100 ng of calf thymus genomic DNA as a template and control primers specific for a 407 bp fragment of the bovine pancreatic trypsin inhibitor (BPTI) gene (0.4  $\mu$ M each). Conditions were set at:

94°C	3 min
30 cycles:	
94°C	30 sec
68°C	1.5 min

5  $\mu$ 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/ $\mu$ l.

### Amplification from a GC-rich template

The GC-Melt Reagent (5M) was tested using the reagents from the Advantage® GC 2 PCR Kit (Cat. Nos. 639120 & 639119). Amplification of a 510 bp GC-rich fragment from the insulin-like growth factor receptor II (IGFR II) gene, of which 110 bp are 90% GC-rich, was performed using the kit reagents in the presence of increasing concentrations of GC-Melt Reagent (0, 0.5, 1.0, 1.5, and 2 M). Reactions were assembled and performed as described in the Advantage GC 2 User Manual, PT3316-1. Cycle parameters were set as follows:

94°C	3 min
25 cycles:	
94°C	30 sec
68°C	1.5 min
68°C	3 min

5  $\mu$ l of each PCR product was electrophoresed on a 2.0% agarose/EtBr gel. The presence of a major band of 0.5 kb was observed in the lanes containing template amplified in the presence of 0.5 M and 1.0 M GC-Melt Reagent.

# Certificate of Analysis

Cat. No. 639240

Titanium DNA Amplification Kit

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## dNTP Mixture (2.5 mM each)

The dNTP Mixture (2.5 mM each) was tested in a PCR reaction using  $\lambda$  DNA as a template. Amplification of both 500 bp and 20 kb fragments was observed. In addition, the dNTP mixture was tested in an RT-PCR reaction using total RNA from HL60 cells as a template. Amplification of a 4.4 kb fragment was observed using a human TFRC gene cDNA template.

**Table 1. Affymetrix and Takara Bio Kit Cat. Nos.**

Product	Affymetrix Cat. Nos.			Takara Bio Cat. Nos.	
	30 Rxn Kits	100 Rxn Kits	User Manual	639240 300 Rxn Kit	639243 400 Rxn Kit
GeneChip Mapping 250K Nsp Assay Kit	900766	900753	701930	✓	—
GeneChip Mapping 250K Sty Assay Kit	900765	900754	701930	✓	—
Affymetrix Genome-Wide Human SNP Nsp/Sty Assay Kit 5.0	901013	901015	702419	✓	✓

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

## Titanium® DNA Amplification Kit

### CATALOG NO.

639240

### 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.

Your use of this product is also subject to compliance with the licensing requirements, listed below if applicable, and described on the product's web page at <http://www.takarabio.com>. It is your responsibility to review, understand and adhere to any restrictions imposed by these statements.

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