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# PRODUCT: Tet-Off® Advanced Vector Set

**CATALOG No.** 631126 (Not sold separately)

## **LOT NUMBER**

Specified on product label

# STORAGE CONDITIONS

- Store plasmids at -20°C.
- Spin briefly to recover contents.
- Avoid repeated freeze/thaw cycles.

# **PLASMID STORAGE BUFFER**

10 mM Tris-HCI (pH 8.0) 1 mM EDTA (pH 8.0)

#### **SHELF LIFE**

1 year from date of receipt under proper storage conditions

## **SHIPPING CONDITIONS**

Dry ice (-70°C)

## **DESCRIPTION**

The Tet-Off Advanced Vector Set is used to create powerful, tightly regulated, tetracycline-inducible mammalian expression systems that are controlled by the withdrawal of doxycline from the culture medium.

## PACKAGE CONTENTS

- 20 µl pTet-Off-Advanced Vector  $(500 \text{ ng/}\mu\text{I})$
- 20 μl pTRE-Tight Vector (500 ng/μl)
- 20 µl pTRE-Tight-Luc Vector (500 ng/µl)
- 40 µl Linear Hygromycin Marker (50 ng/µI)

#### **OTHER**

- Tet-Off Advanced Inducible Gene **Expression System User Manual** (PT3945-1)
- pTet-Off-Advanced Vector Information Packet (PT3945-5)
- pTRE-Tight Vector Information Packet (PT3720-5)

## FOR RESEARCH USE ONLY

#### **QUALITY CONTROL DATA**

See back of page.



United States/Canada 800.662.2566

Asia Pacific +1.650.919.7300

Europe +33.(0)1.3904.6880

Japan +81.(0)77.543.6116

Clontech Laboratories, Inc. A Takara Bio Company 1290 Terra Bella Ave. Mountain View, CA 94043 Technical Support (US) E-mail: tech@clontech.com www.clontech.com

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(PA9X3421)

# **QUALITY CONTROL DATA**

# **Identity** and purity

• The identity of each of the following was verified by restriction digestion. Digestion with the indicated restriction enzymes produced fragments of the indicated sizes on a 0.8% agarose/EtBr gel:

Plasmid Name	Enzyme	<u>Fragments</u>
pTet-Off-Advanced	EcoRI BamHI	7.1 kb 2.6 & 4.5 kb
pTRE-Tight	PvuI XhoI	2.6 kb 0.6 & 2.0 kb
pTRE-Tight-Luc	XbaI BamHI & NheI	4.2 kb 1.6 & 2.6 kb
Linear Hygromycin Marker	HindIII & XbaI	0.5, 0.6 & 1.1 kb

A<sub>260</sub>/A<sub>280</sub>: 1.8–2.0

# **Functional Test**

• Linear Hygromycin Marker

HEK 293 cells were transfected with 200 ng Linear Hygromycin Marker using a liposome-mediated transfection method. After 5 hr at 37°C, the transfection solution was removed and the cells were given fresh medium. 48 hr later, the cells were plated in two 10 cm plates. 48 hr after plating, medium containing hygromycin was added to the plates. After 2–3 weeks, >20 clones were identified.

# **Notice to Purchaser**



# **Tet-Off® Advanced Vector Set**

CATALOG NO.

631126

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#### **STATEMENT 42**

Use of the Tetracycline controllable expression systems (the "Tet Technology") is covered by a series of patents including U.S. Patent Nos. 6087166, 6271341, 7541446, 8383364, European Patents: EP 0990030, 1954811, 2050818, 2352833 and corresponding patent claims outside these regions which are proprietary to TET Systems GmbH & Co. KG. Academic research institutions are granted an automatic license with the purchase of this product to use the Tet Technology only for internal, academic research purposes, which license specifically excludes the right to sell, or otherwise transfer, the Tet Technology or its component parts to third parties. Notwithstanding the above, academic and not-for profit research institutions whose research using the Tet Technology is sponsored by for profit organizations, which shall receive ownership to all data and results stemming from the sponsored research, shall need a commercial license agreement from TET Systems in order to use the Tet Technology. In accepting this license, all users acknowledge that the Tet Technology is experimental in nature. TET Systems GmbH & Co. KG makes no warranties, express or implied or of any kind, and hereby disclaims any warranties, representations, or guarantees of any kind as to the Tet Technology, patents, or products. All others are invited to request a license from TET Systems GmbH & Co. KG prior to purchasing these reagents or using them for any purpose. Clontech is required by its licensing agreement to submit a report of all purchasers of the Tet-controllable expression system to TET Systems. For license information, please contact: GSF/CEO TET Systems GmbH & Co. KG, Im Neuenheimer Feld 582 69120 Heidelberg Germany Tel: +49 6221 5880400 Fax: +49 6221 5880404 email: info@tetsystems.com or use the electronic licensing request form via www.tetsystems.com/main inquiry.htm

# TRADEMARKS:

Clontech Laboratories, Inc.

A Takara Bio Company 1290 Terra Bella Avenue, Mountain View, CA 94043, USA U.S. Technical Support: tech@clontech.com

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United States/Canada Asia Pacific Europe Japan

800.662.2566 +1.650.919.7300 +33.(0)1.3904.6880 +81.(0)77.543.6116

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A Takara Bio Company 1290 Terra Bella Avenue, Mountain View, CA 94043, USA U.S. Technical Support: tech@clontech.com

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