

Luminescent β -galactosidase Reporter System 3

Catalog No(s).	Amount	Lot Number
631713	each	Specified on product label.

Description

The Luminescent β -galactosidase Reporter System 3 is a complete system for the analysis of transcriptional regulation using the reporter enzyme β -galactosidase. The kit contains two pLacZ reporter vectors and reagents necessary for 100 chemiluminescent assays to detect enzyme activity. The protocol describes chemiluminescent detection of β -galactosidase via a scintillation counter, tube or plate luminometer, or x-ray film.

Package Contents

Box 1:

- 10 μ g pLacZ-Basic Vector (500 ng/ μ l)
- 10 μ g pLacZ-Control Vector (500 ng/ μ l)
- 30 μ l Positive Control β -galactosidase

Box 2:

- 25 ml Reaction Buffer
- 500 ml Reaction Substrate

Storage Conditions

- Store pLacZ Vectors and Positive Control β -galactosidase at -20°C
- Store all other components at 4°C . **DO NOT FREEZE.**

Shelf Life

- 6 months from date of receipt under proper storage conditions for Positive Control β -galactosidase, Reaction Buffer, and Reaction Substrate.
- 1 year from date of receipt under proper storage conditions for all other components.

Shipping Conditions

- Blue ice (4°C)

Product Documents

Documents for Clontech® products are available for download at www.clontech.com/manuals

The following documents apply to this product:

- Luminescent Beta-Gal User Manual (PT2106-1)
- Luminescent Beta-Gal Protocol-At-A-Glance (PT2106-2)

Clontech Laboratories, Inc.

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Quality Control Data

- Each pLacZ reporter vector was tested for purity and identity by restriction enzyme digestion:

	Enzyme(s)	Fragment(s)*
pLacZ-Basic Vector	Hae II	1.6, 1.1, & 0.9 kb
pLacZ-Control Vector	Hae II	1.8, 1.6, 1.2 & 0.9 kb

*Additional fragments <0.9 kb could not be observed for either vector.

- Plasmid purity was also verified by A_{260}/A_{280} .
- Enzyme assay: 1 μ l (1 U) of the Positive Control β -galactosidase was added to 19 μ l of a cell lysate prepared from untransfected CHO cells according to the protocol. The enzyme mixture was then assayed according to the protocol as described in the User Manual.

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