

U.S. Scientific Support: 800-521-0390 scientific.support@lonza.com EU/ROW Scientific Support: +49-221-99199-400 scientific.support.eu@lonza.com Document # Inst-LT07-517 07/11 © 2011 Lonza Walkersville, Inc.

# ToxiLight<sup>™</sup> 100% lysis reagent set

# Safety

THESE PRODUCTS ARE FOR RESEARCH USE ONLY. Not approved for human or veterinary use, for application to humans or animals, or for use in clinical or *in vitro* procedures.

The ToxiLight<sup>™</sup> 100% lysis reagent is intended for use with ToxiLight<sup>™</sup> non-destructive cytotoxicity assay (products LT07-117, LT07-217 and LT17-217). The ToxiLight<sup>™</sup> 100% lysis reagent will provide a total adenylate kinase control when used in conjunction with the kits.

# Intended use

- To produce a 100% adenylate kinase control when using the ToxiLight<sup>™</sup> non-destructive cytotoxicity assay.
- The ToxiLight<sup>™</sup> 100% lysis reagent should be included as a control at each time point within your cytotoxicity assay.
- After addition proceed with the ToxiLight<sup>™</sup> assay according to the instructions in the ToxiLight<sup>™</sup> non-destructive cytotoxicity assay kit.

# Instructions for use

For research use only. Not for use in diagnostic procedures.

LT07-517 Kit contents sufficient for 200 assay points. Store contents at 2℃-8℃. Do not freeze.

#### **Kit contents**

- ToxiLight<sup>™</sup> 100% lysis reagent (LT27-239). 1 x 10 ml bottle.
- Tris acetate buffer (LT27-003). 1 x 50 ml bottle.

The kit contents should be stored at 2 °C-8 °C.

# **Reagent storage**

#### ToxiLight<sup>™</sup> 100% lysis reagent (LT27-239)

This is provided ready for use. Store at  $2^{\circ}$ -8° when not in use.

#### Tris acetate buffer (LT27-003)

This is provided ready for use. Store at  $2^{\circ}$ -8°C when not in use.

#### Instructions for use

# Protocol for 96 well plate using recommended culture volume.

- To achieve a total adenylate kinase control, add 50 µl of the ToxiLight<sup>™</sup> 100% lysis reagent to 100 µl of the culture controls.
- To the remaining wells, add 50 µl of tris acetate buffer. (This is a required volume correction to ensure that total volumes in sample and control wells are equal).
- 3. Incubate at room temperature for 10 minutes to ensure complete lysis.
- Add 100 µl AK detection reagent to all wells and wait 5 minutes.
- 5. Read luminescence.

For other culture volumes use at a ratio of 1:2, ToxiLight<sup>™</sup> 100% lysis reagent to cell culture sample volume. e.g. for a 384 well plate using the recommended sample volume of 25 µl add 12.5 µl of the ToxiLight<sup>™</sup> 100% lysis reagent.

Care should be taken to avoid cross contamination of reagents with the ToxiLight<sup>™</sup> 100% lysis reagent. Employ good laboratory practice at all times.