Regencedlab

USER MANUAL

Product: Trypsinizer

Components: EDTA, NaCl, KCl, Na₂HPO4, K₂HPO4, Recombinant Trypsin

Reference number: 126 (100 mL); **127** (500 mL); **412** (100 mL bottle)

Size: 100 mL bottle, 500 mL bottle.

Intended Use:

For research or manufacturing purposes only.

Overview Product:

Trypsinizer is a solution used to dissociate loosely adherent cells in cell cultures without causing cell damage and the aging ensuing trypsinization.

The solution has a low endotoxin level ($\leq 1.0 \text{ EU/mL}$), is free from mycoplasma contamination, meets the sterility criteria (0 CFU), has a pH range of 6.5 – 7.5, and an osmotic pressure of 286 - 356 (mOsm/kg).

The solution contains recombinant trypsin.

Known Applications:

Trypsinizer has been evaluated on mesenchymal stem cells, fibroblasts, keratinocytes, neural stem cells, and cancer cell lines MCF-7 and HepG2, demonstrating effective dissociation of adherent cells into single cell suspensions.

Reconstitution, Dilution, and Mixing:

The product is supplied in a 1X form, requiring no further dilution or addition of any components prior to usage.

Materials and Reagents Required but Not Provided:

Not applicable

Handling and Storage:

Storage temperature: -20° C - 8° C.

Expiration date: best before 18 months from manufacture date.

Instructions for Use:

- 1. Warm the Trypsinizer solution before use. It is recommended to aliquot the solution into a sterile container and incubate it in a heat block or a water bath at 37°C.
- 2. Inspect the cell cultures using a microscope.
- **3.** Remove the old culture medium from the cell cultures.
- **4.** Gently wash the cells by adding the Washing Buffer. Then, remove the washing solution.
- **5.** Add an appropriate amount of Trypsinizer into the culture vessels. Gently swirl to evenly coat the surface of the cell cultures with the Trypsinizer solution.
- 6. Incubate the culture vessels in a 37°C incubator for 1-5 minutes.
- 7. Observe the cells under a microscope (spherical single cell suspensions should be achieved at this point).
- 8. Transfer the cell suspension into centrifuge tubes. Centrifuge at 1500 RPM for 5 minutes.
- **9.** Remove the supernatant and resuspend the cell pellet in an appropriate solution for the intended purpose.

Precautions:

Do not use the product if the packaging is damaged or cracked, or if the solution appears turbid.

First Aid Measure:

Not applicable

Hazards Identification

The symbols present on the kit are explained below

	LOT	漆	REF
Use By:	Batch code	Keep away from light	Catalog number
X	i	\wedge	STERILE A
Temperature Limitation	Consult instructions for use	Caution, consult accompanying documents	Sterilized using aseptic processing techniques

Related-products

Product Name	Reference Number	
Trypsin/EDTA 0,25%		
100 mL	431	
500 mL	430	
Deattachment		
100 mL	120; 411	
500 mL	121	
MSCCult I		
500 mL	108; 408	
MSCCult II		
500 mL	296; 409	
Washing Buffer		
100 mL	149	
500 mL	150	
ThawBest		
100 mL	142	
500 mL	143	

To purchase other products, please visit: http://biomedmart.org

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