

**Neo Trypsin-EDTA (0.5 %) in DPBS (10x)**

#Cat: NB-58-0109      Size: 100ml

**General Information**

Neo Trypsin-EDTA solutions are used to detach adherent cells from culture surfaces. They are composed of natural porcine pancreas-derived trypsin and EDTA. The concentration of trypsin necessary to dislodge cells from their substrate is dependent primarily on the cell type and the age of the culture. Various formulations should be tested to determine the best product for a specific application.

**Product Specifications**

<b>Appearance</b>	Clear frozen liquid
<b>Storage and shelf life</b>	Store at $\leq -15^{\circ}\text{C}$ . Avoid repeated freeze-thaw cycles. Preparation of aliquots recommended. Once opened, store at $4^{\circ}\text{C}$ and use within 2-4 weeks.
<b>Shipping conditions</b>	Frozen (Dry ice)
<b>Thawing</b>	$+37^{\circ}\text{C}$ water bath or overnight at $+2^{\circ}\text{C}$ to $+8^{\circ}\text{C}$ . Swirl gently to homogenize.
<b>Working Concentration</b>	Recommended final concentration: 1 x

**Formulation**

<b>Components</b>	<b>Concentration mg/L</b>
EDTA 2Na	2200.00
KCl	200.00
$\text{KH}_2\text{PO}_4$	200.00
NaCl	8000.00
$\text{Na}_2\text{HPO}_4$	1150.00
Trypsin	5000.00

**Instructions for Use****Prepare 1x solutions from 10x concentrates:**

To prepare an acceptable final 1x solution, perform the following procedure under aseptic conditions.

1. The product can either be thawed in a  $+37^{\circ}\text{C}$  water bath or overnight at  $+2^{\circ}\text{C}$  to  $+8^{\circ}\text{C}$ .
2. Aseptically dilute 100 ml of 10x concentrate with approximately 850 ml of a sterile  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$ -free salt solution (see related products). Mix completely.
3. If necessary, adjust the pH as necessary with 1 N HCl or 1 N NaOH to pH 7.2 – 7.8.
4. Adjust the final volume with the sterile  $\text{Ca}^{2+}$  and  $\text{Mg}^{2+}$ -free salt solution.
5. Dispense the solution into sterile containers. Cap the bottles tightly with sterile closures and store at  $\leq -15^{\circ}\text{C}$ .

## Detachment of adherent cells using Neo Trypsin-EDTA:

This entire procedure should be done in a laminar flow hood using proper aseptic technique.

1. The product can either be thawed in a +37°C water bath or overnight at +2°C to +8°C.
2. Carefully aspirate all of the media from the cell culture flask.
3. Rinse cells with Ca<sup>2+</sup> and Mg<sup>2+</sup>-free salt solution (see related products), aspirate, and discard.
4. Prewarm the 1x trypsin solution in a +37°C water bath. Add enough 1x trypsin solution to completely cover the cells.
5. Incubate the flask at +37°C, or for more sensitive cultures, at room temperature or +2°C to +8°C.
6. When the trypsinization process is complete, cells will appear rounded upon microscopic examination and the solution in the flask will appear cloudy. Check the flask often to avoid overexposure. Trypsin can cause cellular damage and time of exposure should be kept to a minimum. The time required to detach cells from the culture surface is dependent on the cell type, the age of the culture, population density, serum concentration in the growth medium and time since last subculture.
7. Neutralize trypsin either with serum containing medium or trypsin inhibitor. Gently centrifuge the cell suspension and discard the trypsin-containing supernatant.
8. Resuspend the cell pellet with fresh medium and count or culture as desired.

## Related Products

Product	Cat. No.
Neo Dulbecco's PBS (1x), w/o Ca & Mg, w/o Phenol Red	NB-58-0022
Neo Hank's Balanced Salts, w/o Ca & Mg, w/o Phenol Red	NB-58-0072

## Precautions and Disclaimer

The product is for research and further manufacturing only.

## Help Needed?

If you have any further questions regarding this product, please do not hesitate to contact our cell culture experts by email ([info@neo-biotech.com](mailto:info@neo-biotech.com)) or phone (+33 9 77 40 09 09).