

en

TONGUE DEPRESSOR

Product used as a tongue depressor to achieve correct visualization of the oral area and to prevent contamination during throat and nasopharynx sampling. Single-use product. Tongue depressors are medical devices (MD). Only for professional users.

Intended purpose:

Product intended for the displacement of the tongue to facilitate the exploration of adjacent organs or tissues. The depressor is intended to be used in clinical and hospital environments.










Presentation:

Available in two materials:

- High-quality wood, with a shatterproof consistency and rounded edges to prevent aggression in the oral area. They are packaged in cardboard boxes
- Non-toxic, biocompatible plastic (polystyrene) with rounded edges to prevent aggression in the oral area. They are packaged in minigrip lockable bags.


General precautions:

1. Reuse of this product may cause infections.
2. The use of the depressor does not present any difficulties for the intended users.
3. Not suitable for any application other than its intended use.
4. It should not be used if dirt is observed in the depressor.
5. Discard the product if any is found to be broken or chipped.
6. It has a sufficient mechanical consistency to ensure the impossibility of breaking during use. Although unlikely, applying excessive pressure during use could break the depressor. In case of breakage, discard immediately.
7. Depressants do not require special storage conditions.
8. Due to the intended use of the product, the recipients belong to the healthcare sector and have biowaste containers, where the used product must be disposed of and managed as biowaste. Depressor, once used, must be decontaminated or treated as contaminated hospital waste. It is the responsibility of each laboratory to handle, treat and dispose of waste according to regulations. Unused depressants may be considered non-hazardous goods and may be disposed of based on this criterion.

 Do not re-use	 CE marking	 Manufacturer
 Medical Device	 Consult instructions for use on the website www.deltalab.es/eifus	 Use-by-date
 Catalogue number	 Unique device identifier	 Batch code

In case of a serious incident* related to the product, notify to Deltalab, S.L. as well as the competent authority of the State in which the user is established.

*A "serious incident" is understood as one that entails the death, or serious deterioration of the health of the patient or user or a serious threat to public health.



DELTALAB, S.L.
Plaza de La Verneda 1, Pol. Ind. La Llana,
08191 Rubí, Barcelona. España/Spain.
info@deltalabgroup.com - www.deltalab.es