



GERMOCID LINE - TECHNICAL DATASHEET

GERMOCID ENZIMATICO

GERMOCID ENZIMATICO is a Class I Medical Device in compliance with Regulation (UE) 745/2017.

GERMOCID ENZIMATICO is a concentrated solution based on proteolytic enzymes studied for surgical and orthodontic instruments cleansing, before disinfection/sterilization procedures.

GERMOCID ENZIMATICO is also effective on dry residues and on persistent dirt. It can be used on glassware, plastic and metal, probes, optical fibers instruments, equipment for anesthesia, inhalant therapy, hemodialysis, endoscopy and urology.

GERMOCID ENZIMATICO can also be used in ultrasound equipment.



APPLICATION

Enzyme-based detergent for invasive surgical/dental instruments.

INSTRUCTIONS FOR USE

Dilute GERMOCID ENZIMATICO in a proportion of 0,6-10 % in one liter of tap water, possibly warm, and mix thoroughly.

Before dipping the instruments in the prepared solution, it is advised to decontaminate them with a specific product and rinse with cold water, then dip them in the solution of GERMOCID ENZIMATICO for 10 minutes; in presence of dry residues or persistent dirt extend immersion time to 1 hour, and if necessary, use a brush. Complete the processes with a series of rinses, the last one with sterile water.

PACKAGING

Internal Code:	Primary packaging:	Secondary packaging:
B411	1000 ml bottle	Box containing 12 bottles
B413	3000 ml canister	Box containing 4 canisters

Barcode	ITF	Box size (mm)	Pallet composition
8009110025097	08009110025097	335 w x 253 d x 274 h	40 pcs (4 floors x 10)
8009110026193	08009110026193	331 w x 307 d x 268 h	40 pcs (4 floors x 10)

All primary packaging is according to the specifications provided by the Pharmacopoeia. The materials do not contain allergens and are perfectly compatible with all components of the formulation.



Germodis®
S.P.A.

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COMPOSITION

Non-ionic surfactants 15-30%; anionic surfactants 5-15%; enzymes; perfume (Limonene, Citral), preservative (Benzisothiazolinone).

PRODUCT FORM

Clear green liquid with lemon fragrance.

VALIDITY

3 years. The indicated period of validity refers to the product stored in its container and properly used and stored.

HOW TO STORAGE

Store in a cool, dry place, away from heat sources.

QUALITY CONTROL

The components (raw materials, containers, labels, etc.) and the processing steps of each lot are regularly and carefully checked internally following the procedures of the Organization Quality Management System certified UNI EN ISO 9001 and UNI EN ISO 13485.

AUTHORIZATIONS

Class I Medical Device in compliance with Regulation (UE) 745/2017.

CONFIDENTIAL INFORMATION FOR HEALTH WORKERS AND PROFESSIONAL USERS