



SPORIGERM[®]MANI

Antiseptic soap
for the disinfection of the hands
**wide range
disinfecting action**

Rev. 0 – 09/2019

INDICATIONS

SPORIGERM MANI disinfectant soap is a liquid antiseptic with a disinfectant action, removing and controlling the bacterial load present on the skin. SPORIGERM MANI disinfectant soap respects the hydrolipidic film, without altering the skin physiological balance (pH 5.5). It therefore allows frequent washing.

APPLICATIONS

SPORIGERM MANI disinfectant soap is also suitable for the disinfection of surgeons hands before operating, and the disinfection and cleansing of the hands of medical personnel, in order to avoid cross infections.

INSTRUCTIONS FOR USE

Use as it is. Apply about 5ml of product on wet hands for at least 5 minutes and, after washing thoroughly (using a brush for a thorough cleaning of the nails), rinse accurately.

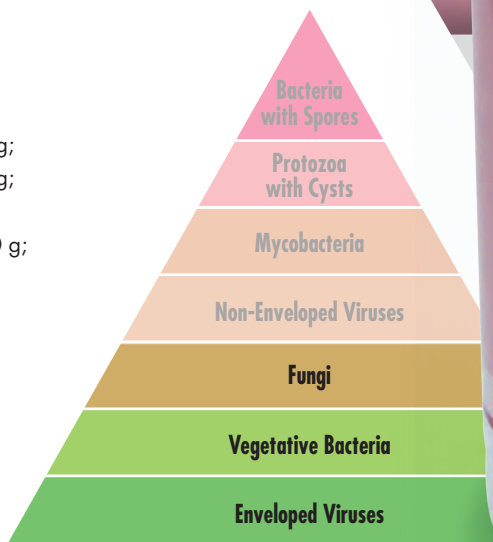
PACKAGING

SporigerM Mani
Code 500007 6, 1,000 ml bottles + 2 dispensers

All the bottles are in high density polyethylene (HDPE) in compliance with the technical specifications established by Farmacopea. This material does not contain latex and is fully compatible with all the components of the formulated.

COMPOSITION 100 grams of product

Alkyl-benzyl-di-methyl-ammoniumchlorid 0.5 g;
Orthophenyl phenol 0.5 g;
Fragrance; Co-formulants;
Purified water, q.s. at 100.00 g;



INFORMATION FOR THE EXCLUSIVE USE OF HEALTHCARE PROFESSIONALS



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ACTION

The alkyl-benzyl-di-methyl-ammoniumchlorid, together with the Orthophenyl phenol carries out a high bactericidal and fungicidal action guaranteeing a full range of action towards Gram-positive, Gram-negative, bacteria and fungi.

TEST

EN1500; EN13727; EN1650; PATCH TEST.

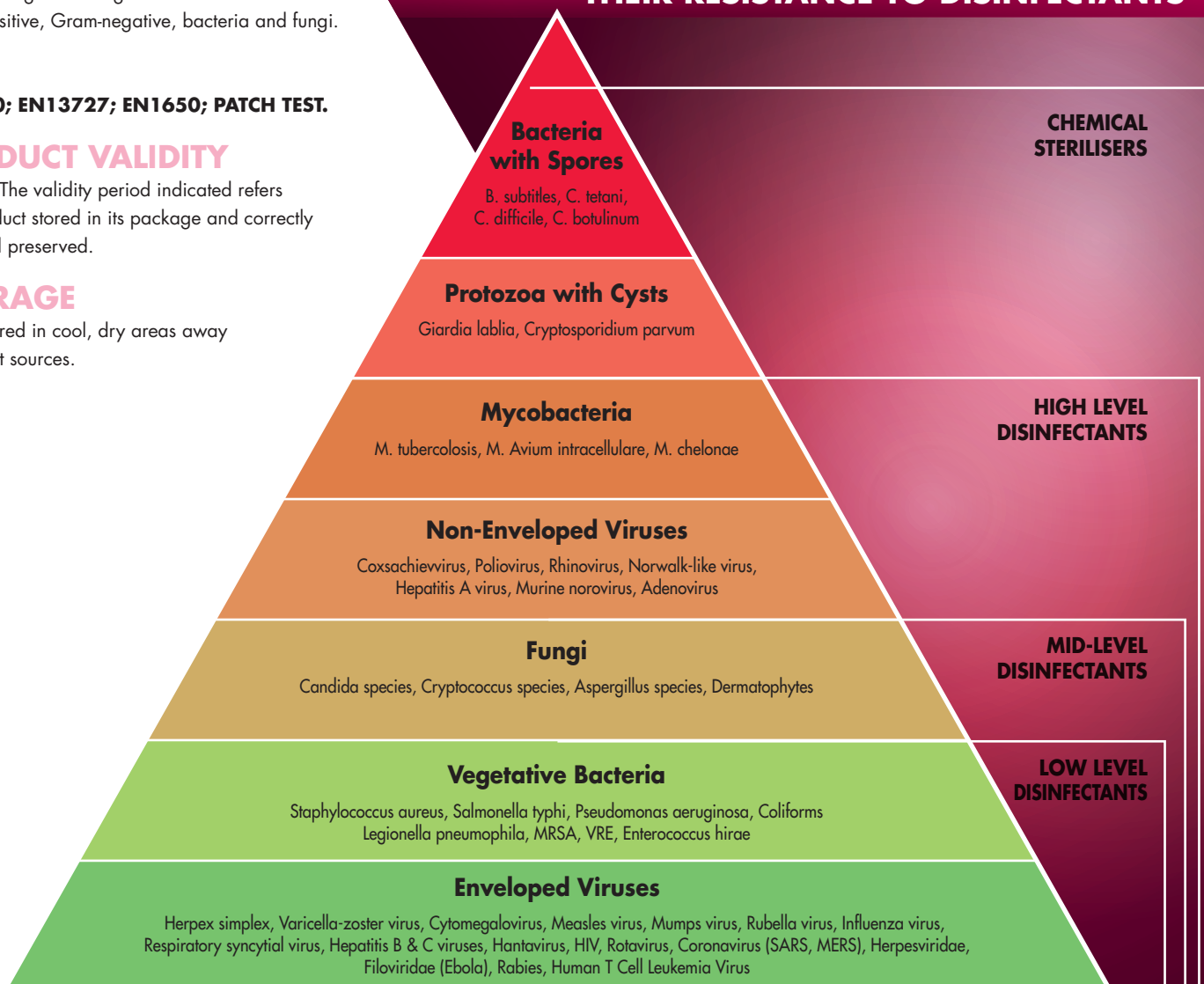
PRODUCT VALIDITY

5 years. The validity period indicated refers to a product stored in its package and correctly used and preserved.

STORAGE

To be stored in cool, dry areas away from heat sources.

PATHOGENIC AGENTS CLASSIFIED ACCORDING TO THEIR RESISTANCE TO DISINFECTANTS



PRECAUTIONS

External use only. Do not use on damaged skin and mucous membranes. Do not swallow. The product is not compatible with surfactants and anionics.

QUALITY CONTROL

All components (raw materials, containers, labels etc.), together with the processing phases of each batch are punctually and thoroughly checked internally according to the corporate UNI EN ISO 9001 certified Quality Management System.

AUTHORISATIONS

Medical surgical device
– Health Ministry Registration
n. 19.570

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