

SPORIGERM®MAN

Rev. 0 - 09/2019

Antiseptic soap for the disinfection of the hands

wide range disinfecting action

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

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

 Protozoa with Cysts

Mycobacteria

Non-Enveloped Viruses

Fungi

Vegetative Bacteria

Enveloped Viruses

Antiseptic liquid for the disinfection of hands and skin for surgical use

Antisettico liquido per la disinfezione delle mani e della cute per uso chirurgico

INFORMATION FOR THE EXCLUSIVE USE OF HEALTHCARE PROFESSIONALS

Manufactured by:

GERMO S.p.A.

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Distributed by:

IDS S.p.A.

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www.idsdental.it - info@idsdental.it



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CHEMICAL

STERILISERS

HIGH LEVEL

MID-LEVEL

LOW LEVEL

DISINFECTANTS

DISINFECTANTS

DISINFECTANTS

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.

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

Bacteria with Spores

B. subtitles, C. tetani, C. difficile, C. botulinum

Protozoa with Cysts

Giardia lablia, Cryptosporidium parvum

Mycobacteria

M. tubercolosis, M. Avium intracellulare, M. chelonae

Non-Enveloped Viruses

Coxsachievvirus, Poliovirus, Rhinovirus, Norwalk-like virus, Hepatitis A virus, Murine norovirus, Adenovirus

Fungi

Candida species, Cryptococcus species, Aspergillus species, Dermatophytes

Vegetative Bacteria

Staphylococcus aureus, Salmonella typhi, Pseudomonas aeruginosa, Coliforms Legionella pneumophila, MRSA, VRE, Enterococcus hirae

Enveloped Viruses

Herpex simplex, Varicella-zoster virus, Cytomegalovirus, Measles virus, Mumps virus, Rubella virus, Influenza virus, Respiratory syncytial virus, Hepatitis B & C viruses, Hantavirus, HIV, Rotavirus, Coronavirus (SARS, MERS), Herpesviridae, Filoviridae (Ebola), Rabies, Human T Cell Leukemia Virus

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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