



SPORIGERM[®] GEL MANI PMC

Rev. 4 - 10/2020

INDICATIONS

SPORIGERM GEL MANI is a Medical Surgical Device used for a fast disinfection of the hands. Very useful while travelling and on any occasion where washing your hands is not possible.

SPORIGERM GEL MANI

Is indicated for the disinfection of the hands of medical and nursing staff before and after contacts with patients and for those operating in hospitals (medical practices, wards, emergency rooms, laboratories). As a prevention measure it can also be used in case of contact with sick patients and potentially infected objects. Also suitable for the food industry for the disinfection of the operators' hands. The alcohol content amounts to 70% in volume (equal to 62% in weight) and it's indicated for the aseptic rubbing of hands in case of lack of water and soap.

SPORIGERM GEL MANI is ready to use, drying quickly and leaving your hands fresh and soft.

DERMATOLOGICALLY TESTED

APPLICATIONS

For the disinfection of hands in the absence of water at home, in the food industry and in hospitals.

INSTRUCTIONS FOR USE

To be used without water. Apply at least 3 ml of pure product directly on the hands and rub until completely dry.

PACKAGING

Sporigerm Gel Mani

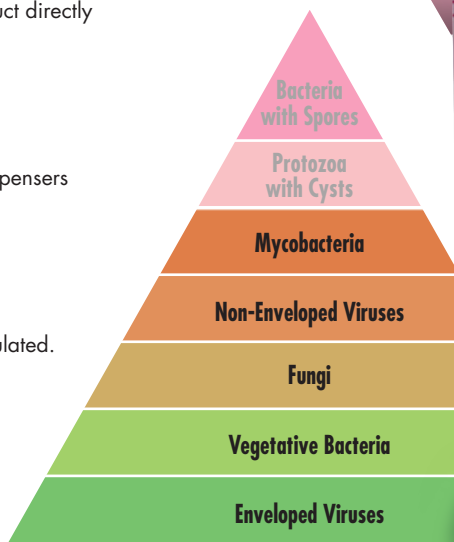
Code 500008 12, 500 ml bottles + 4 dispensers

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 g of product contain:
Denatured ethyl alcohol. 62.00 gr.
Co-formulants, Purified water q.s. for 100 gr.

Antiseptic gel. 70% volume (62% weight) hydroalcoholic solution
No water needed



INFORMATION FOR THE EXCLUSIVE USE OF HEALTHCARE PROFESSIONALS

Manufactured by:
GERMO S.p.A.
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Antiseptic gel. 70% volume (62% weight) hydroalcoholic solution
No water needed

CHARACTERISTICS

Colourless gel.

ACTION

SPORIGERM GEL MANI

Is a disinfectant and detergent with a broad range of action with regards to pathogenic micro-organisms (bacteria, mycobacteria, fungi and viruses), due to the high alcohol content, equal to 70% in volume.

TEST

**EN 12054; EN 1500; EN 1650; EN 14348;
EN 14476; EN 12791; PATCH TEST.**

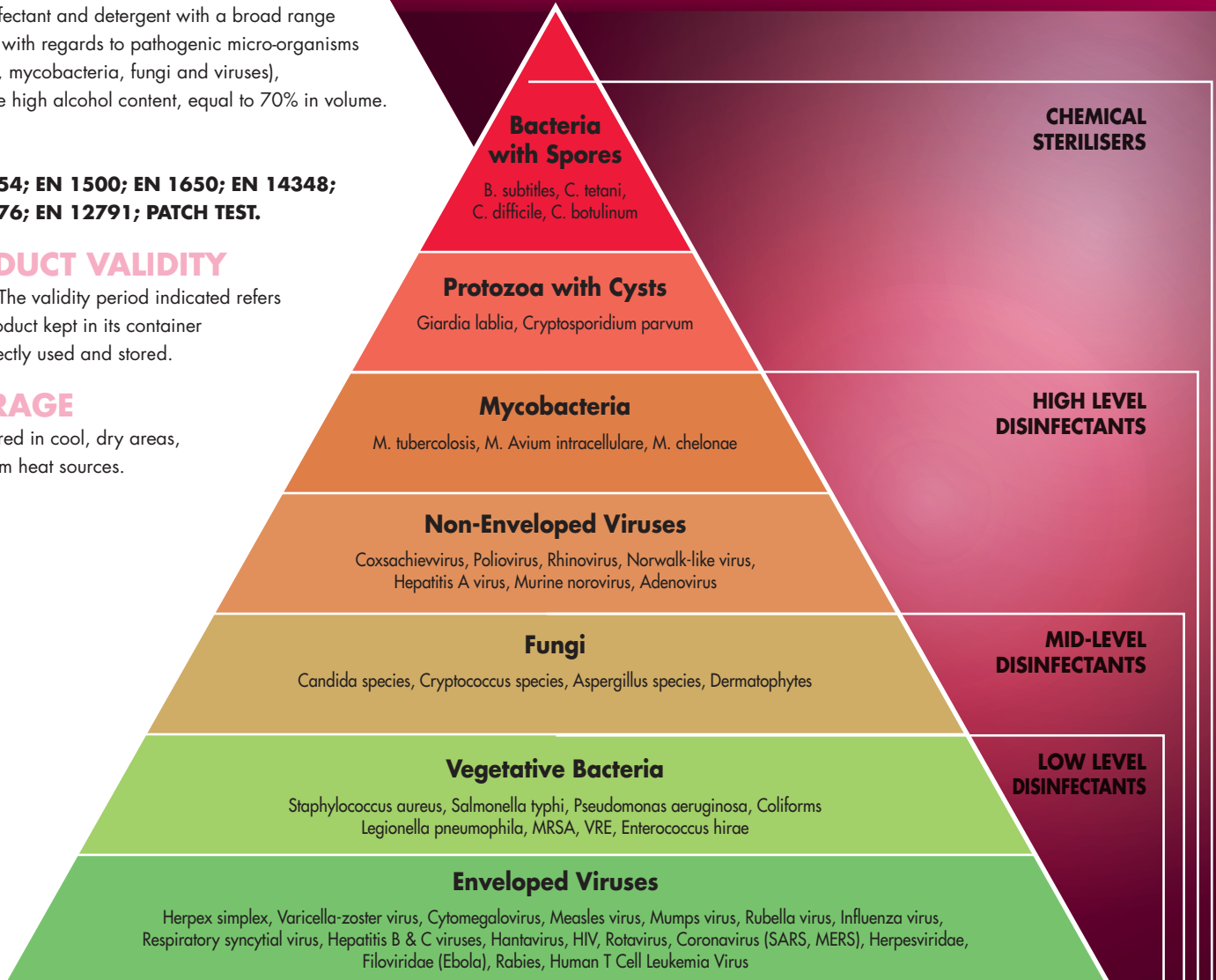
PRODUCT VALIDITY

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

STORAGE

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

PATHOGENIC AGENTS CLASSIFIED ACCORDING TO THEIR RESISTANCE TO DISINFECTANTS



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.
Ministry of Health Registration n. 19.636

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