

Note: Leaflet for RMP is discontinued in the commercial and PS pack.  
This is only relevant for HCP promotion purpose.

For the use only of a Registered Medical Practitioner or a Hospital or a Laboratory

**Spores of Poly-antibiotic resistant *Bacillus Clausii***  
**ENTEROGERMINA®**  
**2 billion / 5 mL oral suspension.**

**FOR ORAL USE**  
**DO NOT INJECT**

#### **COMPOSITION**

One 5mL mini bottle contains spores of polyantibiotic-resistant *Bacillus clausii* - 2 billion (strains: O/C, N/R, SIN and T)

#### **PHARMACEUTICAL FORM**

Oral suspension

#### **CLINICAL PARTICULARS**

##### **Therapeutic indications**

For the treatment of alterations in the intestinal bacterial flora.

##### **Posology and method of administration**

**Adults:** 2-3 mini bottles per day

**Children:** 1-2 mini bottles per day

**Infants:** 1-2 mini bottles per day

unless prescribed otherwise by the doctor.

Administration at regular intervals, Take the mini bottle as it is or dilute it in water or other beverages (e.g., milk, tea, orange juice).

This medicinal product is for oral use only. Do not inject or administer through other routes (see Section Special warnings and special precautions for use).

##### **Contraindications**

Hypersensitivity to the active substance or to any of the excipients.

##### **Special warnings and special precautions for use**

###### **Special warnings**

Severe anaphylactic reactions, such as anaphylactic shock, have occurred with incorrect route of administration.

Any presence of visible corpuscles in the mini bottles of Enterogermina® is due to aggregates of *Bacillus clausii* spores; it does not therefore – suggest that the product has been altered.

Shake the mini bottle before use.

There have been reports of bacteremia, septicemia or sepsis in patients taking *Bacillus clausii* who are immunocompromised or are hospitalized due to a serious illness. Enterogermina® should be used in these patients only if the potential benefits outweigh the potential risks.

During treatment with antibiotics, it is recommended that the preparation be administered between antibiotic doses.

##### **Interactions with other medicinal products and other forms of interaction.**

No interaction studies have been performed.

## **Reproduction**

### **Pregnancy**

Limited data are available on the use of probiotics including Enterogermina® in pregnant women. However, no conclusions can be drawn regarding whether or not Enterogermina® is safe for use during pregnancy. Enterogermina® should be used during pregnancy only if the potential benefits to the mother outweigh the potential risks, including those to the fetus.

### **Lactation**

There are limited available data on the presence of Enterogermina® in human milk, milk production, or the effects on the breastfed infant. However, no conclusions can be drawn regarding whether or not Enterogermina® is safe for use during breastfeeding. Enterogermina® should be used during breastfeeding only if the potential benefits to the mother outweigh the potential risks, including those to the breastfed child.

### **Driving a vehicle or performing other hazardous tasks**

Enterogermina® has no influence on the ability to drive and use machines.

### **Undesirable effects**

The following CIOMS frequency rating is used, when applicable:

Very common  $\geq 10\%$ ; Common  $\geq 1$  and  $< 10\%$ ; Uncommon  $\geq 0.1$  and  $< 1\%$ ;

Rare  $\geq 0.01$  and  $< 0.1\%$ ; Very rare  $< 0.01\%$ ; Not known (cannot be estimated from available data).

Skin and subcutaneous tissue disorders:

During post marketing experience, hypersensitivity reactions, including rash, urticaria and angioedema have been reported.

Infections and infestations:

Not known: Bacteremia, septicemia or sepsis in immunocompromised patients or those hospitalized due to a serious illness.

### **Overdose**

No cases of overdose have been reported.

## **PHARMACOLOGICAL PROPERTIES**

### **Pharmacodynamic properties**

#### **Pharmacotherapeutic category: antidiarrhoeal microorganisms**

Enterogermina® is a preparation consisting of a suspension of 4 spore strains (SIN, O/C, T, N/R) of *Bacillus clausii*, which naturally occur in the intestine and is non-pathogenic.

When administered orally, *Bacillus clausii* spores cross the barrier of the acidic gastric juice due to their high resistance to both chemical and physical agents, and reach the intestinal tract unharmed, where they are transformed into metabolically active vegetative cells.

Spores can survive heat and gastric acidity, by nature. In an in vitro validated model, *Bacillus clausii* spores were shown to survive in a simulated gastric environment (pH 1.4-1.5) up to 120 minutes (survival rate of 96%). In a model that simulates the intestinal environment (bile and pancreatin saline - pH 8), *Bacillus clausii* spores showed their ability to multiply further compared to the initial amount, in a statistically significant way (from 10<sup>9</sup> to 10<sup>12</sup> CFU – Colony-Forming Units), starting from 240 minutes after incubation.

In a study conducted in 20 subjects, it was found that in humans, *Bacillus Clausii* spores persist in the intestine and can be found in faeces until 12 days after a single oral administration.

The administration of Enterogermina® helps to restore the intestinal microbial flora altered by dysmicrobism, also known as dysbiosis, resulting from antibiotic therapy and which may be associated with gastrointestinal symptoms, e.g. diarrhoea, abdominal pain and increased air in the intestine.

In two open-label, randomized, and controlled clinical studies, Enterogermina® was shown to reduce the duration of acute diarrhoea in children older than 6 months.

Taken during antibiotic treatment and 7 to 10 days thereafter, Enterogermina® was shown to reduce the incidence of abdominal pain and diarrhoea associated with antibiotic treatment.

The following 2 main mechanisms contribute to the effect of *Bacillus clausii* in restoring the intestinal bacterial flora.

### **Growth Inhibition of Pathogenic Bacteria**

The three *B. clausii* supposed mechanisms of action are: colonization of free ecological niches, which are made no longer available by the growth of other microorganisms; competition in adhesion to epithelial cells, which is particularly relevant for spores in the early and intermediate stages of germination; production of antibiotics and/or enzymes that are secreted within the intestinal environment. In an in vitro study, *Bacillus clausii* spores were shown to have antagonistic activity against Gram-positive bacteria - *Staphylococcus aureus*, *Clostridium difficile*, *Enterococcus faecium* - by producing bacteriocins and antibiotics such as clausin.

### **Immunomodulatory activity**

Orally administered *Bacillus clausii* spores were shown to stimulate the production of Interferon-gamma and increase the CD4+ T Lymphocyte proliferation, in in vitro and in vivo murine models.

*Bacillus clausii* also showed the ability to produce several B vitamins, helping to correct vitamin deficiencies in the body resulting from an imbalance in the intestinal bacterial flora.

Furthermore, the high level of artificially induced heterologous resistance to antibiotics makes it possible to create the therapeutic conditions to prevent alteration of the intestinal microbial flora following the selective action of antibiotics, especially those with a broad spectrum of action, or to restore it.

Given this antibiotic resistance, Enterogermina® may be administered in between two subsequent administrations of antibiotics.

Antibiotic resistance refers to: penicillins, if not in combination with beta-lactamase inhibitors, cephalosporins (mostly with partial resistance), tetracyclines, macrolides, aminoglycosides (except for gentamicin and amikacin), chloramphenicol, thiamphenicol, lincomycin, clindamycin, isoniazid, cycloserine, novobiocin, rifampicin, nalidixic acid and pipemidic acid (intermediate resistance), metronidazole.

## **PHARMACEUTICAL PARTICULARS**

### **List of excipients**

Purified water Ph. Eur

### **Incompatibilities**

None

### **Storage**

Do not store above 30 °C.

KEEP OUT OF THE REACH OF CHILDREN.

### **Instructions for use**

Shake the mini bottle before use. After mini bottle has been opened, the preparation should be taken shortly to avoid any contamination of the suspension.

For oral use only. Do not inject.

**Manufactured by:**

Sanofi S.R.L. - Viale Europa, 11 - 21040 - Origgio (VA) - Italy

**Importer:**

Sanofi Healthcare India Private Limited, Gala No. 4, Ground Floor, Building No. B1, Citylink Warehousing Complex, S No.121/10/A,121/10/B & 69, NH3, Vadape. Tal: Bhiwandi - 16(Thane Z5) Pin :421302.

**Source:**

1. Bacillus clausii CCSI v3 LRC dated 12<sup>th</sup> March 2020
2. Italian SmPC dated February 2021

**Updated:** April 2022