

Technical Data Sheet

Revamil® Gel

Description

100% pure honey for the treatment of wounds with honey

Intended use

Treatment of chronic, infected, and acute wounds, as well as minor burns

Ingredients

- Mel (honey)

Product specifications

Packaging	Aluminum tube 18 gram (packed individually) / Aluminum tube 5 gram (packed per 27 units)
Composition	INCI: mel (honey)
Shelf life	3 years unopened 3 months after opening
Quality	Revamil® Gel is a CE registered product within the European Union. This registration guarantees production under strict quality measurements.
Pharmaceutical ingredients	Revamil® Gel does not contain pharmaceutical ingredients
Components of animal origin	Revamil® Gel does not contain ingredients of animal origin (definition of animals excludes honeybees, according to EU guidelines)
Preservatives	No preservatives are added to Revamil® Gel.

Application instructions

1. Before applying Revamil Gel first clean the wound.
2. Spread a thin layer of Revamil Gel on the secondary wound dressing.
3. Revamil Gel can also be applied directly to the wound and then covered with a secondary dressing.
4. Attach the secondary dressing in the normal way.
5. Depending on the nature of the wound, replace Revamil Gel every day or every two days.

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Contra indications

Do not use Revamil® Gel if the patient is known to be allergic to honey.

Side effects

- During the first minutes after application Revamil® Gel can cause a light stinging sensation
- In case of unexpected side effects after use contact your doctor or the manufacturer

Safety precautions

- Do not use Revamil® Gel if the tube is damaged
- Always replace the top on the tube after use
- Do not allow the tube to come into direct contact with the surface of the wound
- Once the tube has been opened use the contents within 3 months
- Do not treat different patients with the same tube

Product



Storing conditions

Store between 10°C and 30 °C

Classification

CE Class IIb

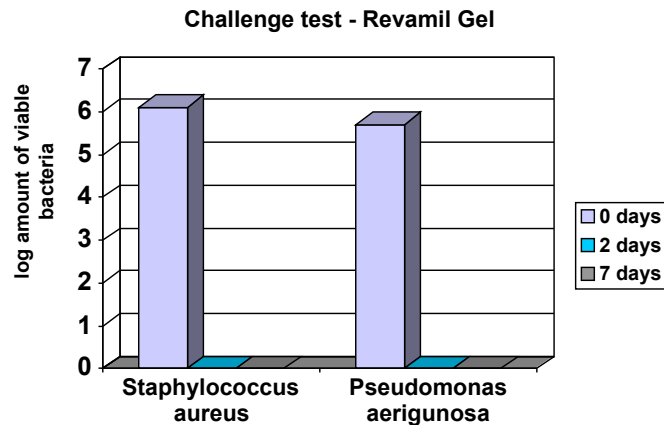
Notified Body

DEKRA Certification BV (No. 0344), Arnhem, the Netherlands

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Antibacterial activity

Challenge test with *Staphylococcus aureus* and *Pseudomonas aeruginosa* showed log reduction of viable bacteria of 6 after 48 hrs.



Manufacturer (official CE status)

Bfactory Health Products B.V.
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3911 TZ Rhenen
The Netherlands

Approved by:

Name: W.J.P. Bosma
Function: Director
Date: 20 July 2024
Signature:

Company stamp:

