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VarroMed 44 mg/ml + 5 mg/ml - bee-hive dispersion

Authorised

- Oxalic acid dihydrate
- Formic acid

Product identification

Medicine name:
VarroMed 44 mg/ml + 5 mg/ml - bee-hive dispersion

Active substance:
Oxalic acid dihydrate

Formic acid

Target species:
Honey bee

Route of administration:
In-hive use

Product details

Active substance and strength:
Oxalic acid dihydrate
43.75 milligram(s) / 1.00 Bottle

Formic acid

5.09 milligram(s) / 1.00 Bottle

Pharmaceutical form:

Bee-hive dispersion

Withdrawal period by route of administration:

In-hive use:

- **Honey bee**
  - Honey. 0 day _ Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AG30

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Croatia , Czechia , Denmark , France , Germany , Greece , Hungary , Ireland , Italy , Lithuania , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:555 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
BeeVital GmbH

Marketing authorisation date:
2/02/2017

Manufacturing sites for batch release:
Lichtenheldt GmbH Pharmazeutische Fabrik
Labor LS SE & Co. KG

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
2/02/2017

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
Published on: 15/10/2026
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