File downloaded on 2026-11-20
Source URL: https://medicines.health.europa.eu/veterinary/en/700000011138

Varroxal 0,71 g/g bee-hive powder

Authorised

- Oxalic acid dihydrate

Product identification

Medicine name:
Varroxal 0,71 g/g bee-hive powder

Active substance:
Oxalic acid dihydrate

Target species:
Honey bee

Route of administration:
Nebulisation use

Product details

Active substance and strength:
Oxalic acid dihydrate
1.00 gram(s) / 1.00 gram(s)

Pharmaceutical form:
Bee-hive powder

Withdrawal period by route of administration:

Nebulisation use:

- **Honey bee**
  - Honey. no withdrawal period Nul dagen

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AG03

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Cardboard box with a 200 g HDPE bottle closed with an ALU/HDPE seal and a HDPE/ PP turn and lock cap and 1 g measuring spoon
Cardboard box with a 75 g HDPE bottle closed with an ALU/HDPE seal and a HDPE/ PP turn and lock cap and 1 g measuring spoon
Cardboard box with a PET/ALU/PE bag containing 50 x 2g sealed paper-PE sachets.
Cardboard box with a PET/ALU/PE bag containing 10 x 2g sealed paper-PE sachets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Andermatt BioVet GmbH

Marketing authorisation date:
8/09/2023

Manufacturing sites for batch release:
Andermatt BioVet GmbH

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/4010336 7/2023

Date of authorisation status change:
8/09/2023

Reference member state:
Netherlands

Procedure number:
NL/V/0402/001

Concerned member states:
Austria Croatia Czechia Estonia France Germany Greece Hungary Latvia Lithuania Poland Portugal Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents