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MAQS Formic Acid 68.2g Beehive Strips for Honey Bees

Not authorised

- Formic acid

Product identification

Medicine name:
MAQS Formic Acid 68.2g Beehive Strips for Honey Bees

Active substance:
Formic acid

Target species:
Honey bee

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Formic acid
68.20 gram(s) / 1.00 Strip

Pharmaceutical form:
Bee-hive strip

Withdrawal period by route of administration:

Cutaneous use:

- **Honey bee**

  - Honey. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AG01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

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Authorisation status:

Surrendered

Authorised in:

France

Package description:

A polypropylene/aluminium foil/polypropylene laminatedsachet containing two strips Pack size Polypropylene tubs containing 2 sachets (4 strips).
A polypropylene/aluminium foil/polypropylene laminatedsachet containing two strips Pack size Polypropylene tubs containing 10 sachets (20 strips).
A polypropylene/aluminium foil/polypropylene laminated sachet containing two strips Pack size Box containing 30 sachets in a plastic liner, packaged in cardboard box (60 strips).
A polypropylene/aluminium foil/polypropylene laminated sachet containing two strips Pack size Box containing 10 sachets in plastic liner, packaged in cardboard box (20 strips).
A polypropylene/aluminium foil/polypropylene laminatedsachet containing two strips Pack size Box containing 2 sachets in plastic liner, packaged in cardboard box (4 strips).

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
NOD Apiary Ireland Limited

Marketing authorisation date:
15/05/2014

Manufacturing sites for batch release:
Animal Health Distributors Limited
Lohmann Pharma Herstellung GmbH

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

Authorisation number:
FR/V/3161438 4/2014

Date of authorisation status change:
4/07/2023

Reference member state:
Ireland

Procedure number:
IE/V/0518/001

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

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 3/05/2024
Updated on: 14/03/2026
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