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)
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