MILBETAB 16 MG/40 MG FILM-COATED TABLETS FOR CATS

Authorised

- Milbemycin oxime
- Praziquantel

Product identification

Medicine name:
MILBETAB 16 MG/40 MG FILM-COATED TABLETS FOR CATS
Milbetab 16 mg/40 mg филмирани таблетки за котки

Active substance:
Milbemycin oxime
Praziquantel

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Milbemycin oxime
16.00 milligram(s) / 1.00 Tablet

Praziquantel
40.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP54AB51

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Cardboard box with 2 tablets (1 blister strip of 2)
Cardboard box with 4 tablets (1 blister strip of 4)
Cardboard box with 10 tablets (1 blister strip of 10)
Cardboard box with 20 tablets (2 blister strips of 10)
Cardboard box with 50 tablets (5 blister strips of 10)
Cardboard box with 100 tablets (10 blister strips of 10)
Multipacks of 10 individual packs of 2 tablets
Multipacks of 10 individual packs of 20 tablets
Multipacks of 10 individual packs of 50 tablets
Cardboard box with 4 tablets (2 blisters of 2)
Cardboard box with 10 tablets (5 blisters of 2)
Cardboard box with 20 tablets (10 blisters of 2)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
8/05/2025

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3296

Date of authorisation status change:
8/05/2025

Reference member state:
France

Procedure number:
FR/V/0475/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Finland Germany Greece Hungary Ireland Italy Netherlands Norway Poland Portugal Romania Spain Sweden

Generic of:
600000042407

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

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

Labelling

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