Metomotyl 5 mg chewable tablets for dogs

Authorised

- Metoclopramide hydrochloride

Product identification

Medicine name:
Metomotyl 5 mg chewable tablets for dogs

Active substance:
Metoclopramide hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Metoclopramide hydrochloride
5.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA03FA01

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

Authorisation status:
Valid

Authorised in:
Croatia

Package description:
Cardboard box containing 100 tablets.
Cardboard box containing 10 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
6/07/2021

Manufacturing sites for batch release:
Lelypharma B.V.
Genera d.d.

Responsible authority:
Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:
UP/I-322-05/21-01/495

Date of authorisation status change:
28/11/2025

Reference member state:
Netherlands

Procedure number:
NL/V/0334/001

Concerned member states:
Austria Belgium Croatia Denmark Finland France Germany Norway Poland Slovenia Sweden

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

Documents

Summary of Product Characteristics

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Combined File of all Documents

English (PDF)
Published on: 27/10/2025
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