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Metomotyl 10 mg chewable tablets for dogs

Authorised

- Metoclopramide hydrochloride

Product identification

Medicine name:
Metomotyl 10 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
10.00 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:
Norway

Available in:
Norway

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Dechra Regulatory B.V.

Marketing authorisation date:
27/04/2022

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

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
20-13491

Date of authorisation status change:
27/04/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0334/002

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

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.

Combined File of all Documents

Final PuAR Metomotyl 5 mg, 10 mg tablets NLV0334001-002_v.09102025.pdf