Vomend 5 mg/ml solution for injection for dogs and cats

Authorised

- Metoclopramide hydrochloride

Product identification

Medicine name:
Vomend 5 mg/ml solution for injection for dogs and cats
Vomend Anti-Emeticum 5 mg/ml Oplossing voor injectie
Vomend Anti-Emeticum 5 mg/ml Solution injectable
Vomend Anti-Emeticum 5 mg/ml Injektionslösung

Active substance:
Metoclopramide hydrochloride

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Metoclopramide hydrochloride

5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA03FA01

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

Authorisation status:
Valid

Authorised in:
Belgium

Package description:
50 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.
5 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.
10 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.
30 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.
25 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.
20 ml vial of clear colourless glass type I. Bromobutyl rubber stoppers type I (the stoppers are secured with aluminium caps). 1 vial in a cardboard box.

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:
1/12/2010

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V381805

Date of authorisation status change:
1/12/2010

Reference member state:
Netherlands

Procedure number:
NL/V/0145/001

Concerned member states:
Austria Belgium Germany Italy Luxembourg Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 16/06/2022
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Package Leaflet

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Labelling

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