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 5 mg/ml oplossing voor injectie voor honden en katten

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

Withdrawal period by route of administration:

Intramuscular use:

- **Dog**
- **Cat**

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA03FA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Netherlands

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:
Eurovet Animal Health B.V.

Marketing authorisation date:
30/11/2010

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

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 106887

Date of authorisation status change:
27/01/2022

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

Combined File of all Documents

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

Summary of Product Characteristics

106887 par.pdf
