

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 Injektionslösung für Hunde und Katzen

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:

Medicinal product on medical prescription for renewable delivery

Authorisation status:

Valid

Authorised in:

Austria

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:

13/12/2010

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

AGES

Authorisation number:

8-00913

Date of authorisation status change:

15/11/2015

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

Package Leaflet

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Summary of Product Characteristics

English (PDF)

Published on: 16/06/2022

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Labelling

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