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Metomotyl 2.5 mg/ml solution for injection for cats and dogs

- Metoclopramide hydrochloride

Authorised

Product identification

Medicine name:
Metomotyl 2.5 mg/ml solution for injection for cats and dogs
Active substance:

- Metoclopramide hydrochloride

Target species:

- Dog
- Cat

Route of administration:

- Intramuscular use
- Subcutaneous use

Product details

Active substance and strength:

- Metoclopramide hydrochloride
  2.50
  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:

- Austria

Package description:

- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 50 ml.
- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 5 ml.
- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 10 ml.
- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 25 ml.
- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 20 ml.
- Clear colourless type I glass vial. Red chlorobutyl 20 mm stopper. Aluminium 20 mm cap. Cardboard box containing 1 vial of 30 ml.

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:

- Le Vet. Beheer B.V.

Marketing authorisation date:

- 23/07/2014

Manufacturing sites for batch release:

- Produlab Pharma B.V.

Responsible authority:

- Austrian Agency For Health And Food Safety

Authorisation number:

- 835668

Date of authorisation status change:

- 23/07/2014

Reference member state:

- Netherlands

Procedure number:

- NL/V/0182/001

Concerned member states:

- Austria
- Belgium
- Czechia
- Denmark
- Estonia
- Finland
- Germany
- Greece
- Hungary
- Iceland
- Ireland
- Italy
- Latvia
- Lithuania
- Luxembourg
- Norway
- Poland
- Portugal
- Romania
- Slovakia
- Spain
- Sweden
- United Kingdom (Northern Ireland)

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

Documents

Product information

Package Leaflet

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