

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

Authorised

- Metoclopramide hydrochloride

## Product identification

**Medicine name:**

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

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**Active substance:**

Metoclopramide hydrochloride

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**Target species:**

Dog

Cat

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**Route of administration:**

Intramuscular use

Subcutaneous use

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## Product details

**Active substance and strength:**

Metoclopramide hydrochloride

2.50 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QA03FA01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Ireland

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

12/09/2014

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**Manufacturing sites for batch release:**

Produlab Pharma B.V.

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10475/009/001

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**Date of authorisation status change:**

12/09/2014

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**Reference member state:**

Netherlands

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**Procedure number:**

NL/V/0182/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)