PRILIUM 150 MG POWDER FOR ORAL SOLUTION FOR DOGS

Authorised

- Imidapril hydrochloride

Product identification

Medicine name:
PRILIUM 150 MG POWDER FOR ORAL SOLUTION FOR DOGS
Prilium 150 mg Poeder voor drank
Prilium 150 mg Poudre pour solution buvable
Prilium 150 mg Pulver zur Herstellung einer Lösung zum Einnehmen

Active substance:
Imidapril hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Imidapril hydrochloride
150.00 milligram(s) / 1.00 Bottle

Pharmaceutical form:
Powder for oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09AA16

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Box containing one 0.880 g powder vial and one 2 ml graduated syringe

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vetoquinol

Marketing authorisation date:
14/07/2003

Manufacturing sites for batch release:
VETOQUINOL

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V254204

Date of authorisation status change:
29/07/2015

Reference member state:
France

Procedure number:
FR/V/0133/002

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Italy Luxembourg Poland Portugal Slovakia 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

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

Labelling

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