Semintra 4 mg/ml - Oral solution

Authorised

- Telmisartan

Product identification

Medicine name:
Semintra 4 mg/ml - Oral solution

Active substance:
Telmisartan

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Telmisartan
4.00 milligram(s) / 1.00 Bottle

Pharmaceutical form:
Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QC09CA07

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

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:
Belgium , Czechia , Denmark , Finland , France , Germany , Iceland , Luxembourg , Norway , Poland , Spain , Sweden

Package description:
Packaging:Bottle (HDPE) + dosing syringe, Package_size:1 bottle, Content:100 ml
Packaging:Bottle (HDPE) + dosing syringe, Package_size:1 bottle, Content:30 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
13/02/2013

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
9/11/2016

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

Documents

Combined File of all Documents

English (PDF)
Published on: 19/11/2025
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