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Senvelgo 15 mg/ml - Oral solution

Authorised

- Velagliflozin proline monohydrate

Product identification

Medicine name:
Senvelgo 15 mg/ml - Oral solution

Active substance:
Velagliflozin proline monohydrate

Target species:
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Velagliflozin proline monohydrate
20.05 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA10BK90

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:
Austria , Belgium , Cyprus , Czechia , France , Greece , Luxembourg , Poland , Slovakia , Spain

Package description:
Packaging:Bottle (HDPE), Package_size:1 bottle + 1 syringe, Content:30 ml
Packaging:Bottle (HDPE), Package_size:1 bottle + 1 syringe, Content:12 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - new active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
20/11/2023

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH
Boehringer Ingelheim Animal Health France

Responsible authority:
European Commission

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

Date of authorisation status change:
20/11/2023

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: 17/04/2026
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