

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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