Source URL: https://medicines.health.europa.eu/veterinary/en/600000107764

Zantoral 30mg/ml oral solution for dogs

Authorised

Ranitidine

Product identification

Medicine name:

Zantoral 30mg/ml oral solution for dogs

Zantoral 30 mg/ml solution buvable pour chiens

Zantoral 30 mg/ml orale oplossing voor honden

Zantoral 30 mg/ml Lösung zum Eingeben für Hunde

Active substance:

Ranitidine

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Ranitidine

30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Oral solution

Withdrawal period by route of administration:

Oral use:

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Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QA02BA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

High density polyethylene bottle 48 ml with high density polyethylene stoppers and a low density polyethylene plug. 3 ml polypropylene/silicone syringe, graduated every 0.1 ml to 3 ml.

High density polyethylene bottle 24 ml with high density polyethylene stoppers and a low density polyethylene plug. 3 ml polypropylene/silicone syringe, graduated every 0.1 ml to 3 ml.

High density polyethylene bottle 12 ml with high density polyethylene stoppers and a low density polyethylene plug. 3 ml polypropylene/silicone syringe, graduated every 0.1 ml to 3 ml.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Emdoka

Marketing authorisation date:

14/12/2023

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V662081

Date of authorisation status change:

14/12/2023

Reference member state:

Italy

Procedure number:

IT/V/0141/001

Concerned member states:

Austria Belgium Germany Luxembourg Netherlands

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

Documents

Summary of Product Characteristics
This document does not exist in this language (English). You can find it in another language below.
Labelling
This document does not exist in this language (English). You can find it in another language below.
Package Leaflet
This document does not exist in this language (English). You can find it in another language below.
Combined File of all Documents