Zantoral 30mg/ml oral solution for dogs

Authorised

- Ranitidine

Product identification

Medicine name:
Zantoral 30mg/ml oral solution for dogs

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:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA02BA02

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

Authorisation status:
Valid

Authorised in:
Italy

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:
20/09/2024

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Ministry Of Health

Authorisation number:
105328

Date of authorisation status change:
20/09/2024

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

Combined File of all Documents

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