File downloaded on 2026-11-24
Source URL: https://medicines.health.europa.eu/veterinary/en/600000085378

ZANTADINE, 30 mg/ml, soluzione orale per cani

Authorised

- Ranitidine hydrochloride

Product identification

Medicine name:
ZANTADINE, 30 mg/ml, soluzione orale per cani

Active substance:
Ranitidine hydrochloride

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Ranitidine hydrochloride
30.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA02BA02

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

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Ceva Salute Animale S.p.A.

Marketing authorisation date:
17/04/2009

Manufacturing sites for batch release:
Vetem S.p.A.
Ceva Sante Animale

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
17/04/2014

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.