

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

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

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