KESIUM 400 MG / 100 MG CHEWABLE TABLETS FOR DOGS

Suspended

This information is not available for this product.

Product identification

Medicine name:
KESIUM 400 MG / 100 MG CHEWABLE TABLETS FOR DOGS
KESIUM 400 mg / 100 mg COMPRIMIDOS MASTICABLES PARA PERROS

Active substance:
This information is not available for this product.

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
This information is not available for this product.

Pharmaceutical form:
Chewable tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Suspended

Authorised in:
Spain

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Box of 80 blisters of 6 chewable breakable tablets
Box of 3 blisters of 4 chewable breakable tablets
Box of 6 blisters of 4 chewable breakable tablets
Box of 9 blisters of 4 chewable breakable tablets
Box of 12 blisters of 4 chewable breakable tablets
Box of 15 blisters of 4 chewable breakable tablets
Box of 18 blisters of 4 chewable breakable tablets
Box of 21 blisters of 4 chewable breakable tablets
Box of 24 blisters of 4 chewable breakable tablets
Box of 60 blisters of 4 chewable breakable tablets

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:
Ceva Salud Animal S.A.

Marketing authorisation date:
16/01/2012

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
2430 ESP

Date of authorisation status change:
25/12/2021

Reference member state:
France

Procedure number:
FR/V/0225/004

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Ireland Italy Luxembourg Netherlands Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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.

Package Leaflet

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.
