EUPENCLAV 500 COMPRIMIDOS

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
EUPENCLAV 500 COMPRIMIDOS

Active substance:
Potassium clavulanate

Amoxicillin trihydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium clavulanate
119.07 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate
459.11 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

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:
Fatro S.p.A.

Marketing authorisation date:
21/12/2007

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
1822 ESP

Date of authorisation status change:
21/12/2007

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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