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AMOXIVAL VET 200 mg tablets for dogs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
AMOXIVAL VET 200 mg tablets for dogs

Active substance:
Amoxicillin trihydrate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate
229.60 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Finland

Available in:
Finland

Package description:
(ID1) 10 Tablet: Box (Cardboard) with 1 Blister (PolyVinyl Chloride; Aluminium) with 10 Tablet
(ID2) 200 Tablet: Box (Cardboard) with 20 Blister (PolyVinyl Chloride; Aluminium) each with 10 Tablet
(ID3) 20 Tablet: Box (Cardboard) with 2 Blister (PolyVinyl Chloride; Aluminium) each with 10 Tablet

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 Sante Animale

Marketing authorisation date:
10/11/2008

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Finnish Medicines Agency

Authorisation number:
23198

Date of authorisation status change:
10/11/2008

Reference member state:
Germany

Procedure number:
DE/V/0164/001

Concerned member states:
Belgium Denmark Finland Sweden

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

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

Combined File of all Documents