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Amoxiclav 200/50 mg tabletta kutyák és macskák részére A.U.V.

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
Amoxiclav 200/50 mg tabletta kutyák és macskák részére A.U.V.

Active substance:
Amoxicillin trihydrate

Potassium clavulanate

Target species:
Dog

Route of administration:
Oral use

Product details

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

Potassium clavulanate

50.00 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:
Hungary

Package description:
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian
Available only in Hungarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Medicus Partner Kft.

Marketing authorisation date:
12/10/2010

Manufacturing sites for batch release:
PenCef Pharma GmbH

Responsible authority:
Directorate Of Veterinary Medicinal Products

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

Date of authorisation status change:
12/10/2010

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.