Clavubactin 250/62.5 mg tablet

Authorised

- Clavulanic acid
- Amoxicillin

Product identification

Medicine name:
Clavubactin 250/62.5 mg tablet
CLAVUBACTIN 250/62,5 mg, tabletės šunims

Active substance:
Clavulanic acid

Amoxicillin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Clavulanic acid
62.50 milligram(s) / 1.00 Tablet

Amoxicillin
250.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

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:

Valid

Authorised in:

Lithuania

Package description:

Cardboard box containing 10 separate cardboard boxes, each containing 5 Aluminium/aluminium strips with 2 tablets each.
Aluminium/aluminium strips with 4 tablets, in a cardboard box containing 50 strips.
Aluminium/aluminium strips with 4 tablets, in a cardboard box containing 5 strips.
Aluminium/aluminium strips with 4 tablets, in a cardboard box containing 25 strips.
Aluminium/aluminium strips with 2 tablets, in a cardboard box containing 5 strips.
Aluminium/aluminium strips with 10 tablets, in a cardboard box containing 25 strips.
Aluminium/aluminium strips with 10 tablets, in a cardboard box containing 10 strips.
Aluminium/aluminium strips with 10 tablets, in a cardboard box containing 1 strip.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Le Vet. B.V.

Marketing authorisation date:
19/03/2009

Manufacturing sites for batch release:
Lelypharma B.V.
Pharmachem Consulting ApS

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/09/1841/001-006

Date of authorisation status change:
28/05/2014

Reference member state:
Netherlands

Procedure number:
NL/V/0110/002

Concerned member states:
Austria Belgium Denmark Finland Greece Lithuania Luxembourg Portugal

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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