Clavaseptin 50 mg Palatable Tablets for Dogs and Cats

Authorised

- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:
Clavaseptin 50 mg Palatable Tablets for Dogs and Cats

Active substance:
Amoxicillin trihydrate
Potassium clavulanate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

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

Potassium clavulanate
11.91 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:
United Kingdom (Northern Ireland)

Available in:
United Kingdom (Northern Ireland)

Package description:
Cardboard box of 1 blister of 10 tablets
Cardboard box of 100 blisters of 10 tablets
Cardboard box of 75 blisters of 10 tablets
Cardboard box of 60 blisters of 10 tablets
Cardboard box of 50 blisters of 10 tablets
Cardboard box of 40 blisters of 10 tablets
Cardboard box of 30 blisters of 10 tablets
Cardboard box of 25 blisters of 10 tablets
Cardboard box of 20 blisters of 10 tablets
Cardboard box of 15 blisters of 10 tablets
Cardboard box of 12 blisters of 10 tablets
Cardboard box of 10 blisters of 10 tablets
Cardboard box of 5 blisters of 10 tablets
Cardboard box of 2 blisters of 10 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Vetoquinol S.A.

Marketing authorisation date:
18/06/2004

Manufacturing sites for batch release:
Vetoquinol S.A.

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 06462/3001

Date of authorisation status change:
4/12/2024

Reference member state:
France

Procedure number:
FR/V/0407/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

eu-puar-frv0407001-mr-rpe869-en.pdf