Clavusan 500 mg + 125 mg tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Clavusan 500 mg + 125 mg tablets for dogs

Active substance:
Potassium clavulanate

Amoxicillin trihydrate

Target species:
Dog

Route of administration:
Oral use

Product details

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

Amoxicillin trihydrate

574.00 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:
Germany

Package description:
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes:Cardboard box of 10 tablets.
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes:Cardboard box of 30 tablets.
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes:Cardboard box of 50 tablets.
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes:Cardboard box of 100 tablets.
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes:Cardboard box of 250 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
22/05/2023

Manufacturing sites for batch release:
Lelypharma B.V.
Alfasan Nederland B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7008812.00.00

Date of authorisation status change:
22/05/2023

Reference member state:
Ireland

Procedure number:
IE/V/0778/003

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway 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

English (PDF)

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