Bioclanic 500 mg + 125 mg flavoured tablets for dogs

Authorised

- Amoxicillin
- Clavulanic acid

Product identification

Medicine name:
Bioclanic 500 mg + 125 mg flavoured tablets for dogs

Active substance:
Amoxicillin

Clavulanic acid

Target species:
Dog

Route of administration:
Oral use

Product details

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

Clavulanic acid

125.00 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:
Italy

Available in:
Italy

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 20 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 40 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 150 tablets.
oPA/Alu/PVC - PVC/Alu heat sealed blister containing 10 tablets each. Package sizes: Cardboard box of 200 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:
Axience

Marketing authorisation date:
8/10/2024

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
Ministry Of Health

Authorisation number:
105779

Date of authorisation status change:
8/10/2024

Reference member state:
Ireland

Procedure number:
IE/V/0788/003

Concerned member states:
Austria Belgium Cyprus Greece Hungary Italy Netherlands Poland Portugal Romania Spain

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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Summary of Product Characteristics

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