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Noroclav 500 mg Palatable Tablets for Dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Noroclav 500 mg Palatable 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
119.13 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate

459.22 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:
Portugal

Available in:
Portugal

Package description:
Aluminium/aluminium blister strips, each containing 5 tablets. Cartons of 100 tablets.
Aluminium/aluminium blister strips, each containing 5 tablets. Cartons of 25 tablets.
Aluminium/aluminium blister strips, each containing 5 tablets. Cartons of 20 tablets.
Aluminium/aluminium blister strips, each containing 5 tablets. Cartons of 10 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
20/08/2007

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
027/01/07RFVPT

Date of authorisation status change:
11/02/2026

Reference member state:
Ireland

Procedure number:
IE/V/0547/001

Concerned member states:
Austria Belgium Denmark France Luxembourg Netherlands Norway Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)

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