Noroclav 250 mg Tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Noroclav 250 mg Tablets for dogs

Active substance:
Potassium clavulanate

Amoxicillin trihydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

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

Amoxicillin trihydrate
229.61 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:
France

Package description:
The product is supplied in high-density polyethylene tubs with a polyethylene screw cap lid containing 100 tablets.
The product is supplied in high-density polyethylene tubs with a polyethylene screw cap lid containing 250 tablets.
The product is presented in packs of 2 blister strips (aluminium-aluminium) each containing 5 tablets per strip.
The product is presented in packs of 4 blister strips (aluminium-aluminium) each containing 5 tablets per strip.
The product is presented in packs of 10 blister strips (aluminium-aluminium) each containing 5 tablets per strip.
The product is presented in packs of 20 strips (aluminium-aluminium) each containing 5 tablets per strip.
The product is also presented in packs of 50 blister strips (aluminium-aluminium) each containing 5 tablets per strip.

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
4/06/2004

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/2868104 9/2004

Date of authorisation status change:
4/06/2009

Reference member state:
Ireland

Procedure number:
IE/V/0546/002

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

English (PDF)

Published on: 13/04/2025
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Combined File of all Documents

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.