Noroclav 50 mg Tablets for Dogs and Cats

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Noroclav 50 mg Tablets for Dogs and Cats
Noroclav Vet. 40+10 mg tabletter

Active substance:
Potassium clavulanate
Amoxicillin trihydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium clavulanate
11.91 milligram(s) / 1.00 Tablet
Amoxicillin trihydrate

45.92 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:
Oral use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
The product is supplied in high-density polyethylene tubs with a polypropylene screw cap lid containing 100 tablets.
The product is presented in packs containing 50 blister strips (aluminium-aluminium) each containing 10 tablets per strip
The product is presented in packs containing 10 blister strips (aluminium-aluminium) each containing 10 tablets per strip
The product is presented in packs containing 2 blister strips (aluminium-aluminium) each containing 10 tablets per strip
The product is supplied in high-density polyethylene tubs with a polyethylene screw cap lid containing 500 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Marketing authorisation holder:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
30/04/2004

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

Responsible authority:
Danish Medicines Agency

Authorisation number:
35924

Date of authorisation status change:
30/04/2004

Reference member state:
Ireland

Procedure number:
IE/V/0546/001

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

English (PDF)
Published on: 2/03/2025
Download

Combined File of all Documents
