Noroclav 50 mg Tablets for Dogs and Cats

- 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

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 <u>www.adrreports.eu/vet</u>

Documents

Summary of Product Characteristics

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

Combined File of all Documents

Source URL: https://medicines.health.europa.eu/veterinary/600000051001