Noroclav 250 mg Tablets for dogs

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Noroclav 250 mg Tablets for dogs
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

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:
Ireland

Available in:
Ireland

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:
28/05/2004

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

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/072/002

Date of authorisation status change:
28/05/2004

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

Combined File of all Documents
