Enrotab

Authorised

- Enrofloxacin

Product identification

Medicine name:
Enrotab

Active substance:
Enrofloxacin

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Enrofloxacin
15.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Box with 2 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (20 tablets)
Box with 15 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (150 tablets)
Box with 10 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (100 tablets)
Box with 6 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (60 tablets)
Box with 5 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (50 tablets)
Box with 3 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (30 tablets)
Box with 1 Alu-PVC PE PVDC blister or Alu-PVC PVDC blister with 10 tablets (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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

15/09/2010

Manufacturing sites for batch release:
Artesan Pharma GmbH & Co. KG

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 104599

Date of authorisation status change:
24/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0136/001

Concerned member states:
Hungary Spain

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

Documents

Combined File of all Documents

English (PDF)
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