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Enrotab

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrotab

Active substance:

Enrofloxacin

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Enrofloxacin
150.00 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:

Hungary

Package description:

Box with 3 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (30 tablets)

Box with 5 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (50 tablets)

Box with 1 Alu-PVC PE PVDC blister or Alu-PVC PVDC blister with 10 tablets (10 tablets)

Box with 15 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (150 tablets)

Box with 2 Alu-PVC PE PVDC blisters or Alu-PVC PVDC blisters with 10 tablets (20 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)

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:

12/10/2010

Manufacturing sites for batch release:

Artesan Pharma GmbH & Co. KG

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

2822/X/10 MgSzH ÁTI

Date of authorisation status change:

12/10/2010

Reference member state:

Netherlands

Procedure number:

NL/V/0136/003

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