XEDEN VET 50 MG TABLET FOR DOGS

Authorised

- Enrofloxacin

Product identification

Medicine name:
XEDEN VET 50 MG TABLET FOR DOGS
Xeden 50 mg tablet voor honden

Active substance:
Enrofloxacin

Target species:
Dog

Route of administration:
Oral use

Product details

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

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

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:
Cardboard box with 10 blisters of 10 tablets
Cardboard box with 1 blister of 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:
CEVA Sante Animale B.V.

Marketing authorisation date:
22/07/2008

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Medicines Evaluation Board

Authorisation number:

REG NL 100584

Date of authorisation status change:
26/04/2022

Reference member state:
France

Procedure number:
FR/V/0186/002

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Italy Luxembourg Netherlands Norway Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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