XEDEN 200 MG TABLET FOR DOGS

Authorised

- Enrofloxacin

Product identification

Medicine name:
XEDEN 200 MG TABLET FOR DOGS
Xeden 200 mg tablet voor honden

Active substance:
Enrofloxacin

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Enrofloxacin
200.00 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 20 blisters of 6 tablets
Cardboard box with 2 blisters of 6 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:
26/08/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Medicines Evaluation Board

Authorisation number:

REG NL 105787

Date of authorisation status change:
26/04/2022

Reference member state:
France

Procedure number:
FR/V/0186/004

Concerned member states:
Austria Belgium Czechia Denmark Finland Germany Greece Hungary Italy Luxembourg Netherlands Portugal Romania Spain

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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