

XEDEN 200 MG TABLET FOR DOGS

Authorised

- Enrofloxacin

Product identification

Medicine name:

XEDEN 200 MG TABLET FOR DOGS

Xeden 200 mg tabletta kutyák részére A.U.V.

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:

Hungary

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-Phylaxia Zrt.

Marketing authorisation date:

22/07/2010

Manufacturing sites for batch release:

Ceva Sante Animale

Responsible authority:

National Food Chain Safety Office

Authorisation number:

2770/X/10 MgSzH ÁTI

Date of authorisation status change:

22/07/2010

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000043503>