

Enrocare 25 mg/ml Solution for Injection for Dogs, Cats, Rabbits, Rodents, Reptiles and Ornamental birds

Authorised

- Enrofloxacin

Product identification

Medicine name:

Enrocare 25 mg/ml Solution for Injection for Dogs, Cats, Rabbits, Rodents, Reptiles and Ornamental birds

Active substance:

Enrofloxacin

Target species:

Reptile

Ornamental bird

Dog

Cat

Rabbit

Chinchilla

Gerbil

Guinea pig

Hamster

Mouse

Rat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Enrofloxacin

25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Rabbit

- Meat and offal. 6 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal. 100 ml vial sold in packs containing 12 x 100 ml cartons.

100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.100 ml vials sold in pack containing 6 x 100 ml cartons.

100 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.100 ml vial sold in packs containing 1 x 100 ml cartons.

50 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.50 ml vial sold in packs containing 12 x 50 ml cartons.

50 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.50 ml vial sold in packs containing 6 x 50 ml cartons.

50 ml Amber Type I multidose glass vial with a grey bromobutyl rubber stopper and aluminium overseal.50 ml vial sold in packs containing 1 x 50 ml cartons.

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:

Emdoka

Marketing authorisation date:

26/03/2010

Manufacturing sites for batch release:

Produlab Pharma B.V.

Responsible authority:

Ministry Of Health And Social Security

Authorisation number:

V 675/10/01/1025

Date of authorisation status change:

22/12/2014

Reference member state:

Ireland

Procedure number:

IE/V/0455/001

Concerned member states:

France Luxembourg Netherlands United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents