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:
Netherlands

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:
4/02/2010

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 104352

Date of authorisation status change:
27/01/2022

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

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics