KARIFLOX 25 mg/ml oral solution for calves

Authorised

- Enrofloxacin

Product identification

Medicine name:
KARIFLOX 25 mg/ml oral solution for calves

Active substance:
Enrofloxacin

Target species:
Cattle

Route of administration:
Oral use

Product details

Active substance and strength:
Enrofloxacin
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Cattle**
  - Meat and offal. 11 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Container Material:High density polyethylene bottlesContainer Closure:Polyethylene screw capContainer Colour:WhiteContainer Volume: 5 litreDosing Device:For all containers a 20 ml measuring device of polypropylene is included.
Container Material:High density polyethylene bottlesContainer Closure:Polyethylene screw capContainer Colour:WhiteContainer Volume: 1 litreDosing Device:For all containers a 20 ml measuring device of polypropylene is included.
Container Material:High density polyethylene bottlesContainer Closure:Polyethylene screw capContainer Colour:WhiteContainer Volume: 500 mlDosing Device:For all containers a 20 ml measuring device of polypropylene is included.
Container Material:High density polyethylene bottlesContainer Closure:Polyethylene screw capContainer Colour:WhiteContainer Volume: 250 mlDosing Device:For all containers a 20 ml measuring device of polypropylene is included.

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:
Laboratorios Karizoo S.A.

Marketing authorisation date:
10/07/2009

Manufacturing sites for batch release:
Laboratorios Karizoo S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10786/004/001

Date of authorisation status change:
10/07/2009

Reference member state:
Ireland

Procedure number:
IE/V/0217/002

Concerned member states:
France Germany Netherlands Spain

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

Documents

Summary of Product Characteristics