

# KARIFLOX 25 mg/ml oral solution for calves

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

KARIFLOX 25 mg/ml oral solution for calves

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**Active substance:**

Enrofloxacin

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**Target species:**

Cattle

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Enrofloxacin

25.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oral solution

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**Withdrawal period by route of administration:****Oral use:**

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**Cattle**

- Meat and offal. 11 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ01MA90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Karizoo S.A.

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**Marketing authorisation date:**

28/04/2009

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**Manufacturing sites for batch release:**

Laboratorios Karizoo S.A.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/7879896 4/2009

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**Date of authorisation status change:**

28/04/2014

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0217/002

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**Concerned member states:**

France Germany Netherlands Spain

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 3/05/2024

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