ADVOCIN 25 mg/ml SOLUCION INYECTABLE

Authorised

- Danofloxacin mesilate

Product identification

Medicine name:
ADVOCIN 25 mg/ml SOLUCION INYECTABLE

Active substance:
Danofloxacin mesilate

Target species:
Cattle
Pig
Cattle (dairy cow)

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Danofloxacin mesilate
31.73 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. no withdrawal period — Carne: 5 días desde la última inyección
- **Pig**
  - Meat and offal. 3 day
- **Cattle (dairy cow)**
  - Meat and offal. no withdrawal period — Carne: 5 días desde la última inyección
- **Cattle**
  - Milk. no withdrawal period — Leche: 48 horas después de la ultima inyección
- **Cattle (dairy cow)**
  - Milk. no withdrawal period — Leche: 48 horas después de la ultima inyección

Intravenous use:

- **Cattle**
  - Meat and offal. no withdrawal period — Carne: 5 días desde la última inyección
- **Cattle (dairy cow)**
  - Meat and offal. no withdrawal period — Carne: 5 días desde la última inyección
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Cattle

- Milk. no withdrawal period	Leche: 48 horas después de la ultima inyección

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Cattle (dairy cow)

- Milk. no withdrawal period	Leche: 48 horas después de la ultima inyección

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA92

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Spain S.L.

Marketing authorisation date:

23/02/1999

Manufacturing sites for batch release:
Fareva Amboise
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Spanish Agency For Medicines And Medical Devices

Authorisation number:
1243 ESP

Date of authorisation status change:
23/02/1999

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

Documents

Summary of Product Characteristics

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Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.