

FATROFLOX 100 mg/ml SOLUCION INYECTABLE

Authorised

- Enrofloxacin

Product identification

Medicine name:

FATROFLOX 100 mg/ml SOLUCION INYECTABLE

Active substance:

Enrofloxacin

Target species:

Cattle

Pig

Route of administration:

Subcutaneous use

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Enrofloxacin

100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

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

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Cattle

- Meat and offal. no withdrawal period Carne: Vía SC 12 días/ Vía IV 5 días

- Milk. no withdrawal period Leche: Vía SC 4 días/Vía IV 3 días

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Cattle

- Meat and offal. no withdrawal period Carne: Vía SC 12 días/ Vía IV 5 días

- Milk. no withdrawal period Leche: Vía SC 4 días/Vía IV 3 días

Intravenous use:

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Cattle

- Meat and offal. no withdrawal period Carne: Vía SC 12 días/ Vía IV 5 días

- Milk. no withdrawal period Leche: Vía SC 4 días/Vía IV 3 días

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Cattle

- Meat and offal. no withdrawal period Carne: Vía SC 12 días/ Vía IV 5 días

- Milk. no withdrawal period Leche: Vía SC 4 días/Vía IV 3 días

Intramuscular use:

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Pig

- Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro Iberica S.L.

Marketing authorisation date:

7/09/2005

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

1648 ESP

Date of authorisation status change:

7/09/2005

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

Documents

Labelling

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

Summary of Product Characteristics

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

Package Leaflet

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