

# XEDEN INIETTABILE, 50 mg/ml, soluzione iniettabile per bovini (vitelli), ovini, suini

Authorised

- Enrofloxacin

## Product identification

**Medicine name:**

XEDEN INIETTABILE, 50 mg/ml, soluzione iniettabile per bovini (vitelli), ovini, suini

**Active substance:**

Enrofloxacin

**Target species:**

Cattle (calf)

Sheep

Pig

**Route of administration:**

Intravenous use

Intramuscular use

Subcutaneous use

## Product details

**Active substance and strength:**

Enrofloxacin

50.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Solution for injection

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**Withdrawal period by route of administration:**

**Intravenous use:**

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**Cattle (calf)**

- Meat and offal. 5 day

Uso non autorizzato in animali che producono latte per consumo umano

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**Sheep**

- Meat and offal. 4 day

- Milk. 3 day

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**Pig**

- Meat and offal. 13 day

**Intramuscular use:**

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**Sheep**

- Meat and offal. 4 day

- Milk. 3 day

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**Pig**

- Meat and offal. 13 day

**Subcutaneous use:**

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**Cattle (calf)**

- Meat and offal. 12 day

Uso non autorizzato in animali che producono latte per consumo umano

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**Pig**

- Meat and offal. 13 day

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**Sheep**

- Meat and offal. 4 day
- Milk. 3 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:**

Italy

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**Package description:**

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Salute Animale S.p.A.

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**Marketing authorisation date:**

20/11/2012

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**Manufacturing sites for batch release:**

Vetem SPA

Ceva Sante Animale

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

20/11/2017

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[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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