

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)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Cattle (calf)

- Meat and offal. 5 day

Uso non autorizzato in animali che producono latte per consumo umano

-

Sheep

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

-

Pig

- Meat and offal. 13 day

Intramuscular use:

-

Sheep

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

-

Pig

- Meat and offal. 13 day

Subcutaneous use:

-

Cattle (calf)

- Meat and offal. 12 day

Uso non autorizzato in animali che producono latte per consumo umano

-

Pig

- Meat and offal. 13 day

-

Sheep

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01MA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in Italian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Ceva Salute Animale S.p.A.

Marketing authorisation date:

20/11/2012

Manufacturing sites for batch release:

Vetem SPA

Ceva Sante Animale

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

20/11/2017

To consult adverse reactions on veterinary medicinal products please go to

www.adrreports.eu/vet

Documents

Combined File of all Documents

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