XEDEN INIETTABILE, 100 mg/ml, soluzione iniettabile per bovini, ovini, suini

Authorised

- Enrofloxacin

Product identification

Medicine name:
XEDEN INIETTABILE, 100 mg/ml, soluzione iniettabile per bovini, ovini, suini

Active substance:
Enrofloxacin

Target species:
Sheep
Pig
Cattle

Route of administration:
Intravenous use
Intramuscular use
Subcutaneous 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:

Intravenous use:

- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 13 day
- **Cattle**
  - Meat and offal. 5 day
  - Milk. 3 day

Intramuscular use:

- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 13 day

Subcutaneous use:

- **Pig**
  - Meat and offal. 13 day
- **Sheep**
  - Meat and offal. 4 day
  - Milk. 3 day
- **Cattle**
  - Meat and offal. 12 day
  - Milk. 4 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
Available only in Italian
Available only in Italian
Available only in Italian
Available only in Italian
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 S.p.A.
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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