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

- Enrofloxacin

Authorised

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

Product information

Combined File of all Documents

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