ENGEMICINA D.D

Authorised

- Oxytetracycline

Product identification

Medicine name:
ENGEMICINA D.D

Active substance:
Oxytetracycline

Target species:
Cattle
Dog
Sheep
Horse (non food-producing)
Pig
Cat

Route of administration:
Intramuscular use
Intravenous use
Subcutaneous use

Product details

Active substance and strength:
Oxytetracycline
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Milk. 96 hour ore (basso dosaggio)
  - Meat and offal. 39 day giorni (basso dosaggio)
  - Meat and offal. 27 day giorni (alto dosaggio)
  - Milk. 120 hour ore (alto dosaggio)
- **Dog**
- **Sheep**
  - Milk. 120 hour ore (basso dosaggio)
  - Meat and offal. 16 day giorni (basso dosaggio)
  - Meat and offal. 18 day giorni (alto dosaggio)
  - Milk. 120 hour ore (alto dosaggio)
- **Horse (non food-producing)**
- **Pig**
  - Meat and offal. 9 day giorni (alto dosaggio)
  - Meat and offal. 13 day giorni (basso dosaggio)

Intravenous use:

- **Cattle**
  - Meat and offal. 39 day giorni (basso dosaggio)
  - Milk. 96 hour ore (basso dosaggio)
- **Sheep**
  - Meat and offal. 16 day giorni (basso dosaggio)
  - Milk. 120 hour ore (basso dosaggio)
- **Horse (non food-producing)**

Subcutaneous use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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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
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:
Intervet International B.V.

Marketing authorisation date:
15/07/1993

Manufacturing sites for batch release:
Intervet Productions S.r.l.
Intervet International GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
15/07/1993

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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