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ACTI-TETRA I

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
ACTI-TETRA I

Active substance:
Oxytetracycline hydrochloride

Target species:
Cattle
Pig
Cat
Horse
Sheep
Goat
Dog

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use
Intraperitoneal use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 14 day
- **Horse**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Goat**
  - Meat and offal. 14 day
  - Milk. 3 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 14 day
- **Horse**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Goat**
  - Meat and offal. 14 day
  - Milk. 3 day

Intravenous use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 14 day
- **Horse**

- Meat and offal. 14 day
- Milk. 3 day

- **Sheep**
  - Meat and offal. 14 day
  - Milk. 3 day

- **Goat**
  - Meat and offal. 14 day
  - Milk. 3 day

Intraperitoneal use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day

- **Pig**
  - Meat and offal. 14 day

- **Horse**
  - Meat and offal. 14 day
  - Milk. 3 day

- **Sheep**
  - Meat and offal. 14 day
  - Milk. 3 day

- **Goat**

- Meat and offal. 14 day
- Milk. 3 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Laboratoires Biove

Marketing authorisation date:
6/05/1988

Manufacturing sites for batch release:
Laboratoires Biove

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6090637 3/1988

Date of authorisation status change:
6/05/2013

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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