OXYTETRACYCLINE 5 % VETOQUINOL

Authorised

- Oxytetracycline dihydrate

Product identification

Medicine name:
OXYTETRACYCLINE 5 % VETOQUINOL

Active substance:
Oxytetracycline dihydrate

Target species:
Cattle
Pig
Cat
Horse
Sheep
Dog

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

Product details

Active substance and strength:
Oxytetracycline dihydrate
53.94 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
  - Milk. 3 day
- **Horse**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Sheep**
  - Meat and offal. 14 day
  - Milk. 3 day

Subcutaneous use:

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

- Milk. 3 day

- **Pig**
  - Meat and offal. 14 hour
  - Milk. 3 day

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

- **Sheep**
  - 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
  - Milk. 3 day

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

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

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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:
Vetoquinol S.A.

Marketing authorisation date:
22/06/1988

Manufacturing sites for batch release:
Vetoquinol S.A.

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

Authorisation number:
FR/V/6437675 6/1988

Date of authorisation status change:
22/06/2013

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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