

VET-OXY 5 % INJECTABLE

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:

VET-OXY 5 % INJECTABLE

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**

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

• **Pig**

- Meat and offal. 14 day

• **Cat**

• **Horse**

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

• **Sheep**

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

• **Goat**

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

• **Dog**

Subcutaneous use:

• **Cattle**

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

• **Pig**

- Meat and offal. 14 day

• **Cat**

• **Horse**

- Milk. 3 day

- Meat and offal. 14 day

• **Sheep**

- Milk. 3 day

- Meat and offal. 14 day

• **Goat**

- Milk. 3 day

- Meat and offal. 14 day

• **Dog**

Intravenous use:

• **Cattle**

- Milk. 3 day

- Meat and offal. 14 day

• **Pig**

- Meat and offal. 14 day

• **Cat**

• **Horse**

- Milk. 3 day

- Meat and offal. 14 day

• **Sheep**

- Milk. 3 day

- Meat and offal. 14 day

• **Goat**

- Milk. 3 day

- Meat and offal. 14 day

• **Dog**

Intraperitoneal use:

• **Cattle**

- Milk. 3 day

- Meat and offal. 14 day

• **Pig**

- Meat and offal. 14 day

- **Cat**

- **Horse**

- Milk. 3 day

- Meat and offal. 14 day

- **Sheep**

- Milk. 3 day

- Meat and offal. 14 day

- **Goat**

- Milk. 3 day

- Meat and offal. 14 day

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratoires Biove

Marketing authorisation date:

31/01/2017

Manufacturing sites for batch release:

Laboratoires Biové

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/7962294 1/2017

Date of authorisation status change:

9/02/2022

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

Documents

Summary of Product Characteristics

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