

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

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

•

Pig

- Meat and offal. 14 day

•

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

Subcutaneous use:

•

Cattle

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

•

Pig

- Meat and offal. 14 day

•

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

Intravenous use:

•

Cattle

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

•

Pig

- Meat and offal. 14 day

•

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

Intraperitoneal use:

•

Cattle

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

•

Pig

- Meat and offal. 14 day

•

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

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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