**Source URL:** https://medicines.health.europa.eu/veterinary/en/60000092008

# TERRAMICINA 100 mg/ml solução injetável para bovinos, ovinos, caprinos, suínos e equinos.

Authorised

• Oxytetracycline hydrochloride

# Product identification

#### **Medicine name:**

TERRAMICINA 100 mg/ml solução injetável para bovinos, ovinos, caprinos, suínos e equinos.

#### **Active substance:**

Oxytetracycline hydrochloride

# **Target species:**

Cattle

Pig

Sheep

Horse

Goat

#### Route of administration:

Intramuscular use

Intravenous use

# **Product details**

# **Active substance and strength:**

Oxytetracycline hydrochloride 100.00 milligram(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

Solution for injection

# Withdrawal period by route of administration: Intramuscular use:

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#### **Cattle**

- Meat and offal. 34 day
- Milk. 120 hour

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# Pig

- Meat and offal. 24 day

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# **Sheep**

- Meat and offal. 15 day
- Milk. 120 hour

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#### Horse

- Meat and offal. 999999 hour

Não administrar a equinos destinados a consumo humano. Não administrar a éguas cujo leite seja destinado a consumo humano

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#### Goat

- Meat and offal. 15 day
- Milk. 120 hour

#### Intravenous use:

•

#### **Cattle**

- Meat and offal. 34 day
- Milk. 120 hour

•

# Pig

- Meat and offal. 24 day

•

# **Sheep**

- Meat and offal. 15 day
- Milk. 120 hour

•

#### Goat

- Meat and offal. 15 day
- Milk. 120 hour

# Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01AA06

# Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Portugal

# Package description:

Available only in **Portuguese** 

Available only in **Portuguese** 

# 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:

Zoetis Portugal Lda.

# Marketing authorisation date:

24/05/1976

# Manufacturing sites for batch release:

Fareva Amboise

## **Responsible authority:**

Directorate General For Food And Veterinary

#### **Authorisation number:**

1047/01/16NFVPT

# **Date of authorisation status change:**

1/02/2017

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

# **Documents**

Combined File of all Documents

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