# Oxyvet 50 mg/ml Roztwór do wstrzykiwań

Authorised

- Oxytetracycline hydrochloride

## Product identification

**Medicine name:**
Oxyvet 50 mg/ml Roztwór do wstrzykiwań

**Active substance:**
Oxytetracycline hydrochloride

**Target species:**
Sheep
Horse
Pig
Cattle

**Route of administration:**
Intramuscular use
Intravenous use

## Product details

**Active substance and strength:**
Oxytetracycline hydrochloride

**Pharmaceutical form:**
Solution for injection

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**Withdrawal period by route of administration:**
**Intramuscular use:**

- **Sheep**
  - Meat and offal. 21 day

- **Horse**
  - All relevant tissues. no withdrawal period

Not to be used in horses intended for human consumption.  Treated horses may never be slaughtered for human consumption.  The horse must have been declared as not intended for human consumption under national horse passport legislation.

- **Pig**
  - Meat and offal. 21 day

- **Cattle**
  - Meat and offal. 24 day

**Intravenous use:**

- **Sheep**
  - Meat and offal. 21 day

- **Horse**
  - All relevant tissues. no withdrawal period

Not to be used in horses intended for human consumption.  Treated horses may never be slaughtered for human consumption.  The horse must have been declared as not intended for human consumption under national horse passport legislation.

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**Pig**

- Meat and offal. 21 day

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**Cattle**

- Meat and offal. 24 day

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01AA06

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

**Authorisation status:**
Valid

**Authorised in:**
Poland

**Package description:**
Available only in Polish
Available only in Polish

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Well-established use application (Article 13a of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Biofaktor Sp. z o.o.

**Marketing authorisation date:**
16/11/1976

**Manufacturing sites for batch release:**

Biowet Pulawy Sp. z o.o.

**Responsible authority:**
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

**Authorisation number:**
0805

**Date of authorisation status change:**
16/11/1976

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

## Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.