

# Oxy-kel 20% Long Acting 200 mg/ml Oplossing voor injectie

Authorised

- Oxytetracycline hydrochloride

## Product identification

**Medicine name:**

Oxy-kel 20% Long Acting 200 mg/ml Oplossing voor injectie

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**Active substance:**

Oxytetracycline hydrochloride

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**Target species:**

Cattle

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**Route of administration:**

Intramuscular use

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## Product details

**Active substance and strength:**

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

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**Pharmaceutical form:**

Solution for injection

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

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**Cattle**

- Meat and offal. 53 day
- Milk. no withdrawal period

Do not use in animals producing milk for human consumption

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

QJ01AA06

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Belgium

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**Available in:**

Belgium

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**Package description:**

Oxy-kel 20% Long Acting 1 Vial with 250 ml Solution for injection

Oxy-kel 20% Long Acting 1 Vial with 100 ml Solution for injection

Oxy-kel 20% Long Acting 1 Vial with 50 ml Solution for injection

Oxy-kel 20% Long Acting 1 Vial with 30 ml Solution for injection

Oxy-kel 20% Long Acting 25 Vials with 30 ml Solution for injection

Oxy-kel 20% Long Acting 12 Vials with 50 ml Solution for injection

Oxy-kel 20% Long Acting 12 Vials with 100 ml Solution for injection

Oxy-kel 20% Long Acting 12 Vials with 250 ml Solution for injection

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

KELA Kempisch Laboratorium Kela Laboratoria

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**Marketing authorisation date:**

11/12/1991

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**Manufacturing sites for batch release:**

KELA Kempisch Laboratorium Kela Laboratoria

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V156213

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**Date of authorisation status change:**

5/12/2023

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.