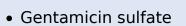
Gentamycin 85 mg/ ml šķīdums injekcijām liellopiem, cūkām, zirgiem, suņiem un kaķiem



# Product identification

#### Medicine name:

Gentamycin 85 mg/ ml šķīdums injekcijām liellopiem, cūkām, zirgiem, suņiem un kaķiem

Authorised

#### **Active substance:**

Gentamicin sulfate

#### **Target species:**

Horse (non food-producing) Cattle (calf) Pig (piglet) Pig (weaned piglet) Dog Cat Cattle

#### Route of administration:

Intravenous use Intramuscular use Subcutaneous use

## Product details

#### Active substance and strength:

Gentamicin sulfate 85.00 milligram(s) / 1.00 millilitre(s)

#### **Pharmaceutical form:**

Solution for injection

#### Withdrawal period by route of administration:

#### Intravenous use:

- Horse (non food-producing)
- . Cattle (calf)
  - Meat and offal. 192 day
- Pig (piglet)
  - Meat and offal. 146 day
- Pig (weaned piglet)
  - Meat and offal. 146 day
- . Dog
- . Cat
- Cattle
  - Meat and offal. 214 day
  - Milk. 7 day

#### Intramuscular use:

- Cattle
  - Meat and offal. 214 day
  - Milk. 7 day
- Cattle (calf)
  - Meat and offal. 192 day
- . Pig (piglet)
  - Meat and offal. 146 day
- Pig (weaned piglet)
  - Meat and offal. 146 day

- . Cat
- . Dog

### Subcutaneous use:

- Dog
- . Cat

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01GB03

### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

### Authorised in:

Latvia

## Available in:

Latvia

## Package description:

Available only in <u>Latvian</u> Available only in <u>Latvian</u> Available only in <u>Latvian</u>

# Additional information

Entitlement type: Marketing Authorisation

Legal basis of product authorisation: Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder: Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

1/03/1995

#### Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

**Responsible authority:** Food And Veterinary Service

Authorisation number: V/NRP/95/0110

Date of authorisation status change:

1/03/1995

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

## 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.

**Source URL:** *https://medicines.health.europa.eu/veterinary/60000014308*