

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

Authorised

- Gentamicin sulfate

Product identification

Medicine name:

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

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 Latvian

Available only in Latvian

Available only in Latvian

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
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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