

Genta 50 mg/ml solution for injection

Authorised

- Gentamicin sulfate

Product identification

Medicine name:

Genta 50 mg/ml solution for injection

Active substance:

Gentamicin sulfate

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Gentamicin sulfate

54.08 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. 214 day

- Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01GB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Type II amber glass vial with bromobutyl rubber stopper, sealed with an aluminium cap, containing 100 ml of a sterile colourless to pale yellow aqueous solution. The vials are packed in a polystyrene box, 12 vials of 100 ml per box.

Type II amber glass vial with bromobutyl rubber stopper, sealed with an aluminium cap, containing 100 ml of a sterile colourless to pale yellow aqueous solution.

Type II amber glass vial with bromobutyl rubber stopper, sealed with an aluminium cap, containing 50 ml of a sterile colourless to pale yellow aqueous solution.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:

KELA Kempisch Laboratorium Kela Laboratoria

Marketing authorisation date:

1/10/2003

Manufacturing sites for batch release:

KELA Kempisch Laboratorium Kela Laboratoria

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10981/011/001

Date of authorisation status change:

1/10/2003

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

Documents

Summary of Product Characteristics