

Gentamicin 50 mg/ml, ενέσιμο διάλυμα για βοοειδή, σκύλους και γάτες

Authorised

- Gentamicin sulfate

Product identification

Medicine name:

Gentamicin 50 mg/ml, ενέσιμο διάλυμα για βοοειδή, σκύλους και γάτες

Active substance:

Gentamicin sulfate

Target species:

Cattle

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Intravenous 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:**Intramuscular use:**

-

Cattle

- Milk. 6 day
- Meat and offal. 95 day

-

Dog

-

Cat**Subcutaneous use:**

-

Cattle

- Meat and offal. 95 day
- Milk. 6 day

-

Dog

-

Cat**Intravenous use:**

-

Cattle

- Meat and offal. 95 day
- Milk. 6 day

-

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:

Cyprus

Package description:

Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Bremer Pharma GmbH

Marketing authorisation date:

12/10/1998

Manufacturing sites for batch release:

Bremer Pharma GmbH

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

11959

Date of authorisation status change:

29/07/2010

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

Documents

Combined File of all Documents

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