Aagent, 50mg/ml, Injekční roztok

Authorised

- Gentamicin

Product identification

Medicine name:
Aagent, 50mg/ml, Injekční roztok

Active substance:
Gentamicin

Target species:
Cattle (calf)
Horse
Pig (piglet)

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Gentamicin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intravenous use:

- **Cattle (calf)**
  - Meat and offal. no withdrawal period

Z důvodu akumulace gentamicinu v játrech, ledvinách a v místě injekčního podání, musí být zamezeno jakémukoli opakování léčby v průběhu ochranné lhůty.,

- **Horse**
  - Meat and offal. no withdrawal period

Nepoužívat u koní, jejichž maso a mléko je určeno pro lidskou spotřebu.,

Intramuscular use:

- **Pig (piglet)**
  - Meat and offal. no withdrawal period

Z důvodu akumulace gentamicinu v játrech, ledvinách a v místě injekčního podání, musí být zamezeno jakémukoli opakování léčby v průběhu ochranné lhůty.,

- **Cattle (calf)**
  - Meat and offal. no withdrawal period

Z důvodu akumulace gentamicinu v játrech, ledvinách a v místě injekčního podání, musí být zamezeno jakémukoli opakování léčby v průběhu ochranné lhůty.,

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Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01GB03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
28/04/1993

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
96/370/93-C

Date of authorisation status change:
15/12/2017

To consult adverse reactions on veterinary medicinal products please go to 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.