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Aagent 50 mg/ml injekčný roztok pre hovädzí dobytok (teľatá), ošípané (ciciaky) a kone (neurčené na produkciu potravín)

Authorised

- Gentamicin

Product identification

Medicine name:
Aagent 50 mg/ml injekčný roztok pre hovädzí dobytok (teľatá), ošípané (ciciaky) a kone (neurčené na produkciu potravín)

Active substance:
Gentamicin

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

Route of administration:
Intramuscular use
Intravenous 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:
Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 103 day

- **Pig (piglet)**
  - Meat and offal. 66 day

Intravenous use:

- **Cattle (calf)**
  - Meat and offal. 103 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01GB03

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

Authorisation status:
Valid

Authorised in:
Slovakia

Available in:

Slovakia

Package description:
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
23/12/1994

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

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

Authorisation number:
96/806/94-S

Date of authorisation status change:
23/12/1994

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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