

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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