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AAGENT, 50 mg/ml ενέσιμο διάλυμα για μοσχάρια και χοιρίδια στον πρώτο μήνα ζωής

Authorised

- Gentamicin

Product identification

Medicine name:
AAGENT, 50 mg/ml ενέσιμο διάλυμα για μοσχάρια και χοιρίδια στον πρώτο μήνα ζωής

Active substance:
Gentamicin

Target species:
Cattle (calf)
Pig (piglet)

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

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

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:
Fatro S.p.A.

Marketing authorisation date:
9/06/1989

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

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
12314

Date of authorisation status change:
9/06/1989

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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