

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:

-

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

Available 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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