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Apravet 552 000 IU/g powder for use in drinking water/milk for pigs, calves, chickens and rabbits

Authorised

- Apramycin sulfate

Product identification

Medicine name:
Apravet 552 000 IU/g powder for use in drinking water/milk for pigs, calves, chickens and rabbits

Active substance:
Apramycin sulfate

Target species:
Pig (weaned piglet)
Rabbit
Cattle (pre-ruminant)
Chicken (broiler)

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:

Apramycin sulfate
1.00 milligram(s) / 1.00 milligram(s)

Pharmaceutical form:
Powder for use in drinking water/milk

Withdrawal period by route of administration:
In drinking water/milk use:

- **Pig (weaned piglet)**
  - Meat and offal. 0 day
- **Rabbit**
  - Meat and offal. 0 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 28 day
- **Chicken (broiler)**
  - Meat and offal. 0 day
  - Eggs. no withdrawal period

Eggs: Not for use in birds producing or intended to produce eggs for human consumption. Do not use within 4 weeks of the start of the laying period.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA92

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
1 Sachet with 1,812g
box containing 50 sachets of 1,812 g
box containing 25 sachets of 1,812 g
bottle containing 90,58 g
bag containing 1811,6 g

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
8/09/2018

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402305.00.00

Date of authorisation status change:
8/09/2018

Reference member state:
Spain

Procedure number:
ES/V/0252/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Netherlands Poland Portugal Romania Slovakia Slovenia United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents

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