File downloaded on 2026-11-23
Source URL: https://medicines.health.europa.eu/veterinary/en/600000093710

Paramore, 200 mg/ml, soluzione per uso orale per vitelli da latte, suini, polli, tacchini e conigli

Authorised

- Paromomycin sulfate

Product identification

Medicine name:
Paramore, 200 mg/ml, soluzione per uso orale per vitelli da latte, suini, polli, tacchini e conigli

Active substance:
Paromomycin sulfate

Target species:
Turkey
Chicken
Rabbit
Pig
Cattle (suckling calf)

Route of administration:
Oral use

Product details

Active substance and strength:
Paromomycin sulfate
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:
Oral use:

- **Turkey**
  - Meat and offal. 2 day

Uso non autorizzato in tacchini che producono uova per il consumo umano

- **Chicken**
  - Meat and offal. 2 day
  - Eggs. 0 day
- **Rabbit**
  - Meat and offal. 3 day
- **Pig**
  - Meat and offal. 44 day
- **Cattle (suckling calf)**
  - Meat and offal. 44 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA07AA06

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

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
11/04/2016

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
11/04/2016

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

Documents

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.