OXITETRACICLINA FP 900 mg/g pulbere pentru utilizare în apa de băut/înlocuitor de lapte

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
OXITETRACICLINA FP 900 mg/g pulbere pentru utilizare în apa de băut/înlocuitor de lapte

Active substance:
Oxytetracycline hydrochloride

Target species:
Pig
Cattle (suckling calf)
Horse (suckling foal)
Sheep (suckling lamb)
Goat (suckling kid)
Rabbit
Dog
Chicken (broiler)
Turkey
Duck
Goose
Chicken

Fish

Route of administration:
Buccal use

Product details

Active substance and strength:
Oxytetracycline hydrochloride
900.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:
Buccal use:

- **Pig**
  - Meat and offal. 7 day
- **Cattle (suckling calf)**
  - Meat and offal. 7 day
- **Horse (suckling foal)**
  - Meat and offal. 7 day
- **Sheep (suckling lamb)**
  - Meat and offal. 7 day
- **Goat (suckling kid)**
  - Meat and offal. 7 day
- **Rabbit**

- Meat and offal. 7 day

- **Chicken (broiler)**
  - Meat and offal. 7 day

- **Turkey**
  - Meat and offal. 7 day

- **Duck**
  - Meat and offal. 7 day

- **Goose**
  - Meat and offal. 7 day

- **Chicken**
  - Meat and offal. 7 day

- **Fish**
  - Meat. 90 day in SPC, nu sunt specificate ”Day degree”

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01AA06

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Available in:

Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Pasteur Filiala Filipesti S.A.

Marketing authorisation date:
11/06/2007

Manufacturing sites for batch release:
Pasteur Filiala Filipesti S.A.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
140004

Date of authorisation status change:
13/06/2024

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

Documents

Summary of Product Characteristics

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