

AMOXICILINA FP 600 mg/g, pulbere pentru utilizare în apa de băut/lapte pentru cai, bovine, oi, capre, porci, găini

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

AMOXICILINA FP 600 mg/g, pulbere pentru utilizare în apa de băut/lapte pentru cai, bovine, oi, capre, porci, găini

Active substance:

Amoxicillin trihydrate

Target species:

Horse

Cattle (newborn calf)

Sheep (lamb)

Goat (kid)

Pig

Chicken (broiler)

Route of administration:

In drinking water/milk use

Product details

Active substance and strength:

Amoxicillin trihydrate

60.00 gram(s) / 100.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water/milk

Withdrawal period by route of administration:

In drinking water/milk use:

-

Horse

- Meat and offal. 5 day

-

Cattle (newborn calf)

- Meat and offal. 5 day

-

Sheep (lamb)

- Meat and offal. 5 day

-

Goat (kid)

- Meat and offal. 5 day

-

Pig

- Meat and offal. 5 day

-

Chicken (broiler)

- Meat and offal. 5 day

Nu se utilizează la păsările care produc ouă pentru consum uman.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

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:

140081

Date of authorisation status change:

9/12/2024

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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