

VETAMPLIUS

Authorised

- Ampicillin
- Ampicillin
- Ampicillin
- Ampicillin

Product identification

Medicine name:

VETAMPLIUS

Active substance:

Ampicillin

Ampicillin

Ampicillin

Ampicillin

Target species:

Cattle

Dog

Sheep

Cat

Pig

Horse

Route of administration:

Intraperitoneal use

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Ampicillin

2.00 gram(s) / 1.00 Bottle

Ampicillin

4.00 gram(s) / 1.00 Bottle

Ampicillin

10.00 gram(s) / 1.00 Bottle

Ampicillin

100.00 gram(s) / 1.00 Bottle

Pharmaceutical form:

Powder and solvent for solution for injection

Withdrawal period by route of administration:

Intraperitoneal use:

-

Cattle

- Meat and offal. 3 day

- Milk. 5 day

-

Sheep

- Meat and offal. 9 day

- Milk. 5 day

-

Pig

- Meat and offal. 9 day

-

Horse

- Meat and offal. 9 day

Uso non autorizzato in equidi che producono latte per il consumo umano

Intramuscular use:

-

Cattle

- Meat and offal. 3 day
- Milk. 5 day

-

Sheep

- Meat and offal. 9 day
- Milk. 5 day

-

Pig

- Meat and offal. 9 day

-

Horse

- Meat and offal. 9 day

Uso non autorizzato in equidi che producono latte per il consumo umano

Intravenous use:

-

Cattle

- Meat and offal. 3 day
- Milk. 5 day

-

Sheep

- Meat and offal. 9 day
- Milk. 5 day

-

Pig

- Meat and offal. 9 day

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Horse

- Meat and offal. 9 day

Usò non autorizzato in equidi che producono latte per il consumo umano

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

9/12/1975

Manufacturing sites for batch release:

Fatro S.p.A.
Bioquim S.A.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

22/09/1986

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.