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
- **Dog**
- **Sheep**
  - Meat and offal. 9 day
  - Milk. 5 day
- **Cat**
- **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

- **Dog**
- **Sheep**
  - Meat and offal. 9 day
  - Milk. 5 day
- **Cat**
- **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
- **Dog**
- **Sheep**
  - Meat and offal. 9 day
  - Milk. 5 day
- **Cat**
- **Pig**
  - Meat and offal. 9 day
- **Horse**
  - Meat and offal. 9 day

Uso 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.
