BACOLAM 100 mg/ml + 250.000 UI/ml sospensione iniettabile per bovini, ovi-caprini, suini, equini e tacchini

Authorised

- Amoxicillin
- COLISTIN SULFATE

Product identification

Medicine name:
BACOLAM 100 mg/ml + 250.000 UI/ml sospensione iniettabile per bovini, ovi-caprini, suini, equini e tacchini

Active substance:
Amoxicillin
COLISTIN SULFATE

Target species:
Cattle
Goat
Sheep
Pig
Horse (non-food-producing)
Turkey

Route of administration:

Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Amoxicillin
100.00 milligram(s) / 1.00 millilitre(s)

COLISTIN SULFATE
250000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

- **Goat**
  - Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

- **Sheep**
  - Meat and offal. 28 day

Non somministrare in animali in lattazione il cui latte o derivati siano destinati al consumo umano

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Pig

- Meat and offal. 13 day

Subcutaneous use:

-

Turkey

- Meat and offal. 25 day

Uso non consentito in animali che producono uova destinati al consumo umano

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RA01

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

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:
14/10/1993

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

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

Date of authorisation status change:
25/10/2013

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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