Vetrimoxin LA 150 mg/ ml suspensija injekcijām liellopiem un cūkām

Authorised

- Amoxicillin

Product identification

Medicine name:
Vetrimoxin LA 150 mg/ ml suspensija injekcijām liellopiem un cūkām

Active substance:
Amoxicillin

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 14 day
  - Milk. 3 day
- **Pig**
  - Meat and offal. 16 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
16/12/2003

Manufacturing sites for batch release:
Vetem S.p.A.
Ceva Sante Animale

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/03/1616

Date of authorisation status change:
16/12/2003

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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