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Vetrimoxin L.A. 150 mg/ml suspension for injection for cattle and pigs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Vetrimoxin L.A. 150 mg/ml suspension for injection for cattle and pigs
Vetrimoxin vet. 150 mg/ml Injektionsvätska, suspension

Active substance:
Amoxicillin trihydrate

Target species:
Cattle
Pig

Route of administration:
Intramuscular use

Product details

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

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Milk. 3 day
  - Meat and offal. 18 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:
Sweden

Available in:
Sweden

Package description:
(ID4) 3000 millilitre(s): unspecified outer container with 12 Vial (Plastic) each with 250 millilitre(s)
(ID3) 250 millilitre(s): unspecified outer container with 1 Vial (Plastic) with 250 millilitre(s)
(ID2) 100 millilitre(s): unspecified outer container with 1 Vial (Plastic) with 100 millilitre(s)
(ID1) 1200 millilitre(s): unspecified outer container with 12 Vial (Plastic) each with 100 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
14/02/2013

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
48416

Date of authorisation status change:
14/02/2013

Reference member state:
Germany

Procedure number:
DE/V/0153/001

Concerned member states:
Austria Denmark Finland Iceland Ireland Netherlands Sweden
United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

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Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.