COBACTAN LA 7.5% w/v suspension for injection for cattle

Authorised

- Cefquinome sulfate

Product identification

Medicine name:
COBACTAN LA 7.5% w/v suspension for injection for cattle

Active substance:
Cefquinome sulfate

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Cefquinome sulfate
88.90 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Cattle**
  - Meat and offal. 13 day
  - Milk. no withdrawal period

Do not use in dairy cows producing milk for human consumption (during lactation or the dry period). Do not use within two months prior to first calving in heifers intended for the production of milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DE90

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

Authorisation status:
Valid

Authorised in:
Greece

Package description:
(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (glass) with 100 millilitre(s)
(ID2) 250 millilitre(s): unspecified outer container with 1 Vial (glass) with 250 millilitre(s)
(ID3) 50 millilitre(s): unspecified outer container with 1 Vial (glass) with 50 millilitre(s)

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:
Intervet International B.V.

Marketing authorisation date:
27/11/2006

Manufacturing sites for batch release:
Intervet International GmbH
Intervet Productions S.r.l.

Responsible authority:
National Organization For Medicines

Authorisation number:
31352/16-05-2012/K-0094206

Date of authorisation status change:
2/05/2017

Reference member state:
Germany

Procedure number:
DE/V/0145/001

Concerned member states:
Austria Belgium Cyprus Czechia Denmark France Greece Hungary Italy Luxembourg Netherlands Poland Portugal Slovakia Slovenia Spain

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents