Emdactilin 150 56.7 mg/ml Oplossing voor injectie

Authorised

- Lincomycin hydrochloride
- SPECTINOMYCIN HYDROCHLORIDE

Product identification

Medicine name:
Emdactilin 150 56.7 mg/ml Oplossing voor injectie
Emdactilin 150 56.7 mg/ml Solution injectable
Emdactilin 150 56.7 mg/ml Injektionslösung

Active substance:
Lincomycin hydrochloride
SPECTINOMYCIN HYDROCHLORIDE

Target species:
Pig
Cattle (calf)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Lincomycin hydrochloride
56.70 milligram(s) / 1.00 millilitre(s)

SPECTINOMYCIN HYDROCHLORIDE
149.01 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 16 day

- **Cattle (calf)**
  - Meat and offal. 24 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01FF52
QJ01XX04

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

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:
Emdactilin 150 12 Vials with 250 ml Solution for injection
Emdactilin 150 6 Vials with 250 ml Solution for injection
Emdactilin 150 1 Vial with 250 ml Solution for injection
Emdactilin 150 12 Vials with 100 ml Solution for injection
Emdactilin 150 6 Vials with 100 ml Solution for injection

Emdactilin 150 1 Vial with 100 ml Solution for injection
Emdactilin 150 12 Vials with 50 ml Solution for injection
Emdactilin 150 6 Vials with 50 ml Solution for injection
Emdactilin 150 1 Vial with 50 ml Solution for injection

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:
Emdoka

Marketing authorisation date:
21/04/1995

Manufacturing sites for batch release:
Produlab Pharma B.V.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V169102

Date of authorisation status change:
3/11/2020

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

Documents

Summary of Product Characteristics

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Package Leaflet

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.
