Novocillin LC 1000 mg intramammary suspension for lactating cow

Authorised

- Oxacillin sodium monohydrate

Product identification

Medicine name:
Novocillin LC 1000 mg intramammary suspension for lactating cow
Novocillin LC 1000 mg интрамамарна суспензия за крави в лактационен период

Active substance:
Oxacillin sodium monohydrate

Target species:
Cattle (dairy cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Oxacillin sodium monohydrate
1042.50 milligram(s) / 10.00 gram(s)

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (dairy cow)**
  - Meat and offal. 6 day
  - Milk. 144 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51CF04

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Available in:
Bulgaria

Package description:
(ID3) 240 gram(s): Box (Cardboard) with 24 Applicator (Linear Low Density PolyEthylene) each with 10 gram(s)
(ID2) 200 gram(s): Box (Cardboard) with 20 Applicator (Linear Low Density PolyEthylene) each with 10 gram(s)
(ID1) 100 gram(s): Box (Cardboard) with 10 Applicator (Linear Low Density PolyEthylene) each with 10 gram(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Pharmanovo Veterinaerarzneimittel GmbH

Marketing authorisation date:
4/10/2021

Manufacturing sites for batch release:
Vet-Agro Trading Sp. z o.o.
Produlab Pharma B.V.
Przedsiebiorstwo Wielobranzowe Vet-Agro Sp. z o.o.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3084

Date of authorisation status change:
4/10/2021

Reference member state:
Germany

Procedure number:
DE/V/0333/001

Concerned member states:
Austria Bulgaria Cyprus Czechia Estonia France Hungary Iceland Ireland Italy Lithuania Poland Portugal Romania Spain
United Kingdom (Northern Ireland)

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 and Labelling

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

Combined File of all Documents