Rilexine 200 suspensija ievadīšanai tesmenī liellopiem

Authorised

- Cefalexin

Product identification

Medicine name:
Rilexine 200 suspensija ievadīšanai tesmenī liellopiem

Active substance:
Cefalexin

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:
Cefalexin
200.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (lactating cow)**
  - Milk. 2 day
    2 dienas (4 slaukšanas reizes)
  - Meat and offal. 4 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51DB01

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

Authorisation status:
Valid

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
27/09/2007

Manufacturing sites for batch release:
Virbac
Haupt Pharma Latina S.r.l.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/07/1702

Date of authorisation status change:
27/09/2007

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

Documents

Combined File of all Documents

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