Kefamast Dry Cow suspensija ievadīšanai tesmenī govīm

Not authorised

- Cefalexin
- Dihydrostreptomycin

Product identification

Medicine name:
Kefamast Dry Cow suspensija ievadīšanai tesmenī govīm

Active substance:
Cefalexin

Dihydrostreptomycin

Target species:
Cattle (dry cow)

Route of administration:
Intramammary use

Product details

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

Dihydrostreptomycin

500.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:
Intramammary suspension

Withdrawal period by route of administration:
Intramammary use:

- **Cattle (dry cow)**
  - Meat and offal. 28 day
  - Milk. 3 day

Ja 40 dienu laikā pēc zāļu lietošanas govs dzemdē, pienu pārtikā nedrīkst lietot ātrāk nekā 42,5 dienas pēc zāļu lietošanas.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ51RD01

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

Authorisation status:
Surrendered

Authorised in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis reviewed according to Acquis communautaire

Marketing authorisation holder:
Bimeda Animal Health Limited

Marketing authorisation date:
18/03/1996

Manufacturing sites for batch release:
Bimeda Animal Health Limited

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/95/0348

Date of authorisation status change:
26/01/2025

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.