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Gamared LC intramammary suspension

Authorised

- Prednisolone
- Novobiocin sodium
- Dihydrostreptomycin sulfate
- NEOMYCIN SULFATE
- Benzylpenicillin (procaine) monohydrate

Product identification

Medicine name:
Gamared LC intramammary suspension

Active substance:
Prednisolone

Novobiocin sodium

Dihydrostreptomycin sulfate

NEOMYCIN SULFATE

Benzylpenicillin (procaine) monohydrate

Target species:
Cattle (lactating cow)

Route of administration:
Intramammary use

Product details

Active substance and strength:

Prednisolone
10.00 milligram(s) / 1.00 Applicator

Novobiocin sodium
100.00 milligram(s) / 1.00 Applicator

Dihydrostreptomycin sulfate
125.00 milligram(s) / 1.00 Applicator

NEOMYCIN SULFATE
102000.00 international unit(s) / 1.00 Applicator

Benzylpenicillin (procaine) monohydrate
100.00 milligram(s) / 1.00 Applicator

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:

Intramammary use:

- **Cattle (lactating cow)**
  - Meat and offal. 7 day
  - Milk. 72 hour Мляко: 72 часа (6 издоявания)

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Informed consent application (Article 13c of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
26/04/2021

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-3052

Date of authorisation status change:
28/05/2023

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 and Labelling

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