

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

Benzympenicillin (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 издоивания)

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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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