

Intramar Lacto, 200 mg + 50 mg + 10mg, Intramammary suspension

Authorised

- Prednisolone
- Amoxicillin trihydrate
- Potassium clavulanate

Product identification

Medicine name:

Intramar Lacto, 200 mg + 50 mg + 10mg, Intramammary suspension

INTRAMAR Lacto 200 mg + 50 mg + 10 mg ενδομαστικό εναιώρημα για βοοειδή

Active substance:

Prednisolone

Amoxicillin trihydrate

Potassium clavulanate

Target species:

Cattle (lactating cow)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Prednisolone

10.00 milligram(s) / 1.00 Applicator

Amoxicillin trihydrate

229.61 milligram(s) / 1.00 Applicator

Potassium clavulanate

59.56 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. 84 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RV01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

Package description:

Plastic (LDPE) Applicator 24 x 1.0 Applicator 24x dezinf. ubrousky

Plastic (LDPE) Applicator 24 x 1.0 Applicator bez ubrousků

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application – bioavailability studies cannot be used to demonstrate bioequivalence (Article 19(1)(b) of Regulation (EU) 2019/6)

Marketing authorisation holder:

Bioveta a.s.

Marketing authorisation date:

13/04/2025

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:

CY00966V

Date of authorisation status change:

13/04/2025

Reference member state:

Czechia

Procedure number:

CZ/V/0188/001

Concerned member states:

Austria Belgium Croatia Cyprus France Germany Greece Hungary Ireland
Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia
Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

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