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:

Greece

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:
22/07/2025

Manufacturing sites for batch release:
Bioveta, a.s.

Responsible authority:
National Organization For Medicines

Authorisation number:
87767/23-07-2025/K-0260001

Date of authorisation status change:
13/03/2025

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

Summary of Product Characteristics

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

Package Leaflet

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