VIMCO emulsion for injection for ewes and female goats

Authorised

- Staphylococcus aureus, strain SP 140, Inactivated

Product identification

Medicine name:
VIMCO emulsion for injection for ewes and female goats
VIMCO ενέσιμο γαλάκτωμα για προβατίνες και θηλυκές αίγες

Active substance:
Staphylococcus aureus, strain SP 140, Inactivated

Target species:
Sheep (ewe)
Goat (adult female)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Staphylococcus aureus, strain SP 140, Inactivated
8.98 cells / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Sheep (ewe)**
  - Meat and offal. 0 day

- **Goat (adult female)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI03AB

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Cyprus

Package description:

box containing 1 PET vial containing 50 doses (100 ml)
box containing 1 PET vial containing 25 doses (50 ml)
box containing 1 PET vial containing 5 doses (10 ml)
box containing 1 glass vial containing 50 doses (100 ml)
box containing 1 glass vial containing 25 doses (50 ml)
box containing 1 glass vial containing 5 doses (10 ml)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
27/11/2023

Manufacturing sites for batch release:
Laboratorios Hipra, S.A.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00965V

Date of authorisation status change:
27/11/2023

Reference member state:
Spain

Procedure number:
ES/V/0209/001

Concerned member states:
Belgium Bulgaria Cyprus France Germany Greece Ireland Italy Netherlands Norway Poland Portugal Romania 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

English (PDF)
Published on: 24/07/2025
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Summary of Product Characteristics

Package Leaflet

Labelling