

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 emulsie voor injectie voor ooien en vrouwelijke geiten

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:

Netherlands

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:

3/01/2018

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 121005

Date of authorisation status change:

17/01/2022

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