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 EMULSION INJECTABLE POUR BREBIS ET CHEVRES

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 except for some pack sizes

Authorisation status:

Valid

Authorised in:

France

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:
14/05/2014

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

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/7140160 7/2014

Date of authorisation status change:
30/04/2019

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

Summary of Product Characteristics

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

Package Leaflet

Labelling

Package Leaflet and Labelling

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

Combined File of all Documents