INMEVA, SUSPENSION FOR INJECTION

Authorised

- Chlamydia abortus, strain A22, Inactivated
- Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated

Product identification

Medicine name:
INMEVA ενέσιμο εναιώρημα για πρόβατα
INMEVA, SUSPENSION FOR INJECTION

Active substance:
Chlamydia abortus, strain A22, Inactivated
Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Chlamydia abortus, strain A22, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Subcutaneous use:

- **Sheep**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Polyethylene (PET) vials of 10ml
Polyethylene (PET) vials of 250ml
Polyethylene (PET) vials of 100ml
Polyethylene (PET) vials of 50ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
22/07/2019

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

Responsible authority:
National Organization For Medicines

Authorisation number:
15839/13-02-2025/K-0235701

Date of authorisation status change:
12/02/2025

Reference member state:
France

Procedure number:
FR/V/0350/001

Concerned member states:
Austria Belgium Denmark Germany Greece Hungary Ireland Italy Luxembourg Netherlands Poland Portugal Romania Spain Sweden 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.