

INMEVA, SUSPENSION FOR INJECTION

Authorised

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

Product identification

Medicine name:

INMEVA, SUSPENSION FOR INJECTION

Inmeva, suspensão injetável

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:

Portugal

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:

18/04/2019

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

969/01/19DIVPT

Date of authorisation status change:

13/10/2022

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