Inmeva, Suspension for Injection

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

Authorised

Product identification

Medicine name:
INMEVA, SUSPENSION FOR INJECTION
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
  /
  1.00
  unit(s)
- Salmonella enterica, subsp. enterica, serovar Abortusovis, strain Sao, Inactivated
  1.00
  relative potency
  /
  1.00
  unit(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:

- United Kingdom (Northern Ireland)

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:

- 10/06/2019

Manufacturing sites for batch release:

- Laboratorios Hipra S.A.

Responsible authority:

- The Veterinary Medicines Directorate

Authorisation number:

- Vm 17533/4019

Date of authorisation status change:

- 15/08/2022

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

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