File downloaded on 2026-11-28
Source URL: https://medicines.health.europa.eu/veterinary/en/600000061750

Porcilis Ery+Parvo+Lepto suspension for injection for pigs

Authorised

- Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated
- Leptospira santarosai, serovar Gatuni, strain S1148/02, Inactivated
- Leptospira interrogans, serovar Pomona, strain Po-01-000, Inactivated
- Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated
- Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
- Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
- Porcine parvovirus, strain 014, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Product identification

Medicine name:
Porcilis Ery+Parvo+Lepto suspension for injection for pigs

Active substance:
Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated

Leptospira santarosai, serovar Gatuni, strain S1148/02, Inactivated

Leptospira interrogans, serovar Pomona, strain Po-01-000, Inactivated

Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated

Porcine parvovirus, strain 014, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Leptospira interrogans, serovar Bratislava, strain As-05-073, Inactivated
1310.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira santarosai, serovar Gatuni, strain S1148/02, Inactivated
276.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira interrogans, serovar Pomona, strain Po-01-000, Inactivated
166.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira kirschneri, serovar Dadas, strain GR-01-005, Inactivated
648.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira interrogans, serovar Copenhageni, strain Ic-02-001, Inactivated
210.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira interrogans, serovar Portlandvere, strain Ca-12-000, Inactivated
2816.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Porcine parvovirus, strain 014, Inactivated
130.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 Protective Dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

(ID6) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)
(ID5) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)
(ID4) 500 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 50 millilitre(s)
(ID3) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)
(ID2) 200 millilitre(s): Box (Cardboard) with 10 Bottle (PolyEthylene TerePhthalate) each with 20 millilitre(s)
(ID1) 20 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 20 millilitre(s)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
6/09/2016

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Danish Medicines Agency

Authorisation number:
56183

Date of authorisation status change:
6/09/2016

Reference member state:
Germany

Procedure number:
DE/V/0268/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain

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.

Combined File of all Documents