

Porcilis Ery+Parvo+Lepto suspension for injection for pigs

Authorised

- *Erysipelothrix rhusiopathiae*, serotype 2, strain M2, Inactivated
- Porcine parvovirus, strain 014, Inactivated
- *Leptospira interrogans*, serogroup Canicola, serovar Portland-vere, strain Ca-12-000, Inactivated
- *Leptospira interrogans*, serogroup Icterohaemorrhagiae, serovar Copenhageni, strain Ic-02-001, Inactivated
- *Leptospira kirschneri*, serogroup Grippotyphosa, serovar Dadas, strain GR-01-005, Inactivated
- *Leptospira interrogans*, serogroup Pomona, serovar Pomona, strain Po-01-000, Inactivated
- *Leptospira santarosai*, Serogroup Tarassovi, serovar Gatuni, strain S1148/02, Inactivated
- *Leptospira interrogans*, serogroup Australis, serovar Bratislava, strain As-05-073, Inactivated

Product identification

Medicine name:

Porcilis Ery+Parvo+Lepto suspension for injection for pigs

Porcilis Ery+Parvo+Lepto, suspensie voor injectie voor varkens

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Porcine parvovirus, strain 014, Inactivated

Leptospira interrogans, serogroup Canicola, serovar Portland-vero, strain Ca-12-000, Inactivated

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

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

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

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

Leptospira interrogans, serogroup Australis, serovar Bratislava, strain As-05-073, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

1.00 Protective Dose / 2.00 millilitre(s)

Porcine parvovirus, strain 014, Inactivated

130.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira interrogans, serogroup Canicola, serovar Portland-vero, strain Ca-12-000, Inactivated

2816.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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

210.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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

648.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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

166.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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

276.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Leptospira interrogans, serogroup Australis, serovar Bratislava, strain As-05-073,
Inactivated

1310.00 enzyme-linked immunosorbent assay unit / 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:

Netherlands

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 Nederland B.V.

Marketing authorisation date:

18/08/2016

Manufacturing sites for batch release:

INTERVET INTERNATIONAL B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 117666

Date of authorisation status change:

8/02/2022

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

Combined File of all Documents

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