

# 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

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**Target species:**

Pig

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**Route of administration:**

Intramuscular use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:****Intramuscular use:**

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**Pig**

- Meat and offal. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AL07

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovenia

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**Available in:**

Slovenia

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

1/12/2016

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

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**Authorisation number:**

DC/V/0554/001

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**Date of authorisation status change:**

1/12/2016

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0268/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.

### Labelling

This document does not exist in this language (English). You can find it in another language below.