# 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 injekčná suspenzia pre ošípané

**Active substance:**
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

**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-vere, 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)

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**Pharmaceutical form:**
Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

- **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:**
Slovakia

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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)

## 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:**
20/04/2017

**Manufacturing sites for batch release:**
INTERVET INTERNATIONAL B.V.

**Responsible authority:**
Institute For State Control Of Veterinary Biologicals And Medicaments

**Authorisation number:**
97/059/DC/16-S

**Date of authorisation status change:**
20/04/2017

**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)

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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

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

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