

# Porcilis Ery+Parvo Suspension for injection

Authorised

- Porcine parvovirus, strain 014, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

## Product identification

**Medicine name:**

Porcilis Ery+Parvo Suspension for injection

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**Active substance:**

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:**

Porcine parvovirus, strain 014, Inactivated

552.00 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:**

QI09AL01

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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:**

Norway

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**Available in:**

Norway

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**Package description:**

(ID44) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)

(ID34) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)

(ID24) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)

(ID14) 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:**

3/05/2005

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

04-3078

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**Date of authorisation status change:**

2/07/2009

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0233/001

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**Concerned member states:**

Norway Portugal

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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

English (PDF)

Published on: 13/03/2026

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### Combined File of all Documents