

# BIOSUIS ParvoEry suspension for injection for pigs

Authorised

- Porcine parvovirus, strain CAPM V198 S-27, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

## Product identification

**Medicine name:**

BIOSUIS ParvoEry suspension for injection for pigs

---

**Active substance:**

Porcine parvovirus, strain CAPM V198 S-27, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated

---

**Target species:**

Pig (sow)

---

**Route of administration:**

Intramuscular use

---

## Product details

**Active substance and strength:**

Porcine parvovirus, strain CAPM V198 S-27, Inactivated

4.00 log<sub>2</sub> haemagglutination inhibiting unit(s) / 1.00 Dose

Erysipelothrix rhusiopathiae, serotype 2, strain 2-64, Inactivated  
1.00 relative potency / 1.00 Dose

---

**Pharmaceutical form:**

Suspension for injection

---

**Withdrawal period by route of administration:**

**Intramuscular use:**

- 

**Pig (sow)**

- Meat and offal. 0 day

---

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

QI09AL01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Netherlands

---

**Package description:**

Glass Vial 1 x 10.0 millilitre(s)  
Glass Vial 10 x 10.0 millilitre(s)  
Glass Vial 1 x 50.0 millilitre(s)  
Plastic Vial 1 x 50.0 millilitre(s)  
Glass Vial 1 x 100.0 millilitre(s)  
Plastic Vial 1 x 100.0 millilitre(s)  
Plastic Vial 1 x 250.0 millilitre(s)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Bioveta a.s.

---

**Marketing authorisation date:**

11/10/2021

---

**Manufacturing sites for batch release:**

Bioveta a.s.

---

**Responsible authority:**

Medicines Evaluation Board

---

**Authorisation number:**

REG NL 127901

---

**Date of authorisation status change:**

28/01/2022

---

**Reference member state:**

Czechia

---

**Procedure number:**

CZ/V/0148/001

---

**Concerned member states:**

Belgium Bulgaria Croatia Denmark Estonia Germany Greece Hungary  
Ireland Latvia Lithuania Netherlands Norway Poland Portugal Spain Sweden  
United Kingdom (Northern Ireland)

---

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

## Documents

Combined File of all Documents

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

Labelling

English (PDF)

Published on: 14/03/2026

[Download](#)

Summary of Product Characteristics

Package Leaflet

eu-puar-czv0148001-mr-biosuis\_parvoery-en.pdf