

# Syvac Ery/Parvo emulsion for injection for pigs

Authorised

- Porcine parvovirus, strain PVP-7, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, Inactivated

## Product identification

**Medicine name:**

Syvac Ery/Parvo emulsion for injection for pigs

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**Active substance:**

Porcine parvovirus, strain PVP-7, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, 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 PVP-7, Inactivated

320.00 antibody unit(s) / 1.00 dose

Erysipelothrix rhusiopathiae, serotype 2, strain SE-9, Inactivated  
7.40 enzyme-linked immunosorbent assay unit / 1.00 dose

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**Pharmaceutical form:**

Emulsion 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:**

Belgium

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**Package description:**

Cardboard box with 1 vial containing 50 ml (25 doses)

Cardboard box with 1 vial containing 100 ml (50 doses)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Syva S.A.

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**Marketing authorisation date:**

10/11/2022

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**Manufacturing sites for batch release:**

Laboratorios Syva S.A.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V660813

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**Date of authorisation status change:**

10/11/2022

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0394/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Czechia Denmark Finland France  
Germany Greece Hungary Ireland Italy Netherlands Poland Portugal  
Romania Slovakia Sweden 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

English (PDF)

Published on: 24/03/2023

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Package Leaflet

English (PDF)

Published on: 20/04/2026

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Combined File of all Documents

Labelling

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