

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

Active substance:

Porcine parvovirus, strain PVP-7, Inactivated

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

Target species:

Pig

Route of administration:

Intramuscular use

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

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Pig

- 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:

Czechia

Available in:

Czechia

Package description:

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

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

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

Marketing authorisation holder:

Laboratorios Syva S.A.

Marketing authorisation date:

8/09/2022

Manufacturing sites for batch release:

Laboratorios Syva S.A.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/033/22-C

Date of authorisation status change:

8/09/2022

Reference member state:

Spain

Procedure number:

ES/V/0394/001

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)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 24/03/2023

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Combined File of all Documents

Package Leaflet

English (PDF)

Published on: 20/04/2026

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Labelling

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