

Authorised

Susvac P+Ery Każda dawka szczepionki (2 ml) zawiera:
Inaktywowany parwowirus świń,
szczep FCV117: nie mniej niż
1:100 HI* Inaktywowane bakterie
Erysipelotrix rhusiopathiae,
serotyp 2: 50 I.U. *HI – jednostki
hamowania hemaglutynacji
(badania odpowiedzi serologicznej
u kawii domowej) Emulsja do
wstrzykiwań

- Porcine parvovirus, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Product identification

Medicine name:

Susvac P+Ery Każda dawka szczepionki (2 ml) zawiera: Inaktywowany parwowirus świń, szczep FCV117: nie mniej niż 1:100 HI* Inaktywowane bakterie Erysipelotrix rhusiopathiae, serotyp 2: 50 I.U. *HI – jednostki hamowania hemaglutynacji (badania

odpowiedzi serologicznej u kawii domowej) Emulsja do wstrzykiwań

Active substance:

Porcine parvovirus, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Porcine parvovirus, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

50.00 international unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection/infusion

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:

Poland

Package description:

Available only in [Polish](#)

Available only in [Polish](#)

Available only in [Polish](#)

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:

Fatro S.p.A.

Marketing authorisation date:

7/03/2024

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

3303

Date of authorisation status change:

7/03/2024

To consult adverse reactions on veterinary medicinal products please go to 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

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

Labelling

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