

PARVORUVAX, sospensione iniettabile per suini

Authorised

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

Product identification

Medicine name:

PARVORUVAX, sospensione iniettabile per suini

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Porcine parvovirus, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

2.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Porcine parvovirus, Inactivated

1.00 log10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Suspension 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:

Italy

Package description:

Available only in [Italian](#)

Available only in [Italian](#)

Available only in [Italian](#)

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:

Ceva Salute Animale S.p.A.

Marketing authorisation date:

9/12/1993

Manufacturing sites for batch release:

Ceva-Phylaxia Zrt.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

9/12/2003

To consult adverse reactions on veterinary medicinal products please go to
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.