

Parvoruvac Injektionssuspension für Schweine

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, strain IM 950, Inactivated
- Porcine parvovirus, strain K22, Inactivated

Product identification

Medicine name:

Parvoruvac Injektionssuspension für Schweine

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, strain IM 950, Inactivated

Porcine parvovirus, strain K22, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain IM 950, Inactivated

1.00 Protective Dose / 2.00 millilitre(s)

Porcine parvovirus, strain K22, Inactivated

2.00 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:

Germany

Package description:

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Ceva Tiergesundheit GmbH

Marketing authorisation date:

23/11/2005

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France
Ceva-Phylaxia Zrt.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

490a/87

Date of authorisation status change:

19/11/2010

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.