

Ruvax

Not authorised

- *Erysipelothrix rhusiopathiae*, serotype 2, Inactivated

Product identification

Medicine name:

Ruvax

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Pig

- Meat and offal. no withdrawal period
Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Netherlands

Package description:

Available only in Dutch

Available only in Dutch

Available only in Dutch

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Boehringer Ingelheim Animal Health Netherlands B.V.

Marketing authorisation date:

14/05/1992

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 2820

Date of authorisation status change:

1/12/2022

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.