

RUVAX

Not
authorised

- *Erysipelothrix rhusiopathiae*, serotype 2, strain IM 950, Inactivated

Product identification

Medicine name:

RUVAX

Active substance:

Erysipelothrix rhusiopathiae, serotype 2, strain IM 950, Inactivated

Target species:

Pig

Sheep (pregnant ewe)

Turkey (for reproduction)

Pig (female)

Pig (male for reproduction)

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Erysipelothrix rhusiopathiae, serotype 2, strain IM 950, Inactivated

1.00 enzyme-linked immunosorbent assay unit / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Pig

- Meat and offal. 0 day

-

Sheep (pregnant ewe)

- Meat and offal. 0 day

-

Turkey (for reproduction)

- Meat and offal. 0 day

-

Pig (female)

- Meat and offal. 0 day

-

Pig (male for reproduction)

- Meat and offal. 0 day

Subcutaneous use:

-

Pig

- Meat and offal. 0 day

-

Sheep (pregnant ewe)

- Meat and offal. 0 day

-

Turkey (for reproduction)

- Meat and offal. 0 day

-

Pig (female)

- Meat and offal. 0 day

•

Pig (male for reproduction)

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Spain

Package description:

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

Available only in [Spanish](#)

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:

Boehringer Ingelheim Animal Health Espana S.A.

Marketing authorisation date:

13/04/1983

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Spanish Agency For Medicines And Health Products

Authorisation number:

2524 ESP

Date of authorisation status change:

1/02/2023

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.

Labelling

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.