Suibiovac Ery Inaktywowane bakterie:-Erysipelothrix rhusiopathiae szczep OV1 (serotyp 1) R.P* ≥ 1-Erysipelothrix rhusiopathiae szczep OV2 (serotyp 2) R.P* ≥ 1(*) względna moc oznaczona metodą in-vitro w porównaniu do szczepionki referencyjnej zgodnie zFarmakopeą Europejską. Zawiesina do wstrzykiwań

Authorised

- Erysipelothrix rhusiopathiae, serotype 1, strain OV1, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain OV2, Inactivated

Product identification

Medicine name:

Suibiovac Ery Inaktywowane bakterie:-Erysipelothrix rhusiopathiae szczep OV1 (serotyp 1) R.P* ≥ 1-Erysipelothrix rhusiopathiae szczep OV2 (serotyp 2) R.P* ≥ 1(*) względna moc oznaczona metodą in-vitro w porównaniu do szczepionki referencyjnej zgodnie zFarmakopeą Europejską. Zawiesina do wstrzykiwań

Active substance:
Erysipelothrix rhusiopathiae, serotype 1, strain OV1, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain OV2, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, serotype 1, strain OV1, Inactivated
1.00 relative potency / 2.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 2, strain OV2, Inactivated
1.00 relative potency / 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:
QI09AB03

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

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Biowet Drwalew Sp. z o.o.

Marketing authorisation date:
29/03/2011

Manufacturing sites for batch release:
Drwalewskie Zaklady Przemyslu Bioweterynaryjnego S.A.

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2077

Date of authorisation status change:
29/03/2011

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

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.

Summary of Product Characteristics

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