BIOSUIS Respi E Emulsja do wstrzykiwań

Authorised

- Actinobacillus pleuropneumoniae, serovar 2, Inactivated
- Actinobacillus pleuropneumoniae, serovars 9 and 11, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid
- Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
- Haemophilus parasuis, serotype 1, 5 and 13, Inactivated

Product identification

Medicine name:
BIOSUIS Respi E Emulsja do wstrzykiwań

Active substance:
Actinobacillus pleuropneumoniae, serovar 2, Inactivated
Actinobacillus pleuropneumoniae, serovars 9 and 11, Inactivated
Actinobacillus pleuropneumoniae, APX I toxoid
Actinobacillus pleuropneumoniae, APX II toxoid
Actinobacillus pleuropneumoniae, APX III toxoid
Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
Haemophilus parasuis, serotype 1, 5 and 13, Inactivated

Target species:

Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, serovar 2, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovars 9 and 11, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid
1.00 relative potency / 1.00 millilitre(s)

Erysipelothrix rhusiopathiae, serotype 1 and 2, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Haemophilus parasuis, serotype 1, 5 and 13, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection/infusion

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB

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
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish
Available only in Polish

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:
Bioveta a.s.

Marketing authorisation date:
15/05/2020

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
2969

Date of authorisation status change:
15/05/2020

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

Documents

Package Leaflet

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.

Summary of Product Characteristics

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