BioSuis Respi E, Injekční emulze

Authorised

- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 11, Inactivated
- Haemophilus parasuis, serotype 1, 5 and 13, Inactivated
- Erysipelothrix rhusiopathiae, Inactivated

Product identification

Medicine name:
BioSuis Respi E, Injekční emulze

Active substance:
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
Actinobacillus pleuropneumoniae, APX I toxoid
Actinobacillus pleuropneumoniae, APX II toxoid
Actinobacillus pleuropneumoniae, APX III toxoid
Actinobacillus pleuropneumoniae, serovar 11, Inactivated
Haemophilus parasuis, serotype 1, 5 and 13, Inactivated
Erysipelothrix rhusiopathiae, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, APX I toxoid
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, APX II toxoid
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, APX III toxoid
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, serovar 11, Inactivated
1.00 relative potency / 1.00 Dose

Haemophilus parasuis, serotype 1, 5 and 13, Inactivated
1.00 relative potency / 1.00 Dose

Erysipelothrix rhusiopathiae, Inactivated
1.00 relative potency / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

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:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

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:
2/04/2012

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/036/12-C

Date of authorisation status change:
21/09/2016

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.

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.