

BIOSUIS APP 2, 9, 11, Emulsion for injection

Authorised

- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
- Actinobacillus pleuropneumoniae, serovar 11, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid

Product identification

Medicine name:

BIOSUIS APP 2, 9, 11, Emulsion for injection

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Active substance:

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated

Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated

Actinobacillus pleuropneumoniae, serovar 11, Inactivated

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

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, serovar 11, 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

Pharmaceutical form:

Emulsion for injection

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

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Pig

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Package description:

Glass Vial 1 x 10.0 Dose

Glass Vial 1 x 50.0 Dose

Glass Vial 1 x 100.0 Dose

Plastic Vial 1 x 10.0 Dose

Plastic Vial 1 x 50.0 Dose

Plastic Vial 1 x 100.0 Dose

Plastic Bottle 1 x 250.0 Dose

Glass Vial 10 x 10.0 millilitre(s) - outer container: cardboard box

Plastic Vial 10 x 10.0 millilitre(s) - outer container: cardboard box

Glass Vial 10 x 10.0 millilitre(s) outer container: plastic box

Plastic Vial 10 x 10.0 millilitre(s) - outer container: plastic box

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:

23/01/2014

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

State Agency Of Medicines

Authorisation number:

1814

Date of authorisation status change:

23/01/2014

Reference member state:

Czechia

Procedure number:

CZ/V/0121/001

Concerned member states:

Croatia Estonia Greece Hungary Latvia Lithuania Poland Romania Slovakia

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.