

Polypleurosin APX Plus, süsteemulsioon sigadele

Authorised

- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Bordetella bronchiseptica, Inactivated
- Pasteurella multocida, serogroup D, Inactivated
- Pasteurella multocida, serogroup A, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, Inactivated
- Actinobacillus pleuropneumoniae, Inactivated

Product identification

Medicine name:

Polypleurosin APX Plus, süsteemulsioon sigadele

Active substance:

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Bordetella bronchiseptica, Inactivated

Pasteurella multocida, serogroup D, Inactivated

Pasteurella multocida, serogroup A, Inactivated

Actinobacillus pleuropneumoniae, serovar 2, Inactivated

Actinobacillus pleuropneumoniae, Inactivated

Target species:

Pig (piglet)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Actinobacillus pleuropneumoniae, APX III toxoid

1.00 relative potency / 1.00 dose

Actinobacillus pleuropneumoniae, APX II toxoid

1.00 relative potency / 1.00 dose

Actinobacillus pleuropneumoniae, APX I toxoid

1.00 relative potency / 1.00 dose

Bordetella bronchiseptica, Inactivated

1.00 relative potency / 1.00 dose

Pasteurella multocida, serogroup D, Inactivated

1.00 relative potency / 1.00 dose

Pasteurella multocida, serogroup A, Inactivated

1.00 relative potency / 1.00 dose

Actinobacillus pleuropneumoniae, serovar 2, Inactivated

1.00 relative potency / 1.00 dose

Actinobacillus pleuropneumoniae, Inactivated

1.00 relative potency / 1.00 dose

Pharmaceutical form:

Emulsion for injection

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

-

Pig (piglet)

- 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:

Available only in [Estonian](#)

Available only in [Estonian](#)

Available only in [Estonian](#)

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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:

25/04/2012

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

State Agency Of Medicines

Authorisation number:

1719

Date of authorisation status change:

25/04/2012

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.