POLYPLEUROSIN APX PLUS IM, Injekční emulze

Authorised

- Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Bordetella bronchiseptica, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid
- Pasteurella multocida, serogroup A, Inactivated
- Pasteurella multocida, serogroup D, Inactivated

Product identification

Medicine name:
POLYPLEUROSIN APX PLUS IM, Injekční emulze

Active substance:
Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
Bordetella bronchiseptica, Inactivated
Actinobacillus pleuropneumoniae, APX I toxoid
Actinobacillus pleuropneumoniae, APX II toxoid
Actinobacillus pleuropneumoniae, APX III toxoid
Pasteurella multocida, serogroup A, Inactivated

Pasteurella multocida, serogroup D, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, serovar 9, strain WSLB 3013, Inactivated
1.00 relative potency / 1.00 Dose

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
1.00 relative potency / 1.00 Dose

Bordetella bronchiseptica, 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

Pasteurella multocida, serogroup A, Inactivated
1.00 relative potency / 1.00 Dose

Pasteurella multocida, serogroup D, 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
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:
18/04/2003

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/032/03-C

Date of authorisation status change:
16/10/2008

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.