

# Suivac APP emulsion for injection for pigs

Authorised

- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid

## Product identification

**Medicine name:**

## Suivac APP emulsion for injection for pigs

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### **Active substance:**

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

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Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

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### **Target species:**

Pig

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### **Route of administration:**

Intramuscular use

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## Product details

### **Active substance and strength:**

Actinobacillus pleuropneumoniae, APX II toxoid

1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid

1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid  
1.00 international unit(s) / 2.00 millilitre(s)

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Pig**

- Meat and offal. 0 day

- Meat and offal. 0 day

- Meat and offal. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AB07

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Germany

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**Package description:**

100 ml (50 doses) vial in a cardboard box

50 ml (25 doses) vial in a cardboard box

500 ml (250 doses) vial in a cardboard box

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Chemvet DK A/S

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**Marketing authorisation date:**

1/09/2016

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**Manufacturing sites for batch release:**

Dyntec spol. s r.o.

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.11863.01.1

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**Date of authorisation status change:**

30/05/2020

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**Reference member state:**

Denmark

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**Procedure number:**

DK/V/0119/001

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**Concerned member states:**

Austria Germany Iceland Ireland Italy Norway Sweden

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.

PI.pdf