

# Porcilis APP suspensija injekcijām cūkām

Not  
authorised

- Actinobacillus pleuropneumoniae, outer membrane protein
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid

## Product identification

**Medicine name:**

Porcilis APP suspensija injekcijām cūkām

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**Active substance:**

Actinobacillus pleuropneumoniae, outer membrane protein

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

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**Target species:**

Pig (weaned piglet)

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**Route of administration:**

Intramuscular use

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## Product details

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

Actinobacillus pleuropneumoniae, outer membrane protein

50.00 unit(s) / 1.00 unit(s)

Actinobacillus pleuropneumoniae, APX I toxoid

50.00 unit(s) / 1.00 unit(s)

Actinobacillus pleuropneumoniae, APX II toxoid

50.00 unit(s) / 1.00 unit(s)

Actinobacillus pleuropneumoniae, APX III toxoid

50.00 unit(s) / 1.00 unit(s)

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### **Pharmaceutical form:**

Suspension for injection

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### **Withdrawal period by route of administration:**

#### **Intramuscular use:**

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#### **Pig (weaned piglet)**

- Not specified. 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:**

Surrendered

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### **Authorised in:**

Latvia

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### **Package description:**

Available only in [Latvian](#)

Available only in [Latvian](#)

Available only in Latvian

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis reviewed according to Acquis communautaire

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

22/03/1996

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/NRP/96/0380

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**Date of authorisation status change:**

1/12/2025

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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.

### Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

### Labelling

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