# Porcilis APP Vet. injektionsvæske, suspension

Authorised

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

## Product identification

**Medicine name:**
Porcilis APP Vet. injektionsvæske, suspension

**Active substance:**
Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, outer membrane protein

Actinobacillus pleuropneumoniae, APX III toxoid

**Target species:**
Pig

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**

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

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

Actinobacillus pleuropneumoniae, outer membrane protein
10000.00 unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid
100000.00 unit(s) / 2.00 millilitre(s)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

- **Pig**
  - Meat and offal. 0 day
  - 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:**

Denmark

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**Available in:**
Denmark

**Package description:**
Available only in Danish

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis reviewed according to Acquis communautaire

**Marketing authorisation holder:**
Intervet International B.V.

**Marketing authorisation date:**
5/12/1997

**Manufacturing sites for batch release:**
Intervet International B.V.

**Responsible authority:**
Danish Medicines Agency

**Authorisation number:**
16742

**Date of authorisation status change:**
5/12/1997

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Summary of Product Characteristics

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