

Porcilis APP suspension for injection for pigs

Authorised

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

Product identification

Medicine name:

Porcilis APP suspension for injection for pigs

Active substance:

Actinobacillus pleuropneumoniae, outer membrane protein

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, Inactivated

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Actinobacillus pleuropneumoniae, outer membrane protein

50.00 unit(s) / 1.00 Dose

Actinobacillus pleuropneumoniae, APX I toxoid

50.00 unit(s) / 1.00 Dose

Actinobacillus pleuropneumoniae, APX II toxoid

50.00 unit(s) / 1.00 Dose

Actinobacillus pleuropneumoniae, APX III toxoid

50.00 unit(s) / 1.00 Dose

Actinobacillus pleuropneumoniae, Inactivated

600.00 milligram(s) / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Pig

- 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:

Norway

Available in:

Norway

Package description:

Carton with one glass bottle type I (Ph. Eur.) or PET bottle with halogenated rubber stoppers and aluminium closures, containing 250 ml (125 doses).

Carton with one glass bottle type I (Ph. Eur.) or PET bottle with halogenated rubber stoppers and aluminium closures, containing 100 ml (50 doses).

Carton with one glass bottle type I (Ph. Eur.) or PET bottle with halogenated rubber stoppers and aluminium closures, containing 50 ml (25 doses).

Carton with one glass bottle type I (Ph. Eur.) or PET bottle with halogenated rubber stoppers and aluminium closures, containing 20 ml (10 doses).

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:

Intervet International B.V.

Marketing authorisation date:

27/01/2017

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:

16-11362

Date of authorisation status change:

17/12/2018

Reference member state:

Ireland

Procedure number:

IE/V/0327/001

Concerned member states:

Croatia Luxembourg Norway Portugal Romania

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

English (PDF)

Published on: 25/09/2024

[Download](#)

Package Leaflet

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