PORCILIS APP SUSPENSION INYECTABLE PARA PORCINO

Authorised

- Actinobacillus pleuropneumoniae, serovar 1, strain APP 1-l-452, Outer membrane protein
- Actinobacillus pleuropneumoniae, serovar 2, strain APP2, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain App2, APX II toxoid and serovar 7, strain App HV143, APX II toxoid
- Actinobacillus pleuropneumoniae, serovar 10, strain APP HV169, APX I toxoid

Product identification

Medicine name:
PORCILIS APP SUSPENSION INYECTABLE PARA PORCINO

Active substance:
Actinobacillus pleuropneumoniae, serovar 1, strain APP 1-l-452, Outer membrane protein

Actinobacillus pleuropneumoniae, serovar 2, strain APP2, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain App2, APX II toxoid and serovar 7, strain App HV143, APX II toxoid

Actinobacillus pleuropneumoniae, serovar 10, strain APP HV169, APX I toxoid

Target species:
Pig (weaned piglet)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, serovar 1, strain APP 1-I-452, Outer membrane protein
10000.00 Protective Dose / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain APP2, APX III toxoid
10000.00 Protective Dose / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2, APX II toxoid and serovar 7, strain App HV143, APX II toxoid
500.00 Protective Dose / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 10, strain APP HV169, APX I toxoid
500.00 Protective Dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig (weaned piglet)**
  - 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:

Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
4/02/1997

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

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3275 ESP

Date of authorisation status change:

7/09/2015

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.