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Porcilis APP injekčná suspenzia pre ošípané

Authorised

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

Product identification

Medicine name:
Porcilis APP injekčná suspenzia pre ošípané

Active substance:
Actinobacillus pleuropneumoniae, Inactivated

Actinobacillus pleuropneumoniae, outer membrane protein

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Actinobacillus pleuropneumoniae, Inactivated
600.00 milligram(s) / 2.00 millilitre(s)

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

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

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

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

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - All relevant tissues. 0 day zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Slovakia

Available in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak

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:
20/11/2002

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

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/054/02-S

Date of authorisation status change:
20/11/2002

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

Documents

Combined File of all Documents

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