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Porcilis APP

Authorised

- 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

Active substance:
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, outer membrane protein

50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid
50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid
50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid
50.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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:
Germany

Available in:
Germany

Package description:
Available only in German
Available only in German
Available only in German
Available only in German
Available only in German

Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
2/01/1996

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
210a/94

Date of authorisation status change:
9/07/2010

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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