Suivac APP emulsion for injection for pigs

Authorised

- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX III toxoid

Product identification

Medicine name:

Suivac APP emulsion for injection for pigs

Active substance:

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX III toxoid

Target species:

Pig

Route of administration:

Intramuscular use

Product details

Active substance and strength:

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

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

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

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

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

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

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

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

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

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
1.00 international unit(s) / 2.00 millilitre(s)

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

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

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

Pharmaceutical form:

Emulsion for injection

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

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
100 ml (50 doses) vial in a cardboard box
50 ml (25 doses) vial in a cardboard box
500 ml (250 doses) vial in a cardboard box

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:
Chemvet DK A/S

Marketing authorisation date:
14/10/2016

Manufacturing sites for batch release:
Dyntec spol. s r.o.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10395/001/001

Date of authorisation status change:
14/10/2016

Reference member state:
Denmark

Procedure number:
DK/V/0119/001

Concerned member states:
Austria Germany Iceland Ireland Italy Norway Sweden

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

Documents

PI.pdf