

SUIVAC APP, Injekční emulze

Authorised

- Actinobacillus pleuropneumoniae, serotype 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, serotype 9, strain App9KL97, Inactivated

Product identification

Medicine name:

SUIVAC APP, Injekční emulze

Active substance:

Actinobacillus pleuropneumoniae, serotype 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, serotype 9, strain App9KL97, Inactivated

Target species:

Pig

Route of administration:

Intradermal use

Intramuscular use

Product details

Active substance and strength:

Actinobacillus pleuropneumoniae, serotype 2, strain App2TR98, Inactivated

1.00 unit(s) / 1.00 Dose

Actinobacillus pleuropneumoniae, serotype 9, strain App9KL97, Inactivated

1.00 unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Intradermal use:****• Pig**

- Meat and offal. 0 day

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:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

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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:

Dyntec spol. s r.o.

Marketing authorisation date:

2/11/2001

Manufacturing sites for batch release:

Dyntec spol. s r.o.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicines

Authorisation number:

97/056/01-C

Date of authorisation status change:

2/11/2001

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

Documents

Summary of Product Characteristics

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Package Leaflet

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Labelling

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