Suivac APP injekčná emulzia

Authorised

- Actinobacillus pleuropneumoniae, Inactivated
- Actinobacillus pleuropneumoniae, Inactivated

Product identification

Medicine name:
Suivac APP injekčná emulzia

Active substance:
Actinobacillus pleuropneumoniae, Inactivated

Actinobacillus pleuropneumoniae, Inactivated

Target species:
Pig
Pig

Route of administration:
Intradermal use
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, Inactivated
1.00 billion colony forming units / 0.20 millilitre(s)

Actinobacillus pleuropneumoniae, Inactivated

1.00 billion colony forming units / 2.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intradermal use:

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

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
Available only in Slovak
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:
Dyntec spol. s r.o.

Marketing authorisation date:
2/05/2002

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

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

Authorisation number:
97/021/02-S

Date of authorisation status change:
2/05/2002

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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