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POLYPLEUROSIN APN PLUS IM

Authorised

- Pasteurella multocida, serogroup D, Inactivated
- Pasteurella multocida, serogroup A, Inactivated
- Bordetella bronchiseptica, Inactivated
- Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid

Product identification

Medicine name:
POLYPLEUROSIN APN PLUS IM

Active substance:
Pasteurella multocida, serogroup D, Inactivated

Pasteurella multocida, serogroup A, Inactivated

Bordetella bronchiseptica, Inactivated

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Pasteurella multocida, serogroup D, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Pasteurella multocida, serogroup A, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Bordetella bronchiseptica, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 9, strain App9KL97, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain App2TR98, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX III toxoid
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX I toxoid
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid
1.00 relative potency / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
28/05/2007

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
120238

Date of authorisation status change:
28/05/2007

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

Documents

Summary of Product Characteristics

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