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Hyobac App Multi Vet., emulsion for injection

Authorised

- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated
- Actinobacillus pleuropneumoniae, APX I toxoid
- Actinobacillus pleuropneumoniae, APX II toxoid
- Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated

Product identification

Medicine name:
Hyobac App Multi Vet., emulsion for injection

Active substance:

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, APX II toxoid

Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Actinobacillus pleuropneumoniae, APX II toxoid
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
1.00 relative potency / 1.00 millilitre(s)

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

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
1.00 relative potency / 1.00 millilitre(s)

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

Actinobacillus pleuropneumoniae, serovar 6, strain WSLB 3075, Inactivated
1.00 relative potency / 1.00 millilitre(s)

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

Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated
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)

Actinobacillus pleuropneumoniae, serovar 5, strain WSLB 3079, Inactivated
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
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - Meat and offal. 0 day
  - 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:
Norway

Available in:
Norway

Package description:
100 ml in vial (HDPE) in paper cardboard
250 ml vial (HDPE) in paper cardboard
100 ml in vial (type II glass) in paper cardboard

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:
Salfarm Danmark A/S

Marketing authorisation date:
15/03/2017

Manufacturing sites for batch release:
Bioveta a.s.
Institute For State Control Of Veterinary Biologicals And Medicaments

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:
16-11342

Date of authorisation status change:
1/12/2021

Reference member state:
Denmark

Procedure number:
DK/V/0121/001

Concerned member states:
Finland Norway Sweden

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 25/01/2022
Updated on: 14/03/2026
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Package Leaflet

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