

Hyobac App Multi Vet., emulsion for injection

Authorised

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

Product identification

Medicine name:

Hyobac App Multi Vet., emulsion for injection

Hyobac App Multi Vet. Injektionsvätska, emulsion

Active substance:

Actinobacillus pleuropneumoniae, APX III toxoid

Actinobacillus pleuropneumoniae, APX I toxoid

Actinobacillus pleuropneumoniae, serotype 5, strain WSLB 3079, Inactivated

Actinobacillus pleuropneumoniae, serotype 6, strain WSLB 3075, Inactivated

Actinobacillus pleuropneumoniae, serotype 2, strain WSLB 3012, Inactivated

Actinobacillus pleuropneumoniae, APX II toxoid

Target species:

Pig

Intramuscular use

Active substance and strength:

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, serotype 5, strain WSLB 3079, Inactivated

1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serotype 6, strain WSLB 3075, Inactivated

1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, serotype 2, strain WSLB 3012, Inactivated

1.00 relative potency / 1.00 millilitre(s)

Actinobacillus pleuropneumoniae, APX II toxoid

1.00 relative potency / 1.00 millilitre(s)

Emulsion for injection

Intramuscular use:

- **Pig**

- [illegible]

- 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:

Sweden

Available in:

Sweden

Package description:

Available only in Swedish

Available only in Swedish

100 ml in vial (HDPE) 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:

3/05/2017

Manufacturing sites for batch release:

Bioveta a.s.
USKVBL

Responsible authority:

Swedish Medical Products Agency

Authorisation number:

55370

Date of authorisation status change:

3/05/2017

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

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Package Leaflet

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