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Hyobac App 2 Vet., emulsion for injection

Authorised

- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, APX III toxoid
- Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, APX II toxoid

Product identification

Medicine name:
Hyobac App 2 Vet., emulsion for injection

Active substance:
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, Inactivated
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, APX III toxoid
Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, APX II toxoid

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:

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

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

Actinobacillus pleuropneumoniae, serovar 2, strain WSLB 3012, 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
  - 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

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Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:
100 ml in vial (type II glass) in paper cardboard
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:
10/04/2012

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Danish Medicines Agency

Authorisation number:
48942

Date of authorisation status change:
10/04/2012

Reference member state:
Denmark

Procedure number:
DK/V/0120/001

Concerned member states:
Sweden

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

Documents

Summary of Product Characteristics

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PI Hyobac App 2.pdf