

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

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

100 ml in 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:

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