Suvaxyn M.Hyo Parasuis Suspension for Injection for Pigs

Authorised

- Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
- Haemophilus parasuis, serotype 4, strain 2170B, Inactivated
- Haemophilus parasuis, serotype 5, strain IA84-29755, Inactivated

Product identification

Medicine name:
Suvaxyn M.Hyo Parasuis Suspension for Injection for Pigs
Suvaxyn M.hyo suspensão injetável para porcos

Active substance:
Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
Haemophilus parasuis, serotype 4, strain 2170B, Inactivated
Haemophilus parasuis, serotype 5, strain IA84-29755, Inactivated

Target species:
Pig (for fattening)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
1.90 unknown / 2.00 millilitre(s)

Haemophilus parasuis, serotype 4, strain 2170B, Inactivated
8.10 unknown / 2.00 millilitre(s)

Haemophilus parasuis, serotype 5, strain IA84-29755, Inactivated
3.40 unknown / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (for fattening)**
  - Meat and offal. no withdrawal period — withdrawal period is 0 days

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Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB17

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Portugal

Package description:

(ID2): 1 Box with 1 Bottle (High Density PolyEthylene) with 60 millilitre(s) (60 millilitre(s))
(ID7): 1 Box with 10 Bottle (High Density PolyEthylene) with 120 millilitre(s) (1200 millilitre(s))

(ID9): 1 Box with 1 Sachet (Low Density PolyEthylene) with 100 millilitre(s) (100 millilitre(s))
(ID10): 1 Box with 10 Sachet (Low Density PolyEthylene) with 100 millilitre(s) (1000 millilitre(s))
(ID3): 1 Box with 1 Bottle (High Density PolyEthylene) with 120 millilitre(s) (120 millilitre(s))
(ID4): 1 Box with 1 Bottle (High Density PolyEthylene) with 250 millilitre(s) (250 millilitre(s))
(ID8): 1 Box with 10 Bottle (High Density PolyEthylene) with 250 millilitre(s) (2500 millilitre(s))
(ID6): 1 Box with 10 Bottle (High Density PolyEthylene) with 60 millilitre(s) (600 millilitre(s))
(ID1): 1 Box with 1 Bottle (High Density PolyEthylene) with 25 millilitre(s) (25 millilitre(s))
(ID5): 1 Box with 10 Bottle (High Density PolyEthylene) with 25 millilitre(s) (250 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Portugal Lda.

Marketing authorisation date:
31/10/1995

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
524/94 DGV

Date of authorisation status change:
1/01/2018

Reference member state:
Germany

Procedure number:
DE/V/0280/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Finland France Greece Ireland Italy Luxembourg Malta Netherlands Norway Poland Portugal Romania Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Combined File of all Documents

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