Suvaxyn MH-One Emulsion for injection for pigs

Authorised

- Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated

Product identification

Medicine name:
Suvaxyn MH-One Emulsion for injection for pigs
Suvaxyn MH-One

Active substance:
Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Mycoplasma hyopneumoniae, strain P-5722-3, Inactivated
1.00 relative unit(s) / 2.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

(ID1): 1 Box with 1 Bottle (High Density PolyEthylene) with 20 millilitre(s) (20 millilitre(s))
(ID6): 1 Box with 10 Bottle (High Density PolyEthylene) with 250 millilitre(s) (2500 millilitre(s))
(ID2): 1 Box with 10 Bottle (High Density PolyEthylene) with 20 millilitre(s) (200 millilitre(s))
(ID3): 1 Box with 1 Bottle (High Density PolyEthylene) with 100 millilitre(s) (100 millilitre(s))
(ID4): 1 Box with 1 Bottle (High Density PolyEthylene) with 250 millilitre(s) (250 millilitre(s))
(ID5): 1 Box with 10 Bottle (High Density PolyEthylene) with 100 millilitre(s) (1000 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 Deutschland GmbH

Marketing authorisation date:
1/04/2005

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.02689.01.1

Date of authorisation status change:
12/08/2013

Reference member state:
Germany

Procedure number:
DE/V/0248/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

English (PDF)
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