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Suvaxyn MH-One Emulsion for injection for pigs

Not authorised

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

Product identification

Medicine name:
Suvaxyn MH-One Emulsion for injection for pigs
SUVAXYN MH-One, injekcinė emulsija kiaulėms

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

Authorisation status:

Surrendered

Authorised in:

Lithuania

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 Belgium

Marketing authorisation date:
28/12/2008

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

Responsible authority:
State Food And Veterinary Service

Authorisation number:
LT/2/08/1822/001-003

Date of authorisation status change:
11/08/2013

Reference member state:
Germany

Procedure number:
DE/V/0248/001

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

RV1822.pdf