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Suvaxyn MH-One Emulsion for injection for pigs

Authorised

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

Product identification

Medicine name:
Suvaxyn MH-One emulsão injetável para suínos
Suvaxyn MH-One Emulsion for injection for pigs

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:
Valid

Authorised in:
Portugal

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 Portugal Lda.

Marketing authorisation date:
18/07/2008

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
804/08 RIVPT

Date of authorisation status change:
18/07/2008

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

Summary of Product Characteristics

Combined File of all Documents

2603675-paren-20250201.pdf