Suvaxyn MH-One Emulsion for injection for pigs

Authorised

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

Product identification

Medicine name:
SUVAXYN MH-ONE ΕΝΕΣΙΜΟ ΓΑΛΑΚΤΩΜΑ
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

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Authorisation status:

Valid

Authorised in:

Greece

Available in:

Greece

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 Hellas S.A.

Marketing authorisation date:
30/11/2008

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

Responsible authority:
National Organization For Medicines

Authorisation number:
14082/21-02-2014/K-0178801

Date of authorisation status change:
20/02/2014

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

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

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