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Porcilis Ery Suspension for injection

Authorised

- Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Product identification

Medicine name:
Porcilis Ery Suspension for injection

Active substance:
Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated
1.00 Protective Dose / 2.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AB03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
(ID6) 100 millilitre(s): Box with 1 Vial (Polyethylenterephthalat) with 100 millilitre(s), closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium)
(ID2) 20 millilitre(s): Box with 1 Vial (Polyethylenterephthalat) with 20 millilitre(s), closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium)
(ID8) 250 millilitre(s): Box with 1 Vial (Polyethylenterephthalat) with 250 millilitre(s), closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium)
(ID4) 50 millilitre(s): Box with 1 Vial (Polyethylenterephthalat) with 50 millilitre(s), closed with Stopfen and Kappe (Halobutyl Rubber, Aluminium)

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:
Intervet International B.V.

Marketing authorisation date:
3/05/2005

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
04-3072

Date of authorisation status change:
27/08/2008

Reference member state:
Germany

Procedure number:
DE/V/0232/001

Concerned member states:
Norway

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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

Combined File of all Documents