Porcilis Ery+Parvo Suspension for injection

Authorised

- Porcine parvovirus, strain 014, Inactivated
- Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Product identification

Medicine name:
Porcilis Ery+Parvo Suspension for injection

Active substance:
Porcine parvovirus, strain 014, Inactivated

Erysipelothrix rhusiopathiae, serotype 2, strain M2, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine parvovirus, strain 014, Inactivated
552.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

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:
QI09AL01

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

Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
(ID44) 250 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 250 millilitre(s)
(ID34) 100 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s)
(ID24) 50 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 50 millilitre(s)
(ID14) 20 millilitre(s): Box (Cardboard) with 1 Bottle (PolyEthylene TerePhthalate) with 20 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:
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:
04-3078

Date of authorisation status change:
2/07/2009

Reference member state:
Germany

Procedure number:
DE/V/0233/001

Concerned member states:
Norway Portugal

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.

Combined File of all Documents