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:
Germany

Available in:
Germany

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 Deutschland GmbH

Marketing authorisation date:
7/07/1999

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
59a/95

Date of authorisation status change:
25/02/2010

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

Combined File of all Documents

English (PDF)
Published on: 27/06/2024
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Package Leaflet