Porcilis PCV (--) - Emulsion for injection

Authorised

- Porcine circovirus type 2, ORF2 capsid protein

Product identification

Medicine name:
Porcilis PCV (--) - Emulsion for injection

Active substance:
Porcine circovirus type 2, ORF2 capsid protein

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine circovirus type 2, ORF2 capsid protein

Presentation_strength:4.5 log2 Ab titre Reference:Hse Comments:II.C.2.1.2 Index:0

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Bulgaria , Czechia , Denmark , Finland , France , Germany , Hungary , Ireland , Italy , Latvia , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (PET), Package_size:10 vials, Content:200 ml

Packaging:Vial (PET), Package_size:10 vials, Content:100 ml

Packaging:Vial (PET), Package_size:10 vials, Content:50 ml

Packaging:Vial (PET), Package_size:10 vials, Content:20 ml

Packaging:Vial (PET), Package_size:1 vial, Content:500 ml

Packaging:Vial (PET), Package_size:1 vial, Content:200 ml

Packaging:Vial (PET), Package_size:1 vial, Content:100 ml

Packaging:Vial (PET), Package_size:1 vial, Content:50 ml

Packaging:Vial (PET), Package_size:10 vials, Content:500 ml

Packaging:Vial (PET), Package_size:1 vial, Content:20 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
12/01/2009

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

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
12/01/2009

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: 29/02/2024

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