

Porcilis PCV ID \geq 1436 AU - Emulsion for injection

Authorised

- Porcine circovirus type 2, ORF2 capsid protein

Product identification

Medicine name:

Porcilis PCV ID \geq 1436 AU - Emulsion for injection

Active substance:

Porcine circovirus type 2, ORF2 capsid protein

Target species:

Pig

Route of administration:

Intradermal use

Product details

Active substance and strength:

Porcine circovirus type 2, ORF2 capsid protein

Presentation_strength: \geq 1436 AU/dose Comments:AU - Antigenic units as determined in the in vitro antigenic mass assay Index:0

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:**Intradermal use:**

-

Pig

- All relevant tissues. 0 day
Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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 , Cyprus , Czechia , Denmark , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:10 vials, Content:10 ml

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

Packaging:Vial (glass), Package_size:1 vial, Content:10 ml

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

28/08/2015

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:

28/08/2015

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

Documents

Combined File of all Documents

English (PDF)

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