

Suvaxyn Circo (--)- Emulsion for injection

Authorised

- Porcine circovirus type 2, ORF2 capsid protein

Product identification

Medicine name:

Suvaxyn Circo (--)- Emulsion for injection

Active substance:

Porcine circovirus type 2, ORF2 capsid protein

Target species:

Pig (for fattening)

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Porcine circovirus type 2, ORF2 capsid protein

Presentation_strength:RP ≥ 0.7 Reference:PCV02-MOP Index:0

Pharmaceutical form:

Emulsion for injection

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

-

Pig (for fattening)

- Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AA

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:

Belgium , Greece , Luxembourg , Netherlands , Poland , Romania , Spain

Package description:

Packaging:Vial (HDPE), Package_size:1 vial, Content:100 ml (50 doses)

Packaging:Vial (HDPE), Package_size:4 vials, Content:250 ml (125 doses)

Packaging:Vial (HDPE), Package_size:1 vial, Content:250 ml (125 doses)

Packaging:Vial (HDPE), Package_size:10 vials, Content:50 ml (25 doses)

Packaging:Vial (HDPE), Package_size:10 vials, Content:100 ml (50 doses)

Packaging:Vial (HDPE), Package_size:1 vial, Content:50 ml (25 doses)

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:

Zoetis Belgium

Marketing authorisation date:

7/02/2018

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

7/02/2018

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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