CircoMax (--) - Emulsion for injection

Authorised

- Porcine circovirus type 1, strain cPCV1-2, expressing the ORF2 protein gene of Porcine circovirus type 2a, Inactivated
- Porcine circovirus type 1, strain cPCV1-2b, expressing the ORF2 protein gene of Porcine circovirus type 2b, Inactivated

Product identification

Medicine name:
CircoMax (--) - Emulsion for injection

Active substance:
Porcine circovirus type 1, strain cPCV1-2, expressing the ORF2 protein gene of Porcine circovirus type 2a, Inactivated

Porcine circovirus type 1, strain cPCV1-2b, expressing the ORF2 protein gene of Porcine circovirus type 2b, Inactivated

Target species:
Pig (for fattening)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Porcine circovirus type 1, strain cPCV1-2, expressing the ORF2 protein gene of Porcine circovirus type 2a, Inactivated

Presentation_strength:1.5-4.9 RP Reference:HSE Index:0

Porcine circovirus type 1, strain cPCV1-2b, expressing the ORF2 protein gene of Porcine circovirus type 2b, Inactivated

Presentation_strength:1.5-5.9 RP Reference:HSE Index:1

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (for fattening)**
  - Not specified. 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 , Estonia , Finland , France , Greece , Hungary , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Poland , Romania ,

Slovakia , Spain , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (HDPE), Package_size:4 vials, Content:250 ml (125 doses)
Packaging:Vial (HDPE), Package_size:10 vials, Content:100 ml (50 doses)
Packaging:Vial (HDPE), Package_size:1 vial, Content:250 ml (125 doses)
Packaging:Vial (HDPE), Package_size:1 vial, Content:100 ml (50 doses)
Packaging:Vial (HDPE), Package_size:10 vials, Content:50 ml (25 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 - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Belgium SA

Marketing authorisation date:
11/01/2022

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:
11/01/2022

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: 11/07/2025
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