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Circovac (--) - Emulsion and suspension for emulsion for injection

Authorised

- Porcine circovirus type 2, strain 1010, Inactivated

Product identification

Medicine name:
Circovac (--) - Emulsion and suspension for emulsion for injection

Active substance:
Porcine circovirus type 2, strain 1010, Inactivated

Target species:
Pig (female)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Porcine circovirus type 2, strain 1010, Inactivated

Presentation_strength:≥ 1.8 log 10 Elisa U Reference:Hse Index:0

Pharmaceutical form:

Emulsion and suspension for emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (female)**
  - Not applicable. 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 , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Italy , Latvia , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Suspension: vial (LDPE); Emulsion: vial (LDPE), Package_size:Suspension: 10 vials + Emulsion: 10 vials, Content:Suspension: 50 ml Emulsion: 100 ml (50 doses for gilts and sows, 200 doses for piglets)
Packaging:Suspension: vial (LDPE); Emulsion: vial (LDPE), Package_size:Suspension: 1 vial + Emulsion: 1 vial, Content:Suspension: 50 ml Emulsion: 100 ml (50 doses for gilts and sows, 200 doses for piglets)
Packaging:Suspension: vial (glass); Emulsion: vial (glass), Package_size:Suspension: 1 vial + Emulsion: 1 vial, Content:Suspension: 5 ml Emulsion: 10 ml (5 doses for gilts and sows, 20 doses for piglets)

Packaging:Suspension: vial (glass); Emulsion: vial (glass), Package_size:Suspension: 10 vials + Emulsion: 10 vials, Content:Suspension: 5 ml Emulsion: 10 ml (5 doses for gilts and sows, 20 doses for piglets)
Packaging:Suspension: vial (glass) Emulsion: vial (glass), Package_size:Suspension: 1 vial + Emulsion: 1 vial, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)
Packaging:Suspension: vial (glass); Emulsion: vial (glass), Package_size:Suspension: 10 vials + Emulsion : 10 vials, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)
Packaging:Suspension: vial (glass); Emulsion: vial (PP), Package_size:Suspension: 1 vial + Emulsion: 1 vial, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)
Packaging:Suspension: vial (glass); Emulsion: vial (PP), Package_size:Suspension: 10 vials + Emulsion: 10 vials, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)
Packaging:Suspension: vial (glass) Emulsion: vial (LDPE), Package_size:Suspension: 10 vials + Emulsion: 10 vials, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)
Packaging:Suspension: vial (glass) Emulsion: vial (LDPE), Package_size:Suspension: 1 vial + Emulsion: 1 vial, Content:Suspension: 20 ml Emulsion: 50 ml (25 doses for gilts and sows, 100 doses for piglets)

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:
Ceva-Phylaxia Zrt.

Marketing authorisation date:
21/06/2007

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
European Commission

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

Date of authorisation status change:
21/06/2007

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Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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