

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: $\geq 1.8 \log_{10}$ Elisa U Reference:Hse Index:0

Pharmaceutical form:

Emulsion and suspension for emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

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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)

Package description:

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)

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 (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 (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 (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 (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: 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: 5 ml Emulsion: 10 ml (5 doses for gilts and sows, 20 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 (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)

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 Co. Ltd

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:

7/12/2021

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: 28/08/2024

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