Respiporc FLU3 (--) - Suspension for injection

Authorised

- Influenza A virus, subtype H1N1, strain A/swine/Haselünne/IDT2617/2003, Inactivated
- Influenza A virus, subtype H3N2, strain A/swine/Bakum/IDT1769/2003, Inactivated
- Influenza A virus, subtype H1N2, strain A/swine/Bakum/1832/2000, Inactivated

Product identification

Medicine name:
Respiporc FLU3 (--) - Suspension for injection

Active substance:
Influenza A virus, subtype H1N1, strain A/swine/Haselünne/IDT2617/2003, Inactivated
Influenza A virus, subtype H3N2, strain A/swine/Bakum/IDT1769/2003, Inactivated
Influenza A virus, subtype H1N2, strain A/swine/Bakum/1832/2000, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H1N1, strain A/swine/Haselünne/IDT2617/2003, Inactivated

Presentation_strength:min. 10.22 log_2 GMNU Reference:Hse Index:0

Influenza A virus, subtype H3N2, strain A/swine/Bakum/IDT1769/2003, Inactivated

Presentation_strength:min. 10.53 log_2 GMNU Reference:Hse Index:1

Influenza A virus, subtype H1N2, strain A/swine/Bakum/1832/2000, Inactivated

Presentation_strength:min. 12.34 log_2 GMNU Reference:Hse Index:2

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AA03

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)

Package description:

Packaging:Vial (PET), Package_size:8 vials, Content:500 ml (250 doses)

Packaging:Vial (PET), Package_size:1 vial, Content:100 ml (50 doses)
Packaging:Vial (PET), Package_size:1 vial, Content:50 ml (25 doses)
Packaging:Vial (PET), Package_size:1 vial, Content:20 ml (10 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:100 ml (50 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:50 ml (25 doses)
Packaging:Vial (glass), Package_size:1 vial, Content:20 ml (10 doses)
Packaging:Bottle (LDPE), Package_size:1 bottle, Content:50 ml (25 doses)
Packaging:Bottle (LDPE), Package_size:1 bottle, Content:100 ml (50 doses)

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 Sante Animale

Marketing authorisation date:
14/01/2010

Manufacturing sites for batch release:
IDT Biologika GmbH
CEVA-Phylaxia Zrt.

Responsible authority:
European Commission

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

Date of authorisation status change:
24/10/2024

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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