Respiporc FLUpan H1N1 (--) - Suspension for injection

Authorised

- Influenza A virus, subtype H1N1, strain A/Jena/VI5258/2009 pdm09, Inactivated

Product identification

Medicine name:
Respiporc FLUpan H1N1 (--) - Suspension for injection

Active substance:
Influenza A virus, subtype H1N1, strain A/Jena/VI5258/2009 pdm09, Inactivated

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Influenza A virus, subtype H1N1, strain A/Jena/VI5258/2009 pdm09, Inactivated
Presentation_strength:≥ 16 HU Reference:HSE Index:0

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

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

Ceva Sante Animale

Marketing authorisation date:
17/05/2017

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:
6/05/2024

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: 30/03/2026
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