

# ProteqFlu (--) - Suspension for injection

Authorised

- Canarypox virus, strain vCP3011, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Richmond/1/2007), Live
- Canarypox virus, strain vCP2242, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Ohio/03), Live

## Product identification

**Medicine name:**

ProteqFlu (--) - Suspension for injection

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**Active substance:**

Canarypox virus, strain vCP3011, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Richmond/1/2007), Live

Canarypox virus, strain vCP2242, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Ohio/03), Live

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**Target species:**

Horse

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**Route of administration:**

Intramuscular use

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## Product details

### Active substance and strength:

Canarypox virus, strain vCP3011, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Richmond/1/2007), Live

Presentation\_strength:vCP3011  $\geq 10^{5.3}$  FAID50 Reference:Hse Index:0

Canarypox virus, strain vCP2242, expressing haemagglutinin gene of Influenza A virus subtype H3N8 (A/equine/Ohio/03), Live

Presentation\_strength:vCP2242  $\geq 10^{5.3}$  FAID50 Reference:Hse Index:1

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### Pharmaceutical form:

Suspension for injection

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### Withdrawal period by route of administration:

#### Intramuscular use:

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#### Horse

- Not applicable. 0 day Zero days

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### Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AD02

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### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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### Authorisation status:

Valid

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### 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)

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### Available in:

Belgium , Czechia , France , Luxembourg , Poland , Slovakia , Spain

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**Package description:**

Packaging:Vial (glass), Package\_size:10 vials, Content:1 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Complete application (stand-alone) - Council Directive 81/851/EEC

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

6/03/2003

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

13/02/2006

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

English (PDF)

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