

# BioEquin FT suspension for injection for horses

Authorised

- Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated
- Clostridium tetani, toxoid
- Influenza A virus, subtype H3N8, strain A/equine/Limerick/2010, Inactivated

## Product identification

**Medicine name:**

BioEquin FT suspension for injection for horses

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**Active substance:**

Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated

Clostridium tetani, toxoid

Influenza A virus, subtype H3N8, strain A/equine/Limerick/2010, Inactivated

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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:**

Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated

5.00 log<sub>2</sub> haemagglutination inhibiting unit(s) / 1.00 Dose

Clostridium tetani, toxoid

30.00 international unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Limerick/2010, Inactivated

5.00 log<sub>2</sub> haemagglutination inhibiting unit(s) / 1.00 Dose

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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**

- Milk. 0 hour

- Meat and offal. 0 day

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

QI05AL01

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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:**

Romania

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**Package description:**

glass vial 2 x 1.0 dose

glass vial 5 x 1.0 dose

glass vial 10 x 1.0 dose

glass vial 1 x 5.0 dose

glass vial 10 x 5.0 dose

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Bioveta a.s.

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**Marketing authorisation date:**

14/05/2025

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Institute For Control Of Biological Products And Veterinary Medicines

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**Authorisation number:**

250062

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**Date of authorisation status change:**

14/05/2025

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0201/001

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**Concerned member states:**

Austria Belgium Bulgaria Denmark Finland France Germany Ireland Italy  
Netherlands Norway Poland Portugal Romania Slovakia Spain Sweden  
United Kingdom (Northern Ireland)

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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

This document does not exist in this language (English). You can find it in another language below.

Labelling

Summary of Product Characteristics

Package Leaflet

eu-puar-czv0201001-mr-bioequin\_ft-en.pdf