

# BioEquin FH, Emulsion for injection

Authorised

- Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Limerick/2010, Inactivated
- Equine herpesvirus 1, Inactivated

## Product identification

**Medicine name:**

BioEquin FH, Emulsion for injection

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**Active substance:**

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

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

Equine herpesvirus 1, 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

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

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

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

Equine herpesvirus 1, Inactivated

2.10 log<sub>10</sub> virus neutralising unit(s) / 1.00 Dose

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Horse**

- Meat. 0 day

- Milk. 0 hour

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

QI05AA04

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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:**

Poland

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**Package description:**

Glass Vial

Glass Vial

Glass Vial

Glass Vial

Glass Vial

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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:**

7/10/2015

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**Manufacturing sites for batch release:**

Bioveta a.s.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2466

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**Date of authorisation status change:**

7/10/2015

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**Reference member state:**

Czechia

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**Procedure number:**

CZ/V/0127/001

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**Concerned member states:**

Estonia Hungary Latvia Lithuania Poland Romania Slovakia Slovenia

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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

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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