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

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

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated
5.00 log2 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 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Milk. 0 hour
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI05AL01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Austria

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

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:
Bioveta a.s.

Marketing authorisation date:
29/09/2025

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
842944

Date of authorisation status change:
29/09/2025

Reference member state:
Czechia

Procedure number:
CZ/V/0201/001

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

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Package Leaflet

English (PDF)
Published on: 13/03/2026
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Labelling

English (PDF)
Published on: 13/03/2026
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Summary of Product Characteristics

English (PDF)
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