BioEquin F suspension for injection for horses

Authorised

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

Product identification

Medicine name:
BioEquin F suspension for injection for horses

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

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

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:
QI05AA01

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

Authorisation status:
Valid

Authorised in:
Belgium

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:
2/10/2025

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
BE-V665134

Date of authorisation status change:
2/10/2025

Reference member state:
Czechia

Procedure number:
CZ/V/0200/001

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

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

Documents

Summary of Product Characteristics

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.

Package Leaflet

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