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
BioEquin FH, injekčná emulzia pre kone

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

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
6.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Limerick/2010, Inactivated
6.00 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Equine herpesvirus 1, Inactivated
2.10 log10 virus neutralising unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Slovakia

Package description:

Glass Vial
Glass Vial
Glass Vial

Glass Vial
Glass Vial

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:
18/03/2015

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/009/MR/15-S

Date of authorisation status change:
15/12/2022

Reference member state:
Czechia

Procedure number:
CZ/V/0127/001

Concerned member states:
Estonia Hungary Latvia Lithuania Poland Romania Slovakia Slovenia

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

Documents

Combined File of all Documents

English (PDF)
Published on: 5/09/2022
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