BioEquin FT suspensija injekcijām zirgiem

Authorised

- Clostridium tetani, toxoid
- Influenza A virus, subtype H3N8, strain A/equine/Brno/08, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated

Product identification

Medicine name:
BioEquin FT suspensija injekcijām zirgiem

Active substance:
Clostridium tetani, toxoid

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

Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Clostridium tetani, toxoid
30.00 international unit(s) / 1.00 unit(s)

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

Viena deva (1 ml) satur: ≥ 5 log2 HIT (specifisko antivielu vidējais ģeometriskais titrs jūrascūciņu asins serumā, noteikts ar hemaglutinācijas inhibīcijas testu)
Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated

Viena deva (1 ml) satur: ≥ 5 log2 HIT (specifisko antivielu vidējais ģeometriskais titrs jūrascūciņu asins serumā, noteikts ar hemaglutinācijas inhibīcijas testu)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Horse**
  - 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:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Bioveta a.s.

Marketing authorisation date:
28/04/2000

Manufacturing sites for batch release:
Bioveta a.s.

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/NRP/20/0021

Date of authorisation status change:
1/05/2000

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.

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.