FLUEQUIN T, Injekční suspenze

Authorised

- Influenza A virus, subtype H7N7, strain A/equine/Prague/1956, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Brno/97, Inactivated
- Clostridium tetani, toxoid

Product identification

Medicine name:
FLUEQUIN T, Injekční suspenze

Active substance:
Influenza A virus, subtype H7N7, strain A/equine/Prague/1956, Inactivated
Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated
Influenza A virus, subtype H3N8, strain A/equine/Brno/97, Inactivated
Clostridium tetani, toxoid

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H7N7, strain A/equine/Prague/1956, Inactivated
160.00 haemagglutinating units / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Morava/95, Inactivated
320.00 haemagglutinating units / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Brno/97, Inactivated
320.00 haemagglutinating units / 1.00 Dose

Clostridium tetani, toxoid
1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

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:

Czechia

Package description:

Available only in Czech

Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

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:
22/01/2004

Manufacturing sites for batch release:
Bioveta a.s.

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

Authorisation number:
97/003/04-C

Date of authorisation status change:
23/02/2009

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.