

Equip FT vet. injektioneste, suspensio

Authorised

- Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
- Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated
- Tetanus toxoid

Product identification

Medicine name:

Equip FT vet. injektioneste, suspensio

Active substance:

Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated

Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated

Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated

Tetanus toxoid

Target species:

Horse

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
1.20 log₁₀ haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
2.10 log₁₀ haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated
2.40 log₁₀ haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Tetanus toxoid

70.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Horse

- Not applicable. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AL01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Finland

Available in:

Finland

Package description:

Available only in [Finnish](#)

Available only in [Finnish](#)

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:

Zoetis Animal Health ApS

Marketing authorisation date:

9/06/1999

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

Finnish Medicines Agency

Authorisation number:

13527

Date of authorisation status change:

9/06/1999

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.