

# EQUIP-FT SUSPENSION INYECTABLE PARA CABALLOS

Authorised

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

## Product identification

**Medicine name:**

EQUIP-FT SUSPENSION INYECTABLE PARA CABALLOS

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**Active substance:**

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

Tetanus toxoid

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

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

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**Target species:**

Horse

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**Route of administration:**

Intramuscular use

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## Product details

### Active substance and strength:

Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated  
1.20 haemagglutination inhibiting unit(s) / 1.00 Dose

Tetanus toxoid

70.00 international unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated  
2.10 haemagglutination inhibiting unit(s) / 1.00 Dose

Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated  
20.00 microgram(s) / 1.00 Dose

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### Pharmaceutical form:

Suspension for injection

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### Withdrawal period by route of administration:

#### Intramuscular use:

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#### Horse

- Meat and offal. 0 day

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### Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AL01

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### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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### Authorisation status:

Valid

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### Authorised in:

Spain

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### Package description:

Available only in [Spanish](#)

Available only in [Spanish](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Zoetis Spain S.L.

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**Marketing authorisation date:**

23/02/1999

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

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**Authorisation number:**

1241 ESP

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**Date of authorisation status change:**

1/01/2016

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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.