EQUIP FT SUSPENSION INJECTABLE POUR CHEVAUX ET PONEYS

Authorised

- Equine influenza virus H3N8, A/equine/Borlange/2/91, Inactivated
- Equine influenza virus H7N7, A/equine/Newmarket/1/77, Inactivated
- Clostridium tetani, toxoid
- Equine influenza virus H3N8, A/equine/Kentucky/2/98, Inactivated

Product identification

Medicine name:
EQUIP FT SUSPENSION INJECTABLE POUR CHEVAUX ET PONEYS

Active substance:
Equine influenza virus H3N8, A/equine/Borlange/2/91, Inactivated

Equine influenza virus H7N7, A/equine/Newmarket/1/77, Inactivated

Clostridium tetani, toxoid

Equine influenza virus H3N8, A/equine/Kentucky/2/98, Inactivated

Target species:
Equid

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Equine influenza virus H3N8, A/equine/Borlange/2/91, Inactivated
2.20 log10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Equine influenza virus H7N7, A/equine/Newmarket/1/77, Inactivated
1.20 log10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Clostridium tetani, toxoid
70.00 international unit(s) / 2.00 millilitre(s)

Equine influenza virus H3N8, A/equine/Kentucky/2/98, Inactivated
2.74 log10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Equid**

  - All relevant tissues. 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:

France

Package description:

Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Zoetis France

Marketing authorisation date:
This information is not available for this product.

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
National Veterinary Medicines Agency

Authorisation number:
FR/V/8448928 9/1995

Date of authorisation status change:
17/11/2010

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.
