Equip F Injektionssuspension für Pferde und Ponys

Authorised

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

Product identification

Medicine name:
Equip F Injektionssuspension für Pferde und Ponys

Active substance:
Influenza A virus, subtype H3N8, strain A/equine/Borlange/91, Inactivated
Influenza A virus, subtype H7N7, strain A/equine/Newmarket/77, Inactivated
Influenza A virus, subtype H3N8, strain A/equine/Kentucky/1/98, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

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

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/Kentucky/1/98, Inactivated
2.64 log10 haemagglutination inhibiting unit(s) / 2.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AA01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Zoetis Deutschland GmbH

Marketing authorisation date:
19/07/2003

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.02824.01.1

Date of authorisation status change:
29/07/2008

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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