Equilis Prequenza Te (--) - Suspension for injection

Authorised

- Influenza A virus, subtype H3N8, strain A/equine/South Africa/4/03, Inactivated
- Tetanus toxoid
- Influenza A virus, subtype H3N8, strain A/equine/Newmarket/2/93, Inactivated

Product identification

Medicine name:
Equilis Prequenza Te (--) - Suspension for injection

Active substance:
Influenza A virus, subtype H3N8, strain A/equine/South Africa/4/03, Inactivated

Tetanus toxoid

Influenza A virus, subtype H3N8, strain A/equine/Newmarket/2/93, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Influenza A virus, subtype H3N8, strain A/equine/South Africa/4/03, Inactivated

Presentation_strength:50 AU Reference:II.C.2.1.1 Index:0

Tetanus toxoid

Presentation_strength:40 Lf Index:1

Influenza A virus, subtype H3N8, strain A/equine/Newmarket/2/93, Inactivated

Presentation_strength:50 AU Reference:II.C.2.1.3 Index:2

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Not applicable. 0 day Zero days

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:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Cyprus , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Ireland , Luxembourg , Netherlands , Norway , Portugal , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Pre-filled syringe (glass), Package_size:5 pre-filled syringes, Content:1 dose
Packaging:Pre-filled syringe (glass), Package_size:1 pre-filled syringe, Content:1 dose
Packaging:Pre-filled syringe (glass), Package_size:10 pre-filled syringes, Content:1 dose

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
8/07/2005

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
8/07/2005

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

Documents

Combined File of all Documents

English (PDF)
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