

Equilis Prequenza Te (--) - Suspension for injection

Authorised

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

Product identification

Medicine name:

Equilis Prequenza Te (--) - Suspension for injection

Active substance:

Tetanus toxoid

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

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

Target species:

Horse

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Tetanus toxoid

Presentation_strength:40 Lf Reference:II. Tetanus toxoid Comments:induction of immunity Index:0

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

Presentation_strength:50 AU Reference:II.C.2.1.1.a Comments:induction of immunity Index:1

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

Presentation_strength:50 AU Reference:II.C.2.1.1.b Comments:induction of immunity Index:3

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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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

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 , Germany , Ireland , Italy , Lithuania , Netherlands , Norway , Poland , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Vial (glass), Package_size:10 vials, 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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