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Equilis West Nile (--) - Suspension for injection

Authorised

- Flavivirus, strain YF-WN, expressing preM and E proteins genes of West Nile virus, Inactivated

Product identification

Medicine name:
Equilis West Nile (--) - Suspension for injection

Active substance:
Flavivirus, strain YF-WN, expressing preM and E proteins genes of West Nile virus, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Flavivirus, strain YF-WN, expressing preM and E proteins genes of West Nile virus, Inactivated

Presentation_strength:1400 AU Reference:Hse Index:0

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:

QI05AA10

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 , Germany , Hungary , Italy , Netherlands

Package description:

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

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

Packaging:Vial (glass), Package_size:10 vials, Content:1 ml (1 dose)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
6/06/2013

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:
6/06/2013

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

Documents

Combined File of all Documents

English (PDF)
Published on: 14/03/2026
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