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# Proteq West Nile (--) - Suspension for injection

Authorised

- Canarypox virus, strain vCP2017, expressing preM and E proteins genes of West Nile virus, Live

## Product identification

**Medicine name:**
Proteq West Nile (--) - Suspension for injection

**Active substance:**
Canarypox virus, strain vCP2017, expressing preM and E proteins genes of West Nile virus, Live

**Target species:**
Horse

**Route of administration:**
Intramuscular use

## Product details

**Active substance and strength:**
Canarypox virus, strain vCP2017, expressing preM and E proteins genes of West Nile virus, Live

Presentation_strength:6.0 ≤ R ≤ 7.8 log 10 CCID50 Reference:Hse Comments:Supply of antigen 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:**

QI05AX

**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 , France , Germany , Luxembourg , Spain

**Package description:**

Packaging:Vial (glass), Package_size:10 vials, Content:1 ml (1 dose)
Packaging:Vial (glass), Package_size:5 vials, Content:1 ml (1 dose)
Packaging:Vial (glass), Package_size:2 vials, Content:1 ml (1 dose)
Packaging:Vial (glass), Package_size:1 vial, Content:1 ml (1 dose)

## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**
Full application (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Boehringer Ingelheim Vetmedica GmbH

**Marketing authorisation date:**
5/08/2011

**Manufacturing sites for batch release:**
Boehringer Ingelheim Animal Health France

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
5/08/2011

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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