

Equip WNV (--)- Emulsion for injection

Authorised

- West Nile virus, strain VM-2, Inactivated

Product identification

Medicine name:

Equip WNV (--)- Emulsion for injection

Active substance:

West Nile virus, strain VM-2, Inactivated

Target species:

Horse

Route of administration:

Intramuscular use

Product details

Active substance and strength:

West Nile virus, strain VM-2, Inactivated

Presentation_strength:RP 1.0-2.2 Reference:HSE Index:0

Pharmaceutical form:

Emulsion 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 , Greece , Portugal , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Single-dose pre-filled syringe (glass) with bromobutyl rubber tip,

Package_size:4 syringes, Content:1 ml

Packaging:Single-dose pre-filled syringe (glass) with bromobutyl rubber tip,

Package_size:10 syringes, Content:1 ml

Packaging:Single-dose pre-filled syringe (glass) with bromobutyl rubber tip,

Package_size:2 syringes, Content:1 ml

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:

Zoetis Belgium

Marketing authorisation date:

21/11/2008

Manufacturing sites for batch release:

Zoetis Belgium

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

18/12/2015

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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