

Zulvac BTV BTV 8 - Suspension for injection

Authorised

- Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

Product identification

Medicine name:

Zulvac BTV BTV 8 - Suspension for injection

Active substance:

Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

Target species:

Cattle

Sheep

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

Presentation_strength:7.3-7.4 log10 TCID₅₀ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Cattle

-

Sheep

- Not applicable. 0 day Zero days

Subcutaneous use:

-

Cattle

-

Sheep

- Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA08

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)

Package description:

Packaging:Vial (HDPE), Package_size:1 vial, Content:240 ml (120 doses)

Packaging:Vial (HDPE), Package_size:1 vial, Content:100 ml (50 doses)

Packaging:Vial (HDPE), Package_size:1 vial, Content:20 ml (10 doses)

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:

Zoetis Belgium SA

Marketing authorisation date:

25/04/2017

Manufacturing sites for batch release:

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

25/04/2017

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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