Zulvac BTV BTV 1 - Suspension for injection

Authorised

- Bluetongue virus, serotype 1, strain ALG2006/01 E1, Inactivated

Product identification

Medicine name:
Zulvac BTV BTV 1 - Suspension for injection

Active substance:
Bluetongue virus, serotype 1, strain ALG2006/01 E1, Inactivated

Target species:
Cattle
Sheep

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Bluetongue virus, serotype 1, strain ALG2006/01 E1, Inactivated

Presentation_strength:6.7-7.4 log 10 $TCID_{50}$ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Sheep**
  - Not applicable. 0 day Zero days

Subcutaneous use:

- **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:20 ml (10 doses)

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

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

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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