Bluevac BTV (--) - BTV1+4+8 - Suspension for injection

Authorised

- Bluetongue virus, serotype 1, strain ALG2006/01 E1, Inactivated
- Bluetongue virus, Serotype 4, strain SPA-1/2004, Inactivated
- Bluetongue virus, serotype 8, strain BEL2006/01, Inactivated

Product identification

Medicine name:
Bluevac BTV (--) - BTV1+4+8 - Suspension for injection

Active substance:
Bluetongue virus, serotype 1, strain ALG2006/01 E1, Inactivated
Bluetongue virus, Serotype 4, strain SPA-1/2004, Inactivated
Bluetongue virus, serotype 8, strain BEL2006/01, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

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

Presentation_strength:≥ 9.06 µg/ml Reference:Hse Index:0

Bluetongue virus, Serotype 4, strain SPA-1/2004, Inactivated

Presentation_strength:≥ 22.06 µg/ml Index:5

Bluetongue virus, serotype 8, strain BEL2006/01, Inactivated

Presentation_strength:≥ 245.67 µg/ml Reference:Hse Index:6

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

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:Bottle (HDPE), Package_size:1 bottle, Content:100 ml

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:252 ml
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:52 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:
CZ Vaccines S.A.U.

Marketing authorisation date:
14/04/2011

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
European Commission

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

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: 25/08/2026

Download