

Bluevac BTV (--)- BTV8 - Suspension for injection

Authorised

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

Product identification

Medicine name:

Bluevac BTV (--)- BTV8 - Suspension for injection

Active substance:

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

Target species:

Cattle
Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Bluetongue virus, serotype 8, strain BEL2006/01, Inactivated
Presentation_strength: $\geq 245.67 \mu\text{g/ml}$ Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Not applicable. 0 day Zero days

-

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: 252 ml

Packaging: Bottle (HDPE), Package_size: 1 bottle, Content: 100 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.

Date of authorisation status change:

14/04/2011

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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