

BTVPUR (--) - BTV 1+4 Suspension for injection

Authorised

- Bluetongue virus, serotype 4, Inactivated
- Bluetongue virus, serotype 1, Inactivated

Product identification

Medicine name:

BTVPUR (--) - BTV 1+4 Suspension for injection

Active substance:

Bluetongue virus, serotype 4, Inactivated

Bluetongue virus, serotype 1, Inactivated

Target species:

Cattle

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Bluetongue virus, serotype 4, Inactivated

Presentation_strength: $\geq 1.86 \log_{10}$ pixels Reference:Hse Index:0

Bluetongue virus, serotype 1, Inactivated

Presentation_strength:≥1.9 log10 pixels Reference:Hse Index:1

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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Cattle

- Not applicable. 0 day Zero days

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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 (PP), Package_size:1 bottle, Content:50 ml (50 doses)

Packaging:Bottle (glass), Package_size:1 bottle, Content:10 ml (10 doses)

Packaging:Bottle (PP), Package_size:10 bottles, Content:100 ml (100 doses)

Packaging:Bottle (PP), Package_size:10 bottles, Content:50 ml (50 doses)

Packaging:Bottle (PP), Package_size:1 bottle, Content:100 ml (100 doses)

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:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

17/12/2010

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

12/04/2016

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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