BTVPUR (--) - BTV 2+8 Suspension for injection

Authorised

- Bluetongue virus, serotype 8, Inactivated
- Bluetongue virus, serotype 1, Inactivated

Product identification

Medicine name:
BTVPUR (--) - BTV 2+8 Suspension for injection

Active substance:
Bluetongue virus, serotype 8, Inactivated
Bluetongue virus, serotype 1, Inactivated

Target species:
Cattle
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Bluetongue virus, serotype 8, Inactivated
Presentation_strength:≥2.12 log10 pixels Reference:Hse Index:0

Bluetongue virus, serotype 1, Inactivated

Presentation_strength:≥1.86 log10 pixels Reference:Hse Index:1

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

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

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:50 ml (50 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:
20/08/2018

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: 11/07/2023

Download

ema-puar-btvpur-v-2231-var-ii-0007-g-en.pdf

ema-puar-btvpur-v-2231-par-en.pdf

ema-puar-btvpur-v-2231-var-ii-0010-en.pdf

ema-puar-btvpur-v-2231-var-s-0002-en.pdf