Syvazul BTV BTV 1 - Suspension for injection

Authorised

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

Product identification

Medicine name:
Syvazul 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:RP ≥ 1 Reference:Hse Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI04AA02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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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)

Available in:
Spain

Package description:
Packaging:Vial (PP), Package_size:1 vial, Content:200 ml (100 sheep doses or 50 cattle doses)
Packaging:Vial (PP), Package_size:1 vial, Content:80 ml (40 sheep doses or 20 cattle 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:
Laboratorios Syva S.A.

Marketing authorisation date:
9/01/2019

Manufacturing sites for batch release:
Laboratorios Syva S.A.

Responsible authority:
European Commission

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

Date of authorisation status change:
9/01/2019

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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