# Syvazul BTV BTV 4 - Suspension for injection

Authorised

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

## Product identification

**Medicine name:**
Syvazul BTV BTV 4 - Suspension for injection

**Active substance:**
Bluetongue virus, Serotype 4, strain SPA-1/2004, Inactivated

**Target species:**
Cattle
Sheep

**Route of administration:**
Intramuscular use
Subcutaneous use

## Product details

**Active substance and strength:**
Bluetongue virus, Serotype 4, strain SPA-1/2004, Inactivated

Presentation_strength:RP ≥ 1 Reference:Hse Index:0

**Pharmaceutical form:**

Suspension for injection

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**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

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI04AA02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**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)

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**Available in:**
Greece , Italy , Portugal , Spain

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