File downloaded on 2026-11-08
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000000028

# Zulvac 8 Ovis (--) - Suspension for injection

Authorised

- Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

## Product identification

**Medicine name:**
Zulvac 8 Ovis (--) - Suspension for injection

**Active substance:**
Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

**Target species:**
Sheep

**Route of administration:**
Subcutaneous use

## Product details

**Active substance and strength:**
Bluetongue virus, Serotype 8, strain BTV-8/BEL2006/02, Inactivated

Presentation_strength:RP* ≥1 Reference:Hse Index:0

**Pharmaceutical form:**
Suspension for injection

**Withdrawal period by route of administration:**

**Subcutaneous use:**

- **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

**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:Vial (glass), Package_size:1 vial, Content:100 ml

Packaging:Vial (glass), Package_size:1 vial, Content:240 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:**

Zoetis Belgium

**Marketing authorisation date:**

15/01/2010

**Manufacturing sites for batch release:**
Zoetis Manufacturing & Research Spain S.L.

**Responsible authority:**
European Commission

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

**Date of authorisation status change:**
15/01/2010

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: 13/03/2026
Download

ema-puar-zulvac-8-ovis-v-147-var-s-0009-en.pdf

ema-puar-zulvac-8-ovis-v-147-par-en.pdf

ema-puar-zulvac-8-ovis-v-147-var-ii-0001-en.pdf