File downloaded on 2025-10-23

Source URL: https://medicines.health.europa.eu/veterinary/en/600000044223

# CZV BOVINE TUBERCULIN PPD SOLUTION FOR INJECTION

Authorised

 Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative

# Product identification

#### **Medicine name:**

CZV BOVINE TUBERCULIN PPD SOLUTION FOR INJECTION CZV Bovines Tuberculin PPD

#### **Active substance:**

Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative

#### **Target species:**

Cattle

#### **Route of administration:**

Intradermal use

# **Product details**

## **Active substance and strength:**

Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative 2500.00 international unit(s) / 0.10 millilitre(s)

#### **Pharmaceutical form:**

# Withdrawal period by route of administration: Intradermal use:

\_\_\_\_

#### **Cattle**

- Meat and offal. 0 day

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AR01

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### Authorised in:

Germany

#### **Available in:**

Germany

#### Package description:

cardboard box of 20 doses with 1 vial of 2 ml cardboard box of 200 doses with 10 vials of 2 ml cardboard box of 500 doses with 25 vials of 2 ml cardboard box of 50 doses with 1 vial of 5 ml cardboard box of 500 doses with 10 vials of 5 ml cardboard box of 1250 doses with 25 vials of 5 ml

# 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: CZ Vaccines S.A.U.
Marketing authorisation date: 28/09/2011
Manufacturing sites for batch release: CZ Vaccines S.A.U.
Responsible authority: Paul-Ehrlich-Institut
Authorisation number: PEI.V.11529.01.1
Date of authorisation status change: 21/03/2020
Reference member state: Spain
Procedure number: ES/V/0180/001
Concerned member states: Belgium Bulgaria France Germany Greece Hungary Ireland Italy Portugal Romania United Kingdom (Northern Ireland)
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Summary of Product Characteristics

Package Leaflet
Labelling
Combined File of all Documents
English (PDF)
Published on: 24/07/2025
<u>Download</u>