

# 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

**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:**

Solution for injection

**Withdrawal period by route of administration:****Intradermal use:**

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**Cattle**

- Meat and offal. 0 day

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

QI02AR01

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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:**

France

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**Available in:**

France

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

CZ Vaccines S.A.U.

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**Marketing authorisation date:**

29/08/2011

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**Manufacturing sites for batch release:**

CZ Vaccines S.A.U.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5653366 6/2011

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**Date of authorisation status change:**

16/09/2016

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0180/001

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**Concerned member states:**

Belgium Bulgaria France Germany Greece Hungary Ireland Italy Portugal  
Romania United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

English (PDF)

Published on: 11/04/2023

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Package Leaflet

Labelling

Combined File of all Documents

Package Leaflet and Labelling

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