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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:

- **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:
Spain

Available in:
Spain

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:
25/10/1982

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3246 ESP

Date of authorisation status change:
18/05/2015

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 www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 14/03/2026
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Package Leaflet

English (PDF)
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Labelling

English (PDF)
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Combined File of all Documents

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