

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AR01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Available in:

United Kingdom (Northern Ireland)

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:

7/12/2011

Manufacturing sites for batch release:

CZ Vaccines S.A.U.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 30824/4001

Date of authorisation status change:

21/10/2024

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

Package Leaflet

Labelling

Combined File of all Documents