

BOVINI TUBERKULIN PPD 28 000 i.j., otopina za injekciju za goveda i svinje

Authorised

- Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative

Product identification

Medicine name:

BOVINI TUBERKULIN PPD 28 000 i.j., otopina za injekciju za goveda i svinje

Active substance:

Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative

Target species:

Cattle

Pig

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative
28000.00 international unit(s) / 1.00 millilitre(s)

Solution for injection

Subcutaneous use:

-

- Meat and offal. 0 day

- Milk. 0 day

-

- Meat and offal. 0 day

QV04CF01

Veterinary medicinal product subject to veterinary prescription

Valid

Croatia

Available only in Croatian

Available only in Croatian

Available only in Croatian

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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:

Genera d.d.

Marketing authorisation date:

26/05/2020

Manufacturing sites for batch release:

Genera d.d.

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/20-01/284

Date of authorisation status change:

30/04/2025

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.