

BIOBOVITUBAL 28000 solutie injectabila pentru bovine, ovine, caprine, porcine, cabaline, caini

Authorised

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

Product identification

Medicine name:

BIOBOVITUBAL 28000 solutie injectabila pentru bovine, ovine, caprine, porcine, cabaline, caini

Active substance:

Mycobacterium bovis, strain AN5, bovine tuberculin purified protein derivative

Target species:

Cattle
Sheep
Goat
Horse
Dog
Pig

Route of administration:

Intradermal 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)

Pharmaceutical form:

Solution for injection

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

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Cattle

- Milk. 0 day
- Meat and offal. 0 day

•

Sheep

- Milk. 0 day
- Meat and offal. 0 day

•

Goat

- Milk. 0 day
- Meat and offal. 0 day

•

Horse

- Meat and offal. 0 day

•

Pig

- 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:

Romania

Package description:

Available only in Romanian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:

Bioveta Romania S.R.L.

Marketing authorisation date:

21/05/2017

Manufacturing sites for batch release:

Bioveta, a.s.

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

230092

Date of authorisation status change:

23/05/2023

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.