

FOR ANIMAL TREATMENT ONLY
RESTRICTED VETERINARY
MEDICINE



TUBERCULIN

25,000 IU per mL

Registered to:ASUREQuality Limited,
Level 1, 7a Pacific Rise, Mt Wellington, 1060
New Zealand
Tel: +64 9 573 8000
Fax: +64 9 573 8118

Registered pursuant to ACVM Act 1997
No. A10883.

See www.foodsafety.govt.nz for
registration conditions



STORE AT 2-8 DEGREES CELSIUS
DO NOT FREEZE
PROTECT FROM LIGHT
STORE OUT OF THE REACH OF CHILDREN

PA2500AU-01-May2021

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TUBERCULIN

25,000 IU per mL

Active Ingredients: Purified Protein Derivative
of Mycobacterium avium, strain D4ER,
25,000 IU/mL.

Indications: For use as an aid in the diagnosis
of mycobacterial infection in cattle.

Directions for use: For use under the authority
or prescription of a veterinary surgeon by
persons approved under the Australian
Biosecurity Act 2015 and Australian Export
Control Act 1982 when applying cattle
tuberculin tests. Shake well before use.

Test Application

The following instructions are guidelines only:
Dose: 0.1mL intra-dermally. It is recommended
syringes are calibrated to 0.1 mL and the
needle of the syringe is 22-26 gauge and 3-4
mm long. Syringes should be kept clean and
sterilized. Contamination of the syringe with
disinfectants or alcohol may interfere with
testing. When filling syringes, care must be
taken to prevent contamination of vials.

Preferred site of injection:

Cattle: the mid cervical site as a comparative
cervical test (CCT);

When using the cervical sites, care is required
to evenly clip hair close to the skin surface
(2mm or less mean length) prior to injecting.
The recommended size of the clipped area is
10 x 10 cm for each injection site.

When applying a CCT use Observe™ Bovine
Tuberculin 30,000 IU per mL in conjunction with
Observe™ Avian Tuberculin 25,000 IU per mL.

Test Interpretation:

It is recommended the test be read 72 hours
after injection.

Skin test reactions are to be classified in
accordance with the requirements of OSPRI, the
TBfree New Zealand programme and the

National Operational Plan pursuant to the
Biosecurity Act 1993 and the Biosecurity
(National Bovine Tuberculosis Pest Management
Plan) Order 1998 (the Order).

Retesting: To avoid desensitization, a retest
using bovine and/or avian PPD tuberculin
should not be applied until 60 days after any
previous tuberculin test.

Product Information

Withholding period: Nil

Storage: Refrigerate (2°C – 8°C). Do not freeze.

Keep the vials in the closed box in order to
protect from light.

Transportation: May be transported at 2°C –
37°C for a period not longer than 14 days.
Do not freeze.

Use: Do not use after the expiry date. Use
immediately after opening. Discard unused
contents. Do not mix with other vaccines or
immunological products.

Method of Disposal: Unused, part-used and
empty vials should be destroyed according to
local regulations.

Safety Precautions: In the case of accidental
self-injection, an area of intense local irritation
may develop for up to 96 hours later. If irritation
occurs, you are advised to consult your medical
practitioner.

Store out of reach of children.

Manufactured by: Prionics Lelystad B.V.,
Platinastraat 33, 8211 AR Lelystad,
The Netherlands

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