

Tuberculin PPD Kit

Authorised

- TUBERCULIN PURIFIED PROTEIN DERIVATIVE, BOVINE
- TUBERCULIN PURIFIED PROTEIN DERIVATIVE, AVIAN

Product identification

Medicine name:

Tuberculin PPD Kit

Active substance:

TUBERCULIN PURIFIED PROTEIN DERIVATIVE, BOVINE

TUBERCULIN PURIFIED PROTEIN DERIVATIVE, AVIAN

Target species:

Cattle

Route of administration:

Intradermal use

Product details

Active substance and strength:

TUBERCULIN PURIFIED PROTEIN DERIVATIVE, BOVINE

30000.00 international unit(s) / 0.10 millilitre(s)

TUBERCULIN PURIFIED PROTEIN DERIVATIVE, AVIAN

25000.00 international unit(s) / 0.10 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intradermal use:

-

Cattle

- Milk. 0 day

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AR01

QI02AR02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

The 20-doses presentation of the Tuberculin PPD Kit: Polystyrene box with 20 vials (hydrolytic Type I) of Bovine Tuberculin PPD 3000 and 20 vials (hydrolytic Type I) of Avian Tuberculin PPD 2500. Each vial contains 20 doses of 0.1 ml.

The 50-doses presentation of the Tuberculin PPD Kit: Polystyrene box with 10 vials (hydrolytic Type I) of Bovine Tuberculin PPD 3000 and 10 vials (hydrolytic Type I) of Avian Tuberculin PPD 2500. Each vial contains 50 doses of 0.1 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:

Prionics Lelystad B.V.

Marketing authorisation date:

20/04/2021

Manufacturing sites for batch release:

Prionics Lelystad B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1007311 1/2021

Date of authorisation status change:

20/04/2021

Reference member state:

Netherlands

Procedure number:

NL/V/0322/001

Concerned member states:

France Hungary Ireland United Kingdom (Northern Ireland)

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

Documents

Labelling

Summary of Product Characteristics

English (PDF)

Published on: 13/06/2022

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Package Leaflet and Labelling

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

Combined File of all Documents

tuberculin pil.pdf

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