Tuberculin PPD Kit

Authorised

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

Product identification

Medicine name:
Tuberculin PPD Kit
Tuberculine PPD Kit

Active substance:
TUBERCULIN PURIFIED PROTEIN DERIVATIVE, AVIAN
TUBERCULIN PURIFIED PROTEIN DERIVATIVE, BOVINE

Target species:
Cattle

Route of administration:
Intradermal use

Product details

Active substance and strength:
TUBERCULIN PURIFIED PROTEIN DERIVATIVE, AVIAN
25000.00 international unit(s) / 0.10 millilitre(s)
TUBERCULIN PURIFIED PROTEIN DERIVATIVE, BOVINE
30000.00 international unit(s) / 0.10 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intradermal use:

- **Cattle**
  - Milk. no withdrawal period zero days
  - Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AR01

QI02AR02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

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.

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.

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:
18/11/2010

Manufacturing sites for batch release:
Prionics Lelystad B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 105356

Date of authorisation status change:
31/01/2022

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

Combined File of all Documents

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

Labelling

Summary of Product Characteristics

tuberculin pil.pdf

tuberculin par.pdf
