Terivac

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Live

Product identification

Medicine name:
Terivac

Active substance:
Turkey rhinotracheitis virus, strain VCO3, Live

Target species:
Turkey

Route of administration:
Nebulisation use
In drinking water/milk use
In drinking water use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain VCO3, Live
199.53 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oral suspension

Withdrawal period by route of administration:

Nebulisation use:

- **Turkey**
  - Meat and offal. 0 day

In drinking water/milk use:

- **Turkey**
  - Meat and offal. 0 day

In drinking water use:

- **Turkey**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01CD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Germany

Package description:

Available only in German
Available only in German
Available only in German
Available only in German
Available only in German
Available only in German

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
18/11/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France SCS

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
106a/90

Date of authorisation status change:
19/06/2008

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.