

POULVAC TRT LYOPHILISATE FOR OCULONASAL SUSPENSION FOR TURKEYS

Authorised

- Turkey rhinotracheitis virus, strain Clone K, Live

Product identification

Medicine name:

POULVAC TRT LYOPHILISATE FOR OCULONASAL SUSPENSION FOR TURKEYS

Active substance:

Turkey rhinotracheitis virus, strain Clone K, Live

Target species:

Turkey

Route of administration:

Nasal use

Ocular use

Nebulisation use

Product details

Active substance and strength:

Turkey rhinotracheitis virus, strain Clone K, Live

3.20 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension

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

-

Turkey

- All relevant tissues. 0 day

Ocular use:

-

Turkey

- All relevant tissues. 0 day

Nebulisation use:

-

Turkey

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01CD01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Germany

Available in:

Germany

Package description:

Cardboard boxes containing 1 vial of 1,000 doses

Cardboard boxes containing 1 vial of 2,000 doses

Cardboard boxes containing 1 vial of 5,000 doses

Cardboard boxes containing 10 vials of 1,000 doses
Cardboard boxes containing 10 vials of 2,000 doses
Cardboard boxes containing 10 vials of 5,000 doses

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis Deutschland GmbH

Marketing authorisation date:

5/12/2000

Manufacturing sites for batch release:

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

PEI.V.01721.01.1

Date of authorisation status change:

13/12/2011

Reference member state:

France

Procedure number:

FR/V/0412/001

Concerned member states:

Austria Belgium Denmark Germany Greece Italy Netherlands Portugal
Spain 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

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