AVIFFA-RTI LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA

Authorised

- Turkey rhinotracheitis virus, strain VCO3, Inactivated

Product identification

Medicine name:
AVIFFA-RTI LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA

Active substance:
Turkey rhinotracheitis virus, strain VCO3, Inactivated

Target species:
Turkey
Chicken (pullet for egg production, future layer)
Future breeder pullet

Route of administration:
Nebulisation use
Ocular use
In drinking water use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain VCO3, Inactivated
4.00 50% cell culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for suspension

Withdrawal period by route of administration:
Nebulisation use:

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

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day

- **Future breeder pullet**
  - Meat and offal. 0 day

Ocular use:

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

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day

- **Future breeder pullet**
  - Meat and offal. 0 day

In drinking water use:

- **Turkey**
  - Meat and offal. 0 day
- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day
- **Future breeder pullet**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish
Available only in Spanish

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:
Boehringer Ingelheim Animal Health Espana S.A.

Marketing authorisation date:
16/07/1991

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2365 ESP

Date of authorisation status change:
7/09/2011

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

Documents

Summary of Product Characteristics

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Labelling

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

Package Leaflet

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