

NOBILIS MA5+CLONE 30 LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA PARA POLLOS

Authorised

- Newcastle disease virus, strain Clone 30, Live
- Infectious bronchitis virus, type Massachusetts, strain Ma5, Live

Product identification

Medicine name:

NOBILIS MA5+CLONE 30 LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA PARA POLLOS

Active substance:

Newcastle disease virus, strain Clone 30, Live

Infectious bronchitis virus, type Massachusetts, strain Ma5, Live

Target species:

Chicken

Route of administration:

Nebulisation use

In drinking water use

Oculonasal use

Product details

Active substance and strength:

Newcastle disease virus, strain Clone 30, Live
10.00 50% Embryo Infective Dose / 1.00 Dose

Infectious bronchitis virus, type Massachusetts, strain Ma5, Live
10.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Nebulisation use:

-

Chicken

- Meat and offal. 0 day
- Egg. 0 day

In drinking water use:

-

Chicken

- Meat and offal. 0 day
- Egg. 0 day

Oculonasal use:

-

Chicken

- Meat and offal. 0 day
 - Egg. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD11

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

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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:

Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:

24/10/2013

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

2906 ESP

Date of authorisation status change:

24/10/2013

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.

Package Leaflet

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

Labelling

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