HIPRAVIAR-S LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA PARA POLLOS

Authorised

- Newcastle disease virus, strain La Sota, Inactivated

Product identification

Medicine name:
HIPRAVIAR-S LIOFILIZADO PARA SUSPENSION OCULONASAL Y PARA ADMINISTRACION EN AGUA DE BEBIDA PARA POLLOS

Active substance:
Newcastle disease virus, strain La Sota, Inactivated

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

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

Product details

Active substance and strength:
Newcastle disease virus, strain La Sota, Inactivated
8.50 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for suspension

Withdrawal period by route of administration:
Nebulisation use:

- **Chicken (broiler)**
  - 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:

- **Chicken (broiler)**
  - 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

Oculonasal use:

- **Chicken (broiler)**
  - 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:
QI01AD06

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

Authorisation status:
Valid

Authorised 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
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:
Laboratorios Hipra S.A.

Marketing authorisation date:
27/07/1972

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2244 ESP

Date of authorisation status change:
16/02/2011

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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