**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000059066

## **AVIPRO ND LASOTA**

Authorised

• Newcastle disease virus, strain La Sota, Inactivated

## Product identification

#### **Medicine name:**

**AVIPRO ND LASOTA** 

#### **Active substance:**

Newcastle disease virus, strain La Sota, Inactivated

#### **Target species:**

Chicken (chick, for replacement) Chicken (broiler)

#### Route of administration:

In drinking water use Intraocular use Nebulisation use

## **Product details**

#### **Active substance and strength:**

Newcastle disease virus, strain La Sota, Inactivated 6.00 50% Embryo Infective Dose / 1.00 Dose

#### **Pharmaceutical form:**

Lyophilisate for suspension

# Withdrawal period by route of administration: In drinking water use: **Chicken (chick, for replacement)** - Meat and offal. 0 day **Chicken (broiler)** - Meat and offal. 0 day Intraocular use: **Chicken (chick, for replacement)** - Meat and offal. 0 day **Chicken (broiler)** - Meat and offal. 0 day **Nebulisation use: Chicken (chick, for replacement)** - Meat and offal. 0 day **Chicken (broiler)** - 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:

#### 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:

Elanco GmbH

## Marketing authorisation date:

11/08/2016

## Manufacturing sites for batch release:

Lohmann Animal Health GmbH

## **Responsible authority:**

Spanish Agency Of Medicines And Medical Devices

#### **Authorisation number:**

3461 FSP

## Date of authorisation status change:

To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>

#### **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.