

# PRIMUM IB-ND DUO Lyophilisate for suspension for chickens

Authorised

- Infectious bronchitis virus, type Massachusetts, strain H120, Live
- Newcastle disease virus, strain B1 Hitchner, Live

## Product identification

**Medicine name:**

PRIMUM IB-ND DUO Lyophilisate for suspension for chickens

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**Active substance:**

Infectious bronchitis virus, type Massachusetts, strain H120, Live  
Newcastle disease virus, strain B1 Hitchner, Live

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**Target species:**

Chicken

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**Route of administration:**

Oculonasal use

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## Product details

**Active substance and strength:**

Infectious bronchitis virus, type Massachusetts, strain H120, Live  
3.00 log 10 50% embryo infective dose / 1.00 Dose

Newcastle disease virus, strain B1 Hitchner, Live  
6.00 log<sub>10</sub> 50% embryo infective dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for ocular nasal suspension

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**Withdrawal period by route of administration:**

**Ocular nasal use:**

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**Chicken**

- Meat and offal. 0 day
  - Egg. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD11

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Package description:**

(ID3): 1 Box with 1 Bottle (Glass) with 2000 Dose (2000 Dose)

(ID4): 1 Box with 10 Bottle (Glass) with 2000 Dose (20000 Dose)

(ID2): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)

(ID1): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Laboratorios Calier S.A.

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**Marketing authorisation date:**

9/09/2019

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**Manufacturing sites for batch release:**

Laboratorios Calier S.A.

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

2906

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**Date of authorisation status change:**

9/09/2019

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0274/001

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**Concerned member states:**

Italy Poland Portugal Spain

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 5/11/2024

Updated on: 14/03/2026

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### Package Leaflet

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### Labelling

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