

# Nobilis IB Primo QX (--) lyophilisate and solvent for oculonasal suspension

Authorised

- Infectious bronchitis virus, type QX, strain D388, Live

## Product identification

**Medicine name:**

Nobilis IB Primo QX (--)  
lyophilisate and solvent for oculonasal suspension

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**Active substance:**

Infectious bronchitis virus, type QX, strain D388, 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 QX, strain D388, Live

Presentation\_strength:  $10^{4.0}$  -  $10^{5.5}$  EID<sub>50</sub> Reference: 2.C.2.1 Index: 0

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**Pharmaceutical form:**

Lyophilisate and solvent for ocular suspension

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**Withdrawal period by route of administration:**

**Ocular use:**

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**Chicken**

- Not applicable. 0 day Zero days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AD07

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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:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging:Lyophilisate: cup (Alu/PP); Solvent: vial (LDPE), Package\_size:In cardboard boxes: Lyophilisate:10 cups; Solvent:10 vials, Content:Lyophilisate:1000 doses; Solvent: 35 ml

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

4/09/2014

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

4/09/2014

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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

Combined File of all Documents

English (PDF)

Published on: 2/05/2024

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