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# 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

**Active substance:**
Infectious bronchitis virus, type QX, strain D388, Live

**Target species:**
Chicken

**Route of administration:**
Oculonasal use

## Product details

**Active substance and strength:**
Infectious bronchitis virus, type QX, strain D388, Live

Presentation_strength:$10^{4.0}$ – $10^{5.5}$ $EID_{50}$ Reference:2.C.2.1 Index:0

**Pharmaceutical form:**

Lyophilisate and solvent for oculonasal suspension

**Withdrawal period by route of administration:**

**Oculonasal use:**

- **Chicken**
  - Not applicable. 0 day Zero days

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

QI01AD07

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**

Valid

**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)

**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

## 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:**

Intervet International B.V.

**Marketing authorisation date:**
4/09/2014

**Manufacturing sites for batch release:**
Intervet International B.V.

**Responsible authority:**
European Commission

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
4/09/2014

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

## Documents

Combined File of all Documents

English (PDF)
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