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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