

NOBILIS MA 5 + CLONE 30 LYOPHILISAT POUR SUSPENSION OCULONASALE / ADMINISTRATION DANS L'EAU DE BOISSON POUR POULES

Authorised

- Infectious bronchitis virus, type Massachusetts, strain Ma5, Live
- Newcastle disease virus, strain Clone 30, Live

Product identification

Medicine name:

NOBILIS MA 5 + CLONE 30 LYOPHILISAT POUR SUSPENSION OCULONASALE /
ADMINISTRATION DANS L'EAU DE BOISSON POUR POULES

Active substance:

Infectious bronchitis virus, type Massachusetts, strain Ma5, Live
Newcastle disease virus, strain Clone 30, Live

Target species:

Chicken (one day-old chick)

Route of administration:

Nebulisation use
Oral use

Oculonasal use

Product details

Active substance and strength:

Infectious bronchitis virus, type Massachusetts, strain Ma5, Live
3.00 log 10 50% embryo infective dose / 1.00 Dose

Newcastle disease virus, strain Clone 30, Live
6.00 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Nebulisation use:

-

Chicken (one day-old chick)

- All relevant tissues. 0 day

Oral use:

-

Chicken (one day-old chick)

- All relevant tissues. 0 day

Oculonasal use:

-

Chicken (one day-old chick)

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD11

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Available only in [French](#)

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Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet

Marketing authorisation date:

31/07/2000

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0323807 5/2000

Date of authorisation status change:

31/07/2010

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

Documents

Summary of Product Characteristics

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

Package Leaflet and Labelling

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