

# 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

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### **Active substance:**

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

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### **Target species:**

Chicken (one day-old chick)

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### **Route of administration:**

Nebulisation use  
Oral use

Oculonasal use

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## Product details

### **Active substance and strength:**

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

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

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### **Pharmaceutical form:**

Lyophilisate for oculonasal suspension/use in drinking water

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### **Withdrawal period by route of administration:**

#### **Nebulisation use:**

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#### **Chicken (one day-old chick)**

- All relevant tissues. 0 day

#### **Oral use:**

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#### **Chicken (one day-old chick)**

- All relevant tissues. 0 day

#### **Oculonasal use:**

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#### **Chicken (one day-old chick)**

- All relevant tissues. 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:**

France

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**Available in:**

France

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**Package description:**

Available only in French

Available only in French

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Available only in French

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Intervet

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**Marketing authorisation date:**

31/07/2000

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/0323807 5/2000

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**Date of authorisation status change:**

31/07/2010

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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

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.