

# Nobilis MS Live, lyophilisate for ocular suspension for chickens

Not  
authorised

- Mycoplasma synoviae, strain MS1, Live

## Product identification

**Medicine name:**

Nobilis MS Live, lyophilisate for ocular suspension for chickens

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**Active substance:**

Mycoplasma synoviae, strain MS1, Live

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**Target species:**

Chicken

Chicken (for reproduction)

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**Route of administration:**

Ocular use

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## Product details

**Active substance and strength:**

Mycoplasma synoviae, strain MS1, Live

100.00 million colony forming units / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate for ocular suspension

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**Withdrawal period by route of administration:****Ocular use:**

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**Chicken**

- Egg. 0 day
- Meat and offal. 0 day

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**Chicken (for reproduction)**

- Egg. 0 day
  - Meat and offal. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AE03

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Belgium

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**Package description:**

- (ID6): 1 Box with 10 Bottle (Glass) with 2000 Dose (20000 Dose)
  - (ID5): 1 Box with 10 Bottle (Glass) with 1000 Dose (10000 Dose)
  - (ID4): 1 Box with 10 Bottle (Glass) with 500 Dose (5000 Dose)
  - (ID3): 1 Box with 1 Bottle (Glass) with 2000 Dose (2000 Dose)
  - (ID2): 1 Box with 1 Bottle (Glass) with 1000 Dose (1000 Dose)
  - (ID1): 1 Box with 1 Bottle (Glass) with 500 Dose (500 Dose)
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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet International B.V.

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**Marketing authorisation date:**

27/06/2014

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Federal Agency For Medicines And Health Products

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**Authorisation number:**

BE-V459671

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**Date of authorisation status change:**

27/10/2023

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0260/001

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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

English (PDF)

Published on: 28/01/2022

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### Package Leaflet

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