

Nobilis Ma5 + Clone30, lyofilisaat en suspenseervloeistof voor oculonasale suspensie/gebruik in drinkwater voor kippen

Authorised

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

Product identification

Medicine name:

Nobilis Ma5 + Clone30, lyofilisaat en suspenseervloeistof voor oculonasale suspensie/gebruik in drinkwater voor kippen

Active substance:

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

Target species:

Chicken

Route of administration:

In drinking water use
Oculonasal use
Nebulisation use

Product details

Active substance and strength:

Infectious bronchitis virus, type Massachusetts, strain Ma5, Live

3.00 log₁₀ 50% embryo infective dose / 1.00 Dose

Newcastle disease virus, strain Clone 30, Live

6.00 log₁₀ 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

In drinking water use:

-

Chicken

- Meat and offal. 7 day

- Egg. no withdrawal period
zero days

Oculonasal use:

-

Chicken

- Meat and offal. 7 day

- Egg. no withdrawal period
zero days

Nebulisation use:

-

Chicken

- Meat and offal. 7 day

- Egg. no withdrawal period
zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD11

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

Available only in [Dutch](#)

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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 Nederland B.V.

Marketing authorisation date:

15/12/1994

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 8104

Date of authorisation status change:

26/03/2021

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

Documents

Combined File of all Documents

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