

AviGate ND QV4 lyophilisate for oculonasal suspension/use in drinking water for chickens

Authorised

- Newcastle disease virus, strain QV4, Live

Product identification

Medicine name:

AviGate ND QV4 lyophilisate for oculonasal suspension/use in drinking water for chickens

Active substance:

Newcastle disease virus, strain QV4, Live

Target species:

Chicken

Route of administration:

Oculonasal use
In drinking water use

Product details

Active substance and strength:

Newcastle disease virus, strain QV4, Live
6.00 log 10 50% embryo infective dose / 1.00 Dose

Pharmaceutical form:

Lyophilisate for ocular nasal suspension/use in drinking water

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

-

Chicken

- All relevant tissues. 0 day

In drinking water use:

-

Chicken

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Carton box with 10 glass vials (type I) containing 1 000 doses, closed with a rubber stopper and sealed with an aluminium cap.

Carton box with 10 glass vials (type I) containing 2 500 doses, closed with a rubber stopper and sealed with an aluminium cap.

Carton box with 10 glass vials (type I) containing 5 000 doses, closed with a rubber stopper and sealed with an aluminium cap.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

HuVepharma

Marketing authorisation date:

5/04/2026

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

National Organization For Medicines

Authorisation number:

42128/06-04-2026/K-0265401

Date of authorisation status change:

15/01/2026

Reference member state:

Netherlands

Procedure number:

NL/V/0435/001

Concerned member states:

Austria Belgium Bulgaria Czechia Greece Italy Poland Portugal Romania
Slovakia Spain

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.