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 oculonasal suspension/use in drinking water

Withdrawal period by route of administration:

Oculonasal 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:
Poland

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:
27/04/2026

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3489

Date of authorisation status change:
27/04/2026

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.

Package Leaflet

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

Labelling

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