# Nobilis gumboro D78 Vet. pulver til opløsning i drikkevand 10.000 TCID50/dosis

Authorised

- Infectious bursal disease virus, strain D78, Inactivated

## Product identification

#### Medicine name:

Nobilis gumboro D78 Vet. 10.000 TCID50/dosis lyofilisat til anvendelse i drikkevand Nobilis gumboro D78 Vet. pulver til opløsning i drikkevand 10.000 TCID50/dosis

#### **Active substance:**

Infectious bursal disease virus, strain D78, Inactivated Infectious bursal disease virus, strain D78, Inactivated

Target species:

Poultry

**Route of administration:** In drinking water/milk use

# Product details

## Active substance and strength:

Infectious bursal disease virus, strain D78, Inactivated 10000.00 tissue culture infective dose 50 / 1.00 Dose Infectious bursal disease virus, strain D78, Inactivated 10000.00 tissue culture infective dose 50 / 1.00 Dose Infectious bursal disease virus, strain D78, Inactivated 10000.00 tissue culture infective dose 50 / 1.00 Dose Infectious bursal disease virus, strain D78, Inactivated 10000.00 tissue culture infective dose 50 / 1.00 Dose Infectious bursal disease virus, strain D78, Inactivated 10000.00 tissue culture infective dose 50 / 1.00 Dose

10000.00 tissue culture infective dose 50 / 1.00 Dose

### Pharmaceutical form:

Lyophilisate for use in drinking water

## Withdrawal period by route of administration: In drinking water/milk use:

### Poultry

- Meat and offal. 0 day

## Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA01

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### Authorisation status:

Valid

Authorised in: Denmark

#### Package description:

Available only in <u>Danish</u> Available only in <u>Danish</u> Available only in <u>Danish</u> Available only in <u>Danish</u> Available only in <u>Danish</u>

# Additional information

#### **Entitlement type:**

Marketing Authorisation

#### Legal basis of product authorisation:

Legal basis reviewed according to Acquis communautaire

#### Marketing authorisation holder:

Intervet International B.V.

#### Marketing authorisation date:

17/09/1999

#### Manufacturing sites for batch release:

Intervet International B.V.

#### **Responsible authority:**

**Danish Medicines Agency** 

# Authorisation number: 14933

# **Date of authorisation status change:** 17/09/1999

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

## Documents

Summary of Product Characteristics

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