

# NOBILIS GUMBORO INAC EMULSION INJECTABLE POUR POULES

Authorised

- Infectious bursal disease virus, strain D78, Inactivated

## Product identification

**Medicine name:**

NOBILIS GUMBORO INAC EMULSION INJECTABLE POUR POULES

---

**Active substance:**

Infectious bursal disease virus, strain D78, Inactivated

---

**Target species:**

Chicken (layer hen)

Chicken (for reproduction)

---

**Route of administration:**

Intramuscular use

Subcutaneous use

---

## Product details

**Active substance and strength:**

Infectious bursal disease virus, strain D78, Inactivated

14.50 log<sub>2</sub> serum neutralising unit(s) / 1.00 Dose

---

**Pharmaceutical form:**

Emulsion for injection

---

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

- 

**Chicken (layer hen)**

- Meat and offal. 0 day
- Eggs. 0 day

- 

**Chicken (for reproduction)**

- Meat and offal. 0 day
- Eggs. 0 day

**Subcutaneous use:**

- 

**Chicken (layer hen)**

- Meat and offal. 0 day
- Eggs. 0 day

- 

**Chicken (for reproduction)**

- Meat and offal. 0 day
- Eggs. 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:**

France

---

**Available in:**

France

---

**Package description:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

---

**Marketing authorisation holder:**

Intervet

---

**Marketing authorisation date:**

10/08/1981

---

**Manufacturing sites for batch release:**

Intervet International B.V.

---

**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

---

**Authorisation number:**

FR/V/4406302 1/1981

---

**Date of authorisation status change:**

10/08/2011

---

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

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

### Package Leaflet and Labelling

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