File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/700000188295

POULVAC POULINAC ND+IB M41+EDS EMULSION INJECTABLE POUR POULETS

Authorised

- Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
- Newcastle disease virus, strain Ulster 2C, Inactivated
- Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated

Product identification

Medicine name:
POULVAC POULINAC ND+IB M41+EDS EMULSION INJECTABLE POUR POULETS

Active substance:
Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Newcastle disease virus, strain Ulster 2C, Inactivated

Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated

Target species:
Chicken (layer hen)
Chicken (for reproduction)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Infectious bronchitis virus, type Massachusetts, strain M41, Inactivated
3.30 log10 ELISA unit / 1.00 Dose

Newcastle disease virus, strain Ulster 2C, Inactivated
24.00 antigen unit(s) / 1.00 Dose

Eggdrop syndrome-1976 virus, strain McFerran 127, Inactivated
9.20 log2 haemagglutination inhibiting unit(s) / 1.00 Dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (layer hen)**
  - All relevant tissues. 0 day

- **Chicken (for reproduction)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AA13

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in French

Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Zoetis France

Marketing authorisation date:
10/09/2025

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/1479875 9/2025

Date of authorisation status change:
10/09/2025

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

Documents

Package Leaflet and Labelling

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

Summary of Product Characteristics

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

fr-puar-700000188295-np-rpe919-fr.pdf