File downloaded on 2026-11-25
Source URL: https://medicines.health.europa.eu/veterinary/en/600000082829

NEO-VAKY DNT EMULSION INYECTABLE PARA PORCINO

Authorised

- Pasteurella multocida, serogroup D, strain CECT 4325, toxoid
- Bordetella bronchiseptica, strain BB4-78, Inactivated
- Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated

Product identification

Medicine name:
NEO-VAKY DNT EMULSION INYECTABLE PARA PORCINO

Active substance:
Pasteurella multocida, serogroup D, strain CECT 4325, toxoid
Bordetella bronchiseptica, strain BB4-78, Inactivated
Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated

Target species:
Pig (piglet)
Pig (pregnant sow)
Pig (sow, multipar)

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Pasteurella multocida, serogroup D, strain CECT 4325, toxoid
8.00 haemagglutinating units / 1.00 dose

Bordetella bronchiseptica, strain BB4-78, Inactivated
7.00 haemagglutinating units / 1.00 dose

Pasteurella multocida, serogroup D, strain CECT 4325, Inactivated
6.00 enzyme-linked immunosorbent assay unit / 1.00 dose

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig (piglet)**
  - Meat and offal. 0 day

- **Pig (pregnant sow)**
  - Meat and offal. 0 day

- **Pig (sow, multipar)**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI09AB04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Ovejero S.A.U.

Marketing authorisation date:
18/10/1978

Manufacturing sites for batch release:
Laboratorios Ovejero S.A.U.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3347 ESP

Date of authorisation status change:
22/12/2015

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.