NOBILIS RISMAVAC CONCENTRADO Y DISOLVENTE PARA SUSPENSION INYECTABLE PARA POLLOS

Authorised

- Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live

Product identification

Medicine name:
NOBILIS RISMAVAC CONCENTRADO Y DISOLVENTE PARA SUSPENSION INYECTABLE PARA POLLOS

Active substance:
Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live

Target species:
Chicken (broiler)
Chicken (pullet for egg production, future layer)
Future breeder pullet

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live 10.00 50% cell culture infectious dose / 0.20 millilitre(s)

Pharmaceutical form:
Suspension and solvent for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day

- **Future breeder pullet**
  - Meat and offal. 0 day

Subcutaneous use:

- **Chicken (broiler)**
  - Meat and offal. 0 day

- **Chicken (pullet for egg production, future layer)**
  - Meat and offal. 0 day

- **Future breeder pullet**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD03

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:
Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:
27/02/1981

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
3334 ESP

Date of authorisation status change:

29/01/2016

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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