Nobilis Rismavac Suspensie en oplosmiddel voor suspensie voor injectie

Authorised

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

Product identification

Medicine name:
Nobilis Rismavac Suspensie en oplosmiddel voor suspensie voor injectie
Nobilis Rismavac Suspension et solvant pour suspension injectable
Nobilis Rismavac Suspension und Lösungsmittel zur Herstellung einer Injektionssuspension

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

Target species:
Chicken (one day-old chick)

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
3.00 log10 50% cell culture infectious dose / 0.20 millilitre(s)

Pharmaceutical form:
This information is not available for this product.

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (one day-old chick)**
  - Meat and offal. no withdrawal period 0 days

Subcutaneous use:

- **Chicken (one day-old chick)**
  - Meat and offal. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Belgium

Available in:
Belgium

Package description:

Nobilis Rismavac 1 MLP Bag with 600 ml of solvent for suspension for injection
Nobilis Rismavac 1 MLP Bag with 500 ml of solvent for suspension for injection
Nobilis Rismavac 1 Ampoule with 5000 doses of Suspension and 1 bag (MLP) with 1000 ml of solvent for suspension for injection
Nobilis Rismavac 1 Ampoule with 4000 doses of Suspension and 1 bag (MLP) with 800 ml of solvent for suspension for injection
Nobilis Rismavac 1 Ampoule with 2000 doses of Suspension and 1 bag (MLP) with 400 ml of solvent for suspension for injection
Nobilis Rismavac 1 Ampoule with 1000 doses of Suspension and 1 bag (MLP) with 200 ml of solvent for suspension for injection

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
25/07/2016

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

Responsible authority:
Federal Agency For Medicines And Health Products

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
27/06/2024

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.