NOBILIS RISMAVAC + CA 126 SUSPENSION A DILUER ET SOLVANT POUR SUSPENSION INJECTABLE POUR POULES

Authorised

- Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live
- Turkey herpesvirus, strain FC-126 (cell-associated), Live

Product identification

Medicine name:
NOBILIS RISMAVAC + CA 126 SUSPENSION A DILUER ET SOLVANT POUR SUSPENSION INJECTABLE POUR POULES

Active substance:
Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live
Turkey herpesvirus, strain FC-126 (cell-associated), Live

Target species:
Chicken (one day-old chick)
Chicken (embryonated eggs)

Route of administration:
Intramuscular use
Subcutaneous use
In ovo

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)/dose / 1.00 Dose

Turkey herpesvirus, strain FC-126 (cell-associated), Live
3.00 log10 (50% cell culture infectious dose)/dose / 1.00 Dose

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Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken (one day-old chick)**
  - All relevant tissues. 0 day

Subcutaneous use:

- **Chicken (one day-old chick)**
  - All relevant tissues. 0 day

In ovo:

- **Chicken (embryonated eggs)**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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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
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:
26/02/2003

Manufacturing sites for batch release:
Merck Sharp & Dohme Animal Health S.L.
Intervet International B.V.

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

Authorisation number:
FR/V/9561987 7/2003

Date of authorisation status change:
26/02/2013

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 and Labelling

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