Nobilis Rismavac + CA 126 koncentrat in vehikel za suspenzijo za injiciranje za piščance

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 koncentrat in vehikel za suspenzijo za injiciranje za piščance

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

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 fluorescent assay infectious dose 50% / 0.20 millilitre(s)

Turkey herpesvirus, strain FC-126 (cell-associated), Live
3.00 log10 fluorescent assay infectious dose 50% / 0.20 millilitre(s)

Pharmaceutical form:

Suspension and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken**
  - Meat and offal. 0 day

Subcutaneous use:

- **Chicken**
  - Meat and offal. 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:

Slovenia

Package description:

Available only in Slovenian

Available only in Slovenian
Available only in Slovenian
Available only in Slovenian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
26/04/2002

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
NP/V/0226/001

Date of authorisation status change:
26/04/2002

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.

Labelling

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.