Nobilis Rismavac + CA126 vet. Koncentrat och vätska till injektionsvätska, suspension

Authorised

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

Product identification

Medicine name:
Nobilis Rismavac + CA126 vet. Koncentrat och vätska till injektionsvätska, suspension

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

Target species:
Chicken

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:

Turkey herpesvirus, strain FC-126 (cell-associated), Live
3.00 log10 plaque forming unit(s) / 1.00 Dose

Marek's disease virus, serotype 1, strain CVI-988 (Rispens, cell-associated), Live
3.00 log10 plaque forming unit(s) / 1.00 Dose

Water
1.00 millilitre(s) / 1.00 Dose

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken**
  - Meat and offal. no withdrawal period

Subcutaneous use:

- **Chicken**
  - Meat and offal. no withdrawal period

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:

Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
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Available only in Swedish
Available only in Swedish
Available only in Swedish
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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:
24/03/2000

Manufacturing sites for batch release:
Oriola Sweden AB
Intervet International B.V.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
14302

Date of authorisation status change:

24/03/2000

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

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