

Nobilis Rismavac+CA 126 vakcina A.U.V.

Authorised

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

Product identification

Medicine name:

Nobilis Rismavac+CA 126 vakcina A.U.V.

Active substance:

Turkey herpesvirus, strain FC-126 (cell-associated), Live

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

Target species:

Chicken (hen)

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 unit(s)/dose

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

Pharmaceutical form:

Suspension and solvent for suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Chicken (hen)

- Meat and offal. 0 day

Subcutaneous use:

-

Chicken (hen)

- 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:

Hungary

Available in:

Hungary

Package description:

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

17/06/1999

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

17/06/1999

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