

CRYOMAREX RISPENS

Authorised

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

Product identification

Medicine name:

CRYOMAREX RISPENS

Active substance:

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

Target species:

Chicken (chick, for replacement)

Chicken (broiler)

Chicken (for reproduction)

Chicken (layer hen)

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
10000.00 plaque forming unit / 1.00 Dose

Pharmaceutical form:

Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

-

Chicken (chick, for replacement)

- Meat and offal. 0 day

-

Chicken (broiler)

- Meat and offal. 0 day

-

Chicken (for reproduction)

- Meat and offal. 0 day

-

Chicken (layer hen)

- Meat and offal. 0 day

Subcutaneous use:

-

Chicken (chick, for replacement)

- Meat and offal. 0 day

-

Chicken (broiler)

- Meat and offal. 0 day

-

Chicken (for reproduction)

- Meat and offal. 0 day

-

Chicken (layer 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:

Croatia

Package description:

6 ampoules of 2000 doses of frozen suspension and 1 bag of 2.400 ml of diluent
2 ampoules of 2000 doses of frozen suspension and 1 bag of 800 ml of diluent
1 ampoule of 2000 doses of frozen suspension and 1 bag of 400 ml of de diluent
9 ampoules of 1000 doses of frozen suspension and 1 bag of 1.800 ml of diluent
7 ampoules of 1000 doses of frozen suspension and 1 bag of 1.400 ml of diluent
6 ampoules of 1000 doses of frozen suspension and 1 bag of 1.200 ml of diluent
3 ampoules of 1000 doses of frozen suspension and 1 bag of 600 ml of diluent
2 ampoules of 1000 doses of frozen suspension and 1 bag of 400 ml of diluent
1 ampoule of 1000 doses of frozen suspension + 1 bag of 200 ml of diluent

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

26/09/2014

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/19-01/598

Date of authorisation status change:

10/06/2022

Reference member state:

Spain

Procedure number:

ES/V/0216/001

Concerned member states:

Bulgaria Croatia Cyprus Czechia Greece Latvia Lithuania Romania Slovenia

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

Documents

Summary of Product Characteristics

English (PDF)

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Package Leaflet