CRYOMAREX RISPENS

Not 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:
Surrendered

Authorised in:
Greece

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:
1/07/2020

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France SCS

Responsible authority:
National Organization For Medicines

Authorisation number:
68464/02-07-2020

Date of authorisation status change:
22/05/2024

Reference member state:
Spain

Procedure number:
ES/V/0216/001

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 12/04/2023
Download

Package Leaflet