Poulvac Marek CVI + HVT

Authorised

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

Product identification

Medicine name:
Poulvac Marek CVI + HVT
Poulvac Marek CVI + HVT Συμπυκνωμένο διάλυμα και διαλύτης για ενέσιμο εναιώρημα, για ορνίθια

Active substance:
Marek's disease virus, serotype 1, strain CVI-988 (Rispens), 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), Live
2.90 log10 cell culture infective dose 50 / 0.20 millilitre(s)

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

1000.00 plaque forming unit / 0.20 millilitre(s)

Pharmaceutical form:
Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Chicken**
  - Meat and offal. no withdrawal period 0 days
  - Egg. no withdrawal period 0 days

Subcutaneous use:

- **Chicken**
  - Meat and offal. no withdrawal period 0 days
  - Egg. no withdrawal period 0 days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
Poulvac Marek Diluent: Type II glass vial containing 400 ml
Poulvac Marek Diluent: Type II glass vial containing 200 ml
Poulvac Marek Diluent: PVC plastic bags containing 200 ml
Poulvac Marek Diluent: PVC plastic bags containing 1000 ml

Poulvac Marek CVI+HVT vaccine concentrate: Liquid nitrogen containers of n 1000-dose ampoules. The ampoules are stored in liquid nitrogen containers in a cane (5 ampoules per cane).
Poulvac Marek CVI+HVT vaccine concentrate: Liquid nitrogen containers of n 2000-dose ampoules. The ampoules are stored in liquid nitrogen containers in a cane (5 ampoules per cane).
Poulvac Marek Diluent: PVC plastic bags containing 400 ml

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Hellas S.A.

Marketing authorisation date:
17/12/2013

Manufacturing sites for batch release:
Zoetis Manufacturing & Research Spain S.L.

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00432V

Date of authorisation status change:
22/11/2018

Reference member state:
Netherlands

Procedure number:
NL/V/0102/001

Concerned member states:
Belgium Cyprus Denmark France Germany Greece Hungary Italy Portugal Slovenia Spain

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

Documents

Combined File of all Documents

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