Poulvac Procerta HVT-IBD (--) - Concentrate and solvent for suspension for injection

Authorised

- Turkey herpesvirus, strain HVT-IBD (cell-associated), expressing VP2 protein gene of Infectious bursal disease virus, Live

Product identification

Medicine name:
Poulvac Procerta HVT-IBD (--) - Concentrate and solvent for suspension for injection

Active substance:
Turkey herpesvirus, strain HVT-IBD (cell-associated), expressing VP2 protein gene of Infectious bursal disease virus, Live

Target species:
Chicken
Chicken (embryonated eggs)

Route of administration:
In ovo
Subcutaneous use

Product details

Active substance and strength:

Turkey herpesvirus, strain HVT-IBD (cell-associated), expressing VP2 protein gene of Infectious bursal disease virus, Live
3580.00 plaque forming unit / 1.00 dose

Pharmaceutical form:
Concentrate and solvent for suspension for injection

Withdrawal period by route of administration:

In ovo:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Subcutaneous use:

- **Chicken**
  - Not applicable. 0 day — Zero days
- **Chicken (embryonated eggs)**
  - Not applicable. 0 day — Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD15

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

Authorisation status:
Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:ampoule (glass), Package_size:1 ampoule, Content:4000 doses

Packaging:ampoule (glass), Package_size:1 ampoule, Content:2000 doses

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:

Zoetis Belgium

Marketing authorisation date:

26/10/2023

Manufacturing sites for batch release:

National Laboratory of health, environment and foodstuff
Institute for State Control of Veterinary Biologicals and Medicaments
National Food Chain Safety Office, Directorate of Veterinary Medicinal Products
Bulgarian Food Safety Agency
Health Products Regulatory Authority
NATIONAL VETERINARY RESEARCH INSTITUTE
Institute for Control of Biological Products & Veterinary Drugs
Danish Medicines Agency
Medical Products Agency
Zoetis Manufacturing & Research Spain S.L.
Sciensano
Paul-Ehrlich-Institute
Institute for State Control of Veterinary Biologicals and Medicaments
Federal Office for Safety in Health Care

EVIRA Virology, Animal Diseases and Food Safety

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/10/2023

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

Documents

Combined File of all Documents

English (PDF)
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