Poulvac IB Primer Oral solution or spray solution after reconstitution of the freeze-dried vaccine in water

Authorised

- Infectious bronchitis virus, strain D274, Live
- Infectious bronchitis virus, type Massachusetts, strain H120, Live

Product identification

Medicine name:
Poulvac IB Primer Oral solution or spray solution after reconstitution of the freeze-dried vaccine in water

Active substance:
Infectious bronchitis virus, strain D274, Live
Infectious bronchitis virus, type Massachusetts, strain H120, Live

Target species:
Chicken

Route of administration:
Nebulisation use
In drinking water use
Oculonasal use

Product details

Active substance and strength:
Infectious bronchitis virus, strain D274, Live
1000.00 50% Embryo Infective Dose / 1.00 Dose

Infectious bronchitis virus, type Massachusetts, strain H120, Live
1000.00 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension/use in drinking water

Withdrawal period by route of administration:
Nebulisation use:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

In drinking water use:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day

Oculonasal use:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Portugal

Available in:
Portugal

Package description:
(ID4) 10000 Dose: Box (board) with 10 Bottle each with 1000 Dose, closed with (Aluminium) and (chlorobutylrubber`)
(ID6) 50000 Dose: Box (board) with 10 Bottle each with 5000 Dose, closed with (Aluminium) and (chlorobutylrubber`)
(ID5) 25000 Dose: Box (board) with 10 Bottle each with 2500 Dose, closed with (Aluminium) and (chlorobutylrubber`)

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:
Zoetis Portugal Lda.

Marketing authorisation date:
28/07/1989

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
275/89 DGV

Date of authorisation status change:
6/02/2023

Reference member state:
Germany

Procedure number:
DE/V/0259/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Greece Hungary Latvia Lithuania Netherlands Portugal Romania Slovakia Slovenia Sweden

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

Documents

Combined File of all Documents