Poulvac AE lyophilisate for suspension for use in drinking water

Authorised

- Avian encephalomyelitis virus, strain Calnek 1143, Live

Product identification

Medicine name:
Poulvac AE lyophilisate for suspension for use in drinking water
Poulvac AE, lüofilisaat joogivees manustamiseks kanadele

Active substance:
Avian encephalomyelitis virus, strain Calnek 1143, Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Avian encephalomyelitis virus, strain Calnek 1143, Live
1258.93 50% Embryo Infective Dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for use in drinking water

Withdrawal period by route of administration:
In drinking water use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD02

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

Authorisation status:
Valid

Authorised in:
Estonia

Available in:
Estonia

Package description:
(ID2) 10000 Dose: Box (board) with 10 Bottle (borosilicate glass) each with 1000 Dose, closed with (chlorobutylrubber`) and (Aluminium)
(ID1) 1000 Dose: Box (board) with 1 Bottle (borosilicate glass) with 1000 Dose, closed with (chlorobutylrubber`) and (Aluminium)
(ID4) 20000 Dose: Box (board) with 10 Bottle (borosilicate glass) each with 2000 Dose, closed with (chlorobutylrubber`) and (Aluminium)
(ID3) 2000 Dose: Box (board) with 1 Bottle (borosilicate glass) with 2000 Dose, closed with (chlorobutylrubber`) and (Aluminium)

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 Belgium

Marketing authorisation date:
5/02/2009

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

Responsible authority:
State Agency Of Medicines

Authorisation number:
1540

Date of authorisation status change:
5/02/2009

Reference member state:
Germany

Procedure number:
DE/V/0277/001

Concerned member states:
Austria Estonia Hungary Italy Latvia Lithuania Luxembourg Poland Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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Combined File of all Documents