Cevac Salmovac Lyophilisate for use in drinking water for chickens

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 441/014 (adenine-histidine auxotrophic), Live

Product identification

Medicine name:
Cevac Salmovac Lyophilisate for use in drinking water for chickens

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 441/014 (adenine-histidine auxotrophic), Live

Target species:
Chicken

Route of administration:
In drinking water use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 441/014 (adenine-histidine auxotrophic), Live
100000000.00 Colony forming unit / 1.00 Dose

Pharmaceutical form:
Lyophilisate for use in drinking water

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

- **Chicken**
  - Egg. 3 week — 3 weeks after third vaccination
  - Meat and offal. 6 week — 6 weeks from last vaccination

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AE01

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

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
(ID5) 2500 Dose: Packung with 1 Vial (Glass type I) with 2500 Dose, closed with Kappe (No information) and Stopper (bromobutyl rubber)
(ID1) 10000 Dose: Packung with 10 Vial (Glass type I) each with 1000 Dose, closed with Stopper (bromobutyl rubber) and Kappe (No information)
(ID2) 5000 Dose: Packung with 1 Vial (Glass type II) with 5000 Dose, closed with Stopper (bromobutyl rubber) and Kappe (No information)
(ID3) 60000 Dose: Packung with 12 Vial (Glass type II) each with 5000 Dose, closed with Stopper (bromobutyl rubber) and Kappe (No information)
(ID4) 1000 Dose: Packung with 1 Vial (Glass type I) with 1000 Dose, closed with Stopper (bromobutyl rubber) and Kappe (No information)

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:
Ceva Animal Health Slovakia s.r.o.

Marketing authorisation date:
4/03/2016

Manufacturing sites for batch release:
IDT Biologika GmbH
Ceva-Phylaxia Zrt.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/006/MR/08-S

Date of authorisation status change:
4/03/2016

Reference member state:
Germany

Procedure number:
DE/V/0208/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal Romania Slovakia Slovenia Spain

United Kingdom (Northern Ireland)

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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