Cevac Salmovac Lyophilisate for Use in Drinking Water

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
Cevac Salmovac Lyophilisate for Use in Drinking Water

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
800000000.00 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
  - Egg. 6 week — 6 weeks after second vaccination

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AE01

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

Authorisation status:
Valid

Authorised in:
United Kingdom (Northern Ireland)

Package description:
(ID4): 1 Box with 1 Vial (Glass) with 1000 vaccine doses (1000 vaccine doses)
(ID3): 1 Box with 12 Vials (Glass) with 5000 vaccine doses (60000 vaccine doses)
(ID2): 1 Box with 1 Vial (Glass) with 5000 vaccine doses (5000 vaccine doses)
(ID1): 1 Box with 10 Vials(Glass) with 1000 vaccine doses (10000 vaccine doses)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ceva Sante Animale

Marketing authorisation date:
20/10/2003

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 14966/3003

Date of authorisation status change:
20/07/2024

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