

# Cevac Salmovac Lyophilisate for use in drinking water for chickens

Authorised

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

## Product identification

### Medicine name:

Cevac Salmovac Lyophilisate for use in drinking water for chickens

Cevac Salmovac, Лиофилизат за прилагане във вода за пиене за пилета

### Active substance:

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

800000000.00 Colony forming unit / 1.00 Dose

**Pharmaceutical form:**

Lyophilisate for use in drinking water

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

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**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
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AE01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Bulgaria

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Ceva Animal Health Bulgaria EOOD

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**Marketing authorisation date:**

14/07/2013

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**Manufacturing sites for batch release:**

IDT Biologika GmbH

Ceva-Phylaxia Zrt.

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**Responsible authority:**

Bulgarian Food Safety Authority

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**Authorisation number:**

0022-2066

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**Date of authorisation status change:**

14/07/2013

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0208/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

### Summary of Product Characteristics

English (PDF)

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### Package Leaflet

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### Labelling

This document does not exist in this language (English). You can find it in another language below.