

# 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

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**Active substance:**

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

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**Target species:**

Chicken

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**Route of administration:**

In drinking water use

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## 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

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**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
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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:**

Ireland

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**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)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Ceva Sante Animale

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**Marketing authorisation date:**

5/03/2004

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**Manufacturing sites for batch release:**

IDT Biologika GmbH

Ceva-Phylaxia Zrt.

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA10815/063/001

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**Date of authorisation status change:**

5/03/2004

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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 [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents