

SALMOVAC SE

Not
authorised

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

Product identification

Medicine name:

SALMOVAC SE

Active substance:

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

Target species:

Chicken

Route of administration:

Oral use

In drinking water/milk use

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:**Oral use:**

-

Chicken

- Egg. 3 week nach 3. Vakzination
- Meat and offal. 6 week nach der 2. Vakzination in der Aufzuchtperiode
- Meat and offal. 3 week nach 3. Vakzination

In drinking water/milk use:

-

Chicken

- Egg. 3 week nach 3. Vakzination
- Meat and offal. 6 week nach der 2. Vakzination in der Aufzuchtperiode
- Meat and offal. 3 week nach 3. Vakzination

In drinking water use:

-

Chicken

- Egg. 3 week nach 3. Vakzination
- Meat and offal. 6 week nach der 2. Vakzination in der Aufzuchtperiode
- Meat and offal. 3 week nach 3. Vakzination

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Germany

Package description:

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

Available only in [German](#)

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 Tiergesundheit GmbH

Marketing authorisation date:

11/11/1999

Manufacturing sites for batch release:

IDT Biologika GmbH

Responsible authority:

Paul-Ehrlich-Institut

Authorisation number:

PEI.V.00671.01.1

Date of authorisation status change:

13/06/2023

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

Documents

Summary of Product Characteristics

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