

# Gallimune Se + St, water-in oil emulsion for injection

Authorised

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated
- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated

## Product identification

### Medicine name:

Gallimune Se + St, water-in oil emulsion for injection  
GALLIMUNE SE+ST, injekciné emulsija

### Active substance:

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated  
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated

### Target species:

Chicken (pullet)  
Chicken

### Route of administration:

Intramuscular use

## Product details

### Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated  
149.00 antibody unit(s) / 0.30 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated  
171.00 antibody unit(s) / 0.30 millilitre(s)

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**Chicken (pullet)**

- Meat and offal. 0 day
- Egg. 0 day

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**Chicken**

- Meat and offal. 0 day
  - Egg. 0 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01AB01

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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:**

Lithuania

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**Package description:**

(ID2) 3000 millilitre(s): Box with 10 Bottle (PolyPropylene) each with 300 millilitre(s), closed with Closure and Cap (Acrylnitril-Butadien-Kautschuk, Aluminium)

(ID1) 300 millilitre(s): Box with 1 Bottle (PolyPropylene) with 300 millilitre(s), closed with Closure and Cap (Acrylnitril-Butadien-Kautschuk, Aluminium)

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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:**

Boehringer Ingelheim Animal Health France

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**Marketing authorisation date:**

17/12/2006

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

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**Responsible authority:**

State Food And Veterinary Service

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**Authorisation number:**

LT/2/06/1721/001-002

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**Date of authorisation status change:**

28/05/2025

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0282/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Denmark France Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Poland Portugal 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

Summary of Product Characteristics

Combined File of all Documents

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