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# 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

**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:**

- **Chicken (pullet)**
  - Meat and offal. 0 day
  - Egg. 0 day
- **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:**
Ireland

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**Package description:**
(ID1) 300 millilitre(s): Box with 1 Bottle (PolyPropylene) with 300 millilitre(s), closed with Closure and Cap (Acrylnitril-Butadien-Kautschuk, Aluminium)
(ID2) 3000 millilitre(s): Box with 10 Bottle (PolyPropylene) each with 300 millilitre(s), closed with Closure and Cap (Acrylnitril-Butadien-Kautschuk, Aluminium)

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## 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:**
Boehringer Ingelheim Vetmedica GmbH

**Marketing authorisation date:**
22/06/2007

**Manufacturing sites for batch release:**
Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

**Responsible authority:**
Health Products Regulatory Authority

**Authorisation number:**
VPA10454/054/001

**Date of authorisation status change:**
22/06/2007

**Reference member state:**
Germany

**Procedure number:**
DE/V/0282/001

**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)

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

## Documents

Combined File of all Documents