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

Not 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)

Pharmaceutical form:
Emulsion for injection

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AB01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Revoked

Authorised in:
Portugal

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)

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 Animal Health Portugal Unipessoal Lda.

Marketing authorisation date:
1/01/2000

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
775/07DIVPT

Date of authorisation status change:
17/12/2019

Reference member state:
Germany

Procedure number:
DE/V/0282/001

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