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

Authorised

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

Product identification

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

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

Target species:
Chicken (pullet)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated
171.00 antibody unit(s) / 0.30 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated 149.00 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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AB01

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
(ID2) 3000 millilitre(s): Box (Cardboard) with 10 Bottle (PolyPropylene) each with 300 millilitre(s)
(ID1) 300 millilitre(s): Box (Cardboard) with 1 Bottle (PolyPropylene) with 300 millilitre(s)

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:
15/06/2007

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health Italia S.p.A.

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.03348.01.1

Date of authorisation status change:
5/01/2021

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

Summary of Product Characteristics

English (PDF)

Published on: 7/02/2024

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