Nobilis Salenvac T suspension for injection for chickens

Authorised

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

Product identification

Medicine name:
Nobilis Salenvac T suspension for injection for chickens
Nobilis Salenvac T

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

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated

1.00 relative potency / 0.50 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated
1.00 relative potency / 0.50 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **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:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
Cardboard box containing a 500 ml (1000 doses) low density polyethylene multi-dose bottle closed with an aluminium ring on top of a rubber stopper
Cardboard box containing a 250 ml (500 doses) low density polyethylene multi-dose bottle closed with an aluminium ring on top of a rubber stopper

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
7/01/2004

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.02970.01.1

Date of authorisation status change:
29/05/2009

Reference member state:
Italy

Procedure number:
IT/V/0137/001

Concerned member states:
Austria Czechia Denmark France Germany Greece Hungary Lithuania Luxembourg Netherlands Portugal Romania Slovakia 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

English (PDF)
Published on: 6/10/2025
Download

Summary of Product Characteristics

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

Package Leaflet

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