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

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:
Austria

Available in:
Austria

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 Ges.m.b.H.

Marketing authorisation date:
1/12/2003

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

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
8-20259

Date of authorisation status change:
1/12/2003

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
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Package Leaflet

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Summary of Product Characteristics

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

Labelling

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