

# 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

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**Active substance:**

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

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**Target species:**

Chicken

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**Route of administration:**

Intramuscular use

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## 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)

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**Pharmaceutical form:**

Suspension for injection

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**Withdrawal period by route of administration:**

**Intramuscular use:**

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**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:**

Germany

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**Available in:**

Germany

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**Package description:**

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

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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Intervet Deutschland GmbH

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**Marketing authorisation date:**

7/01/2004

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Paul-Ehrlich-Institut

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**Authorisation number:**

PEI.V.02970.01.1

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**Date of authorisation status change:**

29/05/2009

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**Reference member state:**

Italy

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**Procedure number:**

IT/V/0137/001

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**Concerned member states:**

Austria Czechia Denmark France Germany Greece Hungary Lithuania  
Luxembourg Netherlands Portugal Romania Slovakia Spain  
United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
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## Documents

Combined File of all Documents

English (PDF)

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