

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:

France

Available in:

France

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

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

Marketing authorisation date:

14/01/2004

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/8276024 1/2004

Date of authorisation status change:

14/01/2004

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)

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Summary of Product Characteristics

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Package Leaflet and Labelling

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