

Nobilis Salenvac T

Not
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

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:

Medicinal product subject to medical prescription

Authorisation status:

Surrendered

Authorised in:

Denmark

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 International B.V.

Marketing authorisation date:

15/06/2005

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Danish Medicines Agency

Authorisation number:

35307

Date of authorisation status change:

15/06/2005

Reference member state:

Italy

Procedure number:

IT/V/0137/001

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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