Nobilis Salenvac T

Not authorised

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

Product identification

Medicine name:
Nobilis Salenvac T

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

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

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated
1.00 relative potency / 0.50 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, 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:
Surrendered

Authorised in:
Luxembourg

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:
25/02/2009

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

Responsible authority:
Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:
V 817/04/02/0784

Date of authorisation status change:
25/02/2009

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

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