Nobilis Salenvac ETC suspension for injection for chickens

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated
- Salmonella enterica, subsp. enterica, serovar Infantis, strain A S03499-06, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain DT104, Inactivated

Product identification

Medicine name:
Nobilis Salenvac ETC suspension for injection for chickens

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain PT4, Inactivated
Salmonella enterica, subsp. enterica, serovar Infantis, strain A S03499-06, 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 Infantis, strain A S03499-06, 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**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Cyprus

Available in:

Cyprus

Package description:

Low density polyethylene bottle containing 1000 doses of vaccine (each dose = 0.5 ml)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
6/04/2020

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

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00788V

Date of authorisation status change:
6/04/2020

Reference member state:
Netherlands

Procedure number:
NL/V/0305/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany Greece Hungary Italy Latvia Lithuania Luxembourg Poland Portugal Romania Slovakia Slovenia 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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