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Cevac Salmune ETI K (--) - Suspension for injection

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 038-90, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 076-94, Inactivated
- Salmonella enterica, subsp. enterica, serovar Infantis, strain SM-595, Inactivated

Product identification

Medicine name:
Cevac Salmune ETI K (--) - Suspension for injection

Active substance:
Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 038-90, Inactivated
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 076-94, Inactivated
Salmonella enterica, subsp. enterica, serovar Infantis, strain SM-595, Inactivated

Target species:
Chicken

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Enteritidis, strain 038-90, Inactivated

Presentation_strength:≥ 122 ELISA units Reference:In house Index:0

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 076-94, Inactivated

Presentation_strength:≥ 212 ELISA units Reference:In house Index:1

Salmonella enterica, subsp. enterica, serovar Infantis, strain SM-595, Inactivated

Presentation_strength:≥ 157 ELISA units Reference:In house Index:2

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Chicken**
  - Not applicable. 0 day Zero days

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:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Belgium , Cyprus , Czechia , Germany , Greece , Hungary , Italy , Luxembourg , Netherlands , Poland , Portugal , Romania , Slovakia , Spain ,

United Kingdom (Northern Ireland)

Package description:
Packaging:Bottle (LDP), Package_size:5 bottles, Content:1000 doses
Packaging:Bottle (LDP), Package_size:1 bottle, Content:1000 doses

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Ceva-Phylaxia Zrt.

Marketing authorisation date:
30/08/2024

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
30/08/2024

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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