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Cevac Meta L

Authorised

- Turkey rhinotracheitis virus, strain CRR126, Live

Product identification

Medicine name:
Cevac Meta L

Active substance:
Turkey rhinotracheitis virus, strain CRR126, Live

Target species:
Chicken

Route of administration:
Ocular use
Nebulisation use

Product details

Active substance and strength:
Turkey rhinotracheitis virus, strain CRR126, Live
3.80 log10 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate for oculonasal suspension

Withdrawal period by route of administration:

Ocular use:

- **Chicken**
  - All relevant tissues. 0 day

Nebulisation use:

- **Chicken**
  - All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AD01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Package description:
Cardboard box with 20 vials of 2500 doses of vaccine.
Cardboard box with 1 vial of 3000 doses of vaccine.
Cardboard box with 10 vials of 3000 doses of vaccine.
Cardboard box with 20 vials of 3000 doses of vaccine.
Cardboard box with 1 vial of 4000 doses of vaccine.
Cardboard box with 10 vials of 4000 doses of vaccine.
Cardboard box with 20 vials of 4000 doses of vaccine.
Cardboard box with 1 vial of 5000 doses of vaccine.
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Cardboard box with 20 vials of 1000 doses of vaccine.
Cardboard box with 10 vials of 1000 doses of vaccine.
Cardboard box with 1 vial of 1000 doses of vaccine.
Cardboard box with 20 vials of 500 doses of vaccine.
Cardboard box with 10 vials of 500 doses of vaccine.
Cardboard box with 1 vial of 500 doses of vaccine.

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:
Ceva Sante Animale

Marketing authorisation date:
22/07/2016

Manufacturing sites for batch release:
Ceva-Phylaxia Zrt.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6805958 3/2016

Date of authorisation status change:
21/09/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0200/001

Concerned member states:

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Poland Portugal Romania Slovakia Slovenia Spain Sweden

United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

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