Livacox Q, Perorální suspenze

Authorised

- Eimeria acervulina, strain CH-P-72/89, Live
- Eimeria maxima, strain J-MN 82/88, Live
- Eimeria necatrix, strain NH-UK 94/95, Live
- Eimeria tenella, strain CH-E-A, Live

Product identification

Medicine name:
Livacox Q, Perorální suspenze

Active substance:
Eimeria acervulina, strain CH-P-72/89, Live
Eimeria maxima, strain J-MN 82/88, Live
Eimeria necatrix, strain NH-UK 94/95, Live
Eimeria tenella, strain CH-E-A, Live

Target species:
Chicken

Route of administration:
In drinking water use
Nebulisation use

Product details

Active substance and strength:

Eimeria acervulina, strain CH-P-72/89, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria maxima, strain J-MN 82/88, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria necatrix, strain NH-UK 94/95, Live
120.00 oocyst(s) / 1.00 Dose

Eimeria tenella, strain CH-E-A, Live
500.00 oocyst(s) / 1.00 Dose

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
In drinking water use:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day

Nebulisation use:

- **Chicken**
  - Egg. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AN01

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

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Authorisation status:
Valid

Authorised in:
Czechia

Package description:
Available only in Czech
Available only in Czech
Available only in Czech
Available only in Czech

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:
BIOPHARM Vyzkumny ustav biofarmacie a veterinarnich leciv a.s.

Marketing authorisation date:
10/09/1999

Manufacturing sites for batch release:
BIOPHARM Vyzkumny ustav biofarmacie a veterinarnich leciv a.s.

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/046/99-C

Date of authorisation status change:
20/10/2016

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.