PARACOX-8 BG

Authorised

- Eimeria tenella, strain HP, Live
- Eimeria praecox, strain HP, Live
- Eimeria necatrix, strain HP, Live
- Eimeria mitis, strain HP, Live
- Eimeria maxima, strain MFP, Live
- Eimeria maxima, strain CP, Live
- Eimeria brunetti, strain HP, Live
- Eimeria acervulina, strain HP, Live

Product identification

Medicine name:
PARACOX-8 BG

Active substance:
Eimeria tenella, strain HP, Live
Eimeria praecox, strain HP, Live
Eimeria necatrix, strain HP, Live
Eimeria mitis, strain HP, Live
Eimeria maxima, strain MFP, Live
Eimeria maxima, strain CP, Live
Eimeria brunetti, strain HP, Live
Eimeria acervulina, strain HP, Live

Target species:
Chicken (chick)

Route of administration:
In drinking water use
Nebulisation use

Product details

Active substance and strength:
Eimeria tenella, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria praecox, strain HP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria necatrix, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria mitis, strain HP, Live
1000.00 oocyst(s) / 1.00 Dose

Eimeria maxima, strain MFP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria maxima, strain CP, Live
200.00 oocyst(s) / 1.00 Dose

Eimeria brunetti, strain HP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria acervulina, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Pharmaceutical form:
Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AN01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

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:
12/08/2009

Manufacturing sites for batch release:
Merck Sharp & Dohme Animal Health S.L.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2300

Date of authorisation status change:
15/01/2019

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.