

Paracox-8 vet., suspensija geriamajai suspensijai ruošti vištoms

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 vet., suspensija geriamajai suspensijai ruošti vištoms

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

Route of administration:

In drinking water use

In-feed use

Coarse spray

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:

Suspension for oral suspension

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

-

Chicken

In-feed use:

-

Chicken

Coarse spray:

-

Chicken

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AN01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:

Valid

Authorised in:

Lithuania

Package description:

Available only in [Lithuanian](#)

Available only in [Lithuanian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

3/09/2001

Manufacturing sites for batch release:

MSD Animal Health UK Limited

Merck Sharp & Dohme Animal Health S.L.

Responsible authority:

State Food And Veterinary Service

Authorisation number:

LT/2/01/1331/001-002

Date of authorisation status change:

2/09/2006

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

Documents

Combined File of all Documents

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