

Paracox 8 suspenzija za peroralno suspenzijo za piščance

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 suspenzija za peroralno suspenzijo za piščance

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-feed use

In drinking water use

Product details

Active substance and strength:

Eimeria tenella, strain HP, Live

500.00 Organisms / 0.00 millilitre(s)

Eimeria praecox, strain HP, Live

100.00 Organisms / 0.00 millilitre(s)

Eimeria necatrix, strain HP, Live

500.00 Organisms / 0.00 millilitre(s)

Eimeria mitis, strain HP, Live

1000.00 Organisms / 0.00 millilitre(s)

Eimeria maxima, strain MFP, Live

100.00 Organisms / 0.00 millilitre(s)

Eimeria maxima, strain CP, Live

200.00 Organisms / 0.00 millilitre(s)

Eimeria brunetti, strain HP, Live

100.00 Organisms / 0.00 millilitre(s)

Eimeria acervulina, strain HP, Live

500.00 Organisms / 0.00 millilitre(s)

Pharmaceutical form:

Suspension for oral suspension

Withdrawal period by route of administration:**In-feed use:**

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Chicken

- Meat and offal. 0 day

In drinking water use:

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Chicken

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AN01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Slovenia

Package description:

Available only in Slovenian

Available only in Slovenian

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:

31/12/2002

Manufacturing sites for batch release:

MSD Animal Health UK Limited
Merck Sharp & Dohme Animal Health S.L.

Responsible authority:

Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:

NP/V/0252/001

Date of authorisation status change:

31/12/2002

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.