File downloaded on 2026-11-24
Source URL: https://medicines.health.europa.eu/veterinary/en/600000027720

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:

- **Chicken**

- Meat and offal. 0 day

In drinking water use:

- **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.