

Paracox-8, suspenzija za oralnu suspenziju za kokoši (piliće)

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 oralnu suspenziju za kokoši (piliće)

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:

Oral use

Product details

Active substance and strength:

Eimeria tenella, strain HP, Live
500.00 oocyst(s) / 0.10 millilitre(s)

Eimeria praecox, strain HP, Live
100.00 oocyst(s) / 0.10 millilitre(s)

Eimeria necatrix, strain HP, Live
500.00 oocyst(s) / 0.10 millilitre(s)

Eimeria mitis, strain HP, Live
1000.00 oocyst(s) / 0.10 millilitre(s)

Eimeria maxima, strain MFP, Live
100.00 oocyst(s) / 0.10 millilitre(s)

Eimeria maxima, strain CP, Live
200.00 oocyst(s) / 0.10 millilitre(s)

Eimeria brunetti, strain HP, Live
100.00 oocyst(s) / 0.10 millilitre(s)

Eimeria acervulina, strain HP, Live
500.00 oocyst(s) / 0.10 millilitre(s)

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:**Oral use:**

-

Chicken

- Meat and offal. 0 day Cjepivo se ne primjenjuje nesilicama.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AN01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Croatia

Package description:

Available only in [Croatian](#)

Available only in [Croatian](#)

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:

Intervet International B.V. Subsidiary In The Republic Of Croatia

Marketing authorisation date:

10/01/2013

Manufacturing sites for batch release:

Schering-Plough Limited

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/17-01/309

Date of authorisation status change:

13/05/2026

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.