

# Paracox-5, suspenzija za oralnu suspenziju, za kokoši (piliće)

Authorised

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

## Product identification

**Medicine name:**

Paracox-5, suspenzija za oralnu suspenziju, za kokoši (piliće)

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**Active substance:**

Eimeria tenella, strain HP, Live

Eimeria mitis, strain HP, Live

Eimeria maxima, strain MFP, Live

Eimeria maxima, strain CP, Live

Eimeria acervulina, strain HP, Live

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**Target species:**

Chicken

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**Route of administration:**

Oral use

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## Product details

### Active substance and strength:

Eimeria tenella, strain HP, Live  
500.00 oocyst(s) / 0.00 millilitre(s)

Eimeria mitis, strain HP, Live  
1000.00 oocyst(s) / 0.00 millilitre(s)

Eimeria maxima, strain MFP, Live  
100.00 oocyst(s) / 0.00 millilitre(s)

Eimeria maxima, strain CP, Live  
200.00 oocyst(s) / 0.00 millilitre(s)

Eimeria acervulina, strain HP, Live  
500.00 oocyst(s) / 0.00 millilitre(s)

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### Pharmaceutical form:

Oral suspension

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### Withdrawal period by route of administration:

#### Oral use:

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#### Chicken

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

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### Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AN01

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### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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### Authorisation status:

Valid

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### Authorised in:

Croatia

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**Package description:**

Available only in [Croatian](#)

Available only in [Croatian](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

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

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**Marketing authorisation date:**

9/01/2013

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Ministry Of Agriculture Veterinary And Food Safety Directorate

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**Authorisation number:**

UP/I-322-05/17-01/313

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**Date of authorisation status change:**

27/12/2024

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

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