

# Paracox-8 suspensie voor suspensie voor oraal gebruik voor kippen

Authorised

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

## Product identification

### **Medicine name:**

Paracox-8 suspensie voor suspensie voor oraal gebruik voor kippen

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### **Active substance:**

Eimeria maxima, strain MFP, Live

Eimeria maxima, strain CP, Live

Eimeria brunetti, strain HP, Live

Eimeria praecox, strain HP, Live

Eimeria mitis, strain HP, Live

Eimeria tenella, strain HP, Live

Eimeria acervulina, strain HP, Live

Eimeria necatrix, 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 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 brunetti, strain HP, Live  
100.00 oocyst(s) / 0.00 millilitre(s)

Eimeria praecox, strain HP, Live  
100.00 oocyst(s) / 0.00 millilitre(s)

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

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

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

Eimeria necatrix, 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. no withdrawal period  
Withdrawal period = zero days

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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:**

Netherlands

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**Available in:**

Netherlands

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**Package description:**

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

Available only in [Dutch](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Legal basis not covered by Directive 2001/82/EC

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**Marketing authorisation holder:**

Intervet Nederland B.V.

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**Marketing authorisation date:**

17/06/1999

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**Manufacturing sites for batch release:**

MERCK SHARP & DOHME ANIMAL HEALTH, S.L.

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**Responsible authority:**

Medicines Evaluation Board

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**Authorisation number:**

REG NL 3070

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**Date of authorisation status change:**

17/10/2024

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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

Combined File of all Documents

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