

# PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS

Authorised

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

## Product identification

**Medicine name:**

PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS

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**Active substance:**

Eimeria maxima, strain CP, Live

Eimeria tenella, strain HP, Live

Eimeria mitis, strain HP, Live

Eimeria maxima, strain MFP, Live

Eimeria acervulina, strain HP, Live

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**Target species:**

Chicken (broiler)

Chicken (one day-old chick)

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**Route of administration:**

Oral use

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## Product details

### **Active substance and strength:**

Eimeria maxima, strain CP, Live  
500.00 Organisms / 0.00 millilitre(s)

Eimeria tenella, strain HP, Live  
1000.00 Organisms / 0.00 millilitre(s)

Eimeria mitis, strain HP, Live  
100.00 Organisms / 0.00 millilitre(s)

Eimeria maxima, strain MFP, Live  
200.00 Organisms / 0.00 millilitre(s)

Eimeria acervulina, strain HP, Live  
500.00 Organisms / 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 (broiler)**

- All relevant tissues. 0 day

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#### **Chicken (one day-old chick)**

- All relevant tissues. 0 day

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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:**

Luxembourg

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**Package description:**

box of 5 vials of 4 mL (1000 doses)

1 vial of 500 mL (solvent)

1 vial of 100 mL (solvent)

box of 5 vials of 20 mL (5000 doses)

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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 International B.V.

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**Marketing authorisation date:**

3/04/2000

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**Manufacturing sites for batch release:**

Merck Sharp & Dohme Animal Health S.L.

MSD Animal Health UK Limited

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**Responsible authority:**

Ministry Of Health And Social Security

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**Authorisation number:**

V 817/00/06/0658

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**Date of authorisation status change:**

3/04/2000

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**Reference member state:**

France

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**Procedure number:**

FR/V/0351/001

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**Concerned member states:**

Austria Belgium Cyprus Czechia Denmark Estonia Finland Germany Greece  
Hungary Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway  
Poland Portugal Slovakia Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

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