PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS

Authorised

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

Product identification

Medicine name:
PARACOX-5, SUSPENSION FOR ORAL SUSPENSION FOR CHICKENS
Paracox-5 suspensija iekšķīgi lietojamas suspensijas pagatavošanai vistām

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Active substance:
Eimeria acervulina, strain HP, Live

Carminic acid

Eimeria maxima, strain CP, Live

Eimeria tenella, strain HP, Live

Eimeria mitis, strain HP, Live

Eimeria maxima, strain MFP, Live

Target species:
Chicken (broiler)
Chicken (one day-old chick)

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria acervulina, strain HP, Live
500.00 Organisms / 1.00 Dose

Carminic acid
0.10 millilitre(s) / 1.00 Dose

Eimeria maxima, strain CP, Live
500.00 Organisms / 1.00 Dose

Eimeria tenella, strain HP, Live
1000.00 Organisms / 1.00 Dose

Eimeria mitis, strain HP, Live
100.00 Organisms / 1.00 Dose

Eimeria maxima, strain MFP, Live
200.00 Organisms / 1.00 Dose

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:

Oral use:

- **Chicken (broiler)**
  - All relevant tissues. 0 day
- **Chicken (one day-old chick)**
  - All relevant tissues. 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:
Latvia

Package description:
Container of 5 plastic vials of 20 mL (5000 doses)
Container of 1 plastic vial of 100 mL (1000 doses)
Container of 1 plastic vial of 500 mL (5000 doses)
Container of 5 plastic vials of 4 mL (1000 doses)

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:
31/05/2006

Manufacturing sites for batch release:
Merck Sharp & Dohme Animal Health S.L.
MSD Animal Health UK Limited

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/MRP/06/1674

Date of authorisation status change:
31/05/2006

Reference member state:

France

Procedure number:
FR/V/0351/001

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)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

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