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
Paracox 5 suspenzija za peroralno suspenzijo za piščance

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

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

Route of administration:
Oral use

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)

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:
Slovenia

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)

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:
16/04/2003

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

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0251/001

Date of authorisation status change:
16/04/2003

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

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.