Paracox 8 oral suspension

Authorised

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

Product identification

Medicine name:
Paracox 8 oral suspension

Active substance:
Eimeria tenella, strain HP, Live

Eimeria praecox, strain HP, Live

Eimeria necatrix, strain HP, Live

Eimeria mitis, strain HP, Live

Eimeria maxima, strain MFP, Live

Eimeria maxima, Live

Eimeria brunetti, strain HP, Live

Eimeria acervulina, Live

Target species:
Chicken (chick)

Route of administration:
In drinking water use
In drinking water/milk use
Topical

Product details

Active substance and strength:
Eimeria tenella, strain HP, Live
500.00 cells / 0.00 millilitre(s)

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

Eimeria necatrix, strain HP, Live
500.00 cells / 0.00 millilitre(s)

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

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

Eimeria maxima, Live
200.00 cells / 0.00 millilitre(s)

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

Eimeria acervulina, Live
500.00 cells / 0.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
In drinking water use:

- **Chicken (chick)**
  - Meat and offal. 0 day
  - Egg. 0 day

In drinking water/milk use:

- **Chicken (chick)**
  - Meat and offal. 0 day
  - Egg. 0 day

Topical:

- **Chicken (chick)**
  - Meat and offal. 0 day
  - Egg. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AN

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

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Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID1): 1 Box with 1 Bottle (Polyethylenterephthalat Glycol-modifiziert) with 4 millilitre(s) (4 millilitre(s))
(ID3): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 100 millilitre(s) (100 millilitre(s))
(ID4): 1 Box with 1 Bottle (PolyEthylene TerePhthalate) with 500 millilitre(s) (500 millilitre(s))
(ID2): 1 Box with 1 Bottle (Polyethylenterephthalat Glycol-modifiziert) with 20 millilitre(s) (20 millilitre(s))

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Intervet Deutschland GmbH

Marketing authorisation date:
10/08/1998

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

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
483a/91

Date of authorisation status change:
24/08/2008

Reference member state:
Germany

Procedure number:
DE/V/0210/001

Concerned member states:
Austria

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 28/01/2022
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