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Paracox-8 vet. Suspension till oral suspension

Authorised

- Water
- 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, strain CP, Live
- Eimeria brunetti, strain HP, Live
- Eimeria acervulina, strain HP, Live

Product identification

Medicine name:
Paracox-8 vet. Suspension till oral suspension

Active substance:
Water
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, strain CP, Live
Eimeria brunetti, strain HP, Live

Eimeria acervulina, strain HP, Live

Target species:
Chicken

Route of administration:
In drinking water use
Nebulisation use

Product details

Active substance and strength:
Water
1.00 millilitre(s) / 1.00 millilitre(s)

Eimeria tenella, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria praecox, strain HP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria necatrix, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Eimeria mitis, strain HP, Live
1000.00 oocyst(s) / 1.00 Dose

Eimeria maxima, strain MFP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria maxima, strain CP, Live
200.00 oocyst(s) / 1.00 Dose

Eimeria brunetti, strain HP, Live
100.00 oocyst(s) / 1.00 Dose

Eimeria acervulina, strain HP, Live
500.00 oocyst(s) / 1.00 Dose

Pharmaceutical form:
Suspension for oral suspension

Withdrawal period by route of administration:

In drinking water use:

- **Chicken**
  - Meat and offal. no withdrawal period

Ej till fjäderfä som producerar ägg för human konsumtion

Nebulisation use:

- **Chicken**
  - Meat and offal. no withdrawal period

Ej till fjäderfä som producerar ägg för human konsumtion

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI01AN01

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

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Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish
Available only in Swedish

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:
25/02/1994

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

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
12039

Date of authorisation status change:
25/02/1994

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

Documents

Package Leaflet

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Summary of Product Characteristics

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.