Paracox-8 vet. suspensjon til mikstur, suspensjon til kylling

Authorised

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

Product identification

Medicine name:
Paracox-8 vet. suspensjon til mikstur, suspensjon til kylling

Active substance:
Eimeria praecox, strain HP, Live

Eimeria necatrix, strain HP, Live

Eimeria acervulina, Live

Eimeria mitis, strain HP, Live

Eimeria maxima, strain CP, Live

Eimeria maxima, strain MFP, Live

Eimeria brunetti, strain HP, Live

Eimeria tenella, strain HP, Live

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria praecox, strain HP, Live
100.00 oocyst(s) / 0.00 millilitre(s)

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

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

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

Eimeria maxima, strain CP, Live
200.00 oocyst(s) / 0.00 millilitre(s)

Eimeria maxima, strain MFP, Live
100.00 oocyst(s) / 0.00 millilitre(s)

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

Eimeria tenella, strain HP, Live
500.00 oocyst(s) / 0.00 millilitre(s)

Pharmaceutical form:
Suspension for oral suspension

Withdrawal period by route of administration:
Oral use:

- **Chicken**
  - 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:
Norway

Available in:
Norway

Package description:
Available only in Norwegian
Available only in Norwegian
Available only in Norwegian
Available only in Norwegian

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:
2/09/1996

Manufacturing sites for batch release:
MSD Animal Health UK Limited

Responsible authority:

Norwegian Medical Products Agency

Authorisation number:
0000-08234

Date of authorisation status change:
1/10/2006

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.