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HuveGuard MMAT suspension for oral suspension for chickens

Authorised

- Eimeria tenella, strain Rt 3+15, Live
- Eimeria acervulina, strain RA 3+20, Live
- Eimeria mitis, strain Jormit 3+9, Live
- Eimeria maxima, strain MCK +10, Live

Product identification

Medicine name:
HuveGuard MMAT suspension for oral suspension for chickens

Active substance:
Eimeria tenella, strain Rt 3+15, Live

Eimeria acervulina, strain RA 3+20, Live

Eimeria mitis, strain Jormit 3+9, Live

Eimeria maxima, strain MCK +10, Live

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria tenella, strain Rt 3+15, Live
417.00 oocyst(s) / 0.03 millilitre(s)

Eimeria acervulina, strain RA 3+20, Live
50.00 unit(s) / 0.03 millilitre(s)

Eimeria mitis, strain Jormit 3+9, Live
278.00 unit(s) / 0.03 millilitre(s)

Eimeria maxima, strain MCK +10, Live
278.00 unit(s) / 0.03 millilitre(s)

Pharmaceutical form:
Suspension for oral suspension

Withdrawal period by route of administration:
Oral use:

- **Chicken**
  - Meat and offal. 0 day
  - Egg. 0 day

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:
Malta

Package description:

Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 10 vials of 5000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 10 vials of 1000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 5 vials of 5000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 5 vials of 1000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 1 vial of 5000 doses.
Low density polyethylene (LDPE) vial of 30 ml with a grey butyl rubber stopper and aluminum cap. Carboard box with 1 vial of 1000 doses.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
2/04/2019

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Veterinary And Phytosanitary Regulation Department

Authorisation number:
VMA75

Date of authorisation status change:
2/04/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0206/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Malta Norway Poland Portugal Romania 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