HuveGuard MMAT

Authorised

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

Product identification

Medicine name:
HuveGuard MMAT
HuveGuard MMAT suspensão para suspensão oral em galinhas

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

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:
Eimeria maxima, strain MCK +10, Live
100.00 unit(s) / 1.00 Dose

Eimeria mitis, strain Jormit 3+9, Live
100.00 unit(s) / 1.00 Dose

Eimeria acervulina, strain RA 3+20, Live
50.00 unit(s) / 1.00 Dose

Eimeria tenella, strain Rt 3+15, Live
150.00 unit(s) / 1.00 Dose

Pharmaceutical form:
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:
Portugal

Available in:
Portugal

Package description:
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.
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 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 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 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 10 vials of 5000 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:
3/06/2016

Manufacturing sites for batch release:
Biovet J.S.C.

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
928/01/16RIVPT

Date of authorisation status change:
13/04/2022

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

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