

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 vakcina A.U.V.

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

Authorisation status:

Valid

Authorised in:

Hungary

Available in:

Hungary

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:

13/07/2016

Manufacturing sites for batch release:

Biovet J.S.C.

Responsible authority:

National Food Chain Safety Office

Authorisation number:

3791/X/16 NÉBIH ÁTI

Date of authorisation status change:

13/07/2016

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

Source URL: <https://medicines.health.europa.eu/veterinary/600000036046>