Evant (--) - Suspension and solution for oral spray

Authorised

- Eimeria acervulina, strain 003, Live
- Eimeria maxima, strain 013, Live
- Eimeria mitis, strain 006, Live
- Eimeria praecox, strain 007, Live
- Eimeria tenella, strain 004, Live

Product identification

Medicine name:
Evant (--) - Suspension and solution for oral spray

Active substance:
Eimeria acervulina, strain 003, Live

Eimeria maxima, strain 013, Live

Eimeria mitis, strain 006, Live

Eimeria praecox, strain 007, Live

Eimeria tenella, strain 004, Live

Target species:
Chicken

Route of administration:
Oral use

Product details

Active substance and strength:

Eimeria acervulina, strain 003, Live

Presentation_strength:332-450 sporulated oocysts Reference:Ph. Eur. Index:0

Eimeria maxima, strain 013, Live

Presentation_strength:196-265 sporulated oocysts Reference:Ph. Eur. Index:1

Eimeria mitis, strain 006, Live

Presentation_strength:293-397 sporulated oocysts Reference:Ph. Eur. Index:2

Eimeria praecox, strain 007, Live

Presentation_strength:293-397 sporulated oocysts Reference:Ph. Eur. Index:3

Eimeria tenella, strain 004, Live

Presentation_strength:276-374 sporulated oocysts Reference:Ph. Eur. Index:4

Pharmaceutical form:

This information is not available for this product.

Withdrawal period by route of administration:

Oral use:

- **Chicken**
  - Not applicable. 0 day Zero days

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:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein ,

Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial solvent, Content:Suspension: 7 ml (1000 doses); Solvent: 50 ml
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial solvent, Content:Suspension: 70 ml (10000 doses); Solvent: 500 ml
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial solvent, Content:Suspension: 35 ml (5000 doses); Solvent: 250 ml
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial Solvent (Hipracell), Content:Suspension: 70 ml (10000 doses); Solvent: 500 ml
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial Solvent (Hipracell), Content:Suspension: 35 ml (5000 doses); Solvent: 250 ml
Packaging:Vial (glass); Solvent: vial (PP), Package_size:1 vial Suspension; 1 vial Solvent (Hipracell), Content:Suspension: 7 ml (1000 doses); Solvent: 50 ml

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:
Laboratorios Hipra, S.A.

Marketing authorisation date:
5/02/2019

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
5/02/2019

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

Documents

Combined File of all Documents

English (PDF)
Published on: 4/12/2025
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