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Enterisol Ileitis lyophilisate and diluent for oral suspension for pigs

Authorised

- Lawsonia intracellularis, strain MS B3903, Live

Product identification

Medicine name:
Enterisol Ileitis lyophilisate and diluent for oral suspension for pigs

Active substance:
Lawsonia intracellularis, strain MS B3903, Live

Target species:
Pig

Route of administration:
Oral use
In-feed use
In drinking water use

Product details

Active substance and strength:
Lawsonia intracellularis, strain MS B3903, Live
79432.00 50% tissue culture infectious dose / 1.00 Dose

Pharmaceutical form:
Lyophilisate and solvent for oral suspension

Withdrawal period by route of administration:

Oral use:

- **Pig**
  - Meat and offal. 0 day

In-feed use:

- **Pig**
  - Meat and offal. 0 day

In drinking water use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI09AE04

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
(ID2) 10 Dose; 20 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 10 Dose, closed with Stopper (Bromobutyl Rubber) and Cap (Aluminium) and 1 Vial (High Density PolyEthylene) with 20 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)
(ID4) 50 Dose; 100 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 50 Dose, closed with Stopper (Bromobutyl Rubber) and Cap (Aluminium) and 1 Vial (High

Density PolyEthylene) with 100 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)
(ID6) 100 Dose; 200 millilitre(s): Box (Cardboard) with 1 Vial (Glass type I) with 100 Dose, closed with Cap (Aluminium) and Stopper (Bromobutyl Rubber) and 1 Vial (High Density PolyEthylene) with 200 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)
(ID8) 1200 Dose; 2400 millilitre(s): Box (Cardboard) with 12 Vial (Glass type I) each with 100 Dose, closed with Stopper (Bromobutyl Rubber) and Cap (Aluminium) and 12 Vial (High Density PolyEthylene) each with 200 millilitre(s), closed with Stopper (Chlorobutyl Rubber) and Cap (Aluminium)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:
28/09/2008

Manufacturing sites for batch release:
Boehringer Ingelheim Vetmedica GmbH

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1254

Date of authorisation status change:
28/09/2008

Reference member state:

Germany

Procedure number:
DE/V/0236/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands 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

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.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents