

# 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

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**Active substance:**

Lawsonia intracellularis, strain MS B3903, Live

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**Target species:**

Pig

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Lawsonia intracellularis, strain MS B3903, Live

79433.00 50% tissue culture infectious dose / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate and solvent for oral suspension

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**Withdrawal period by route of administration:**

**Oral use:**

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**Pig**

- Meat and offal. no withdrawal period  
Withdrawal period is 0 days

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI09AE04

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**Package description:**

(ID2): 1 Box with (1 Bottle (Glass) with 10 Dose and 1 Bottle (High Density PolyEthylene) with 20 millilitre(s)) (10.0 Dose, 20.0 millilitre(s))

(ID4): 1 Box with (1 Bottle (Glass) with 50 Dose and 1 Bottle (High Density PolyEthylene) with 100 millilitre(s)) (50.0 Dose, 100.0 millilitre(s))

(ID6): 1 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (High Density PolyEthylene) with 200 millilitre(s)) (100.0 Dose, 200.0 millilitre(s))

(ID8): 1 Box with 12 Box with (1 Bottle (Glass) with 100 Dose and 1 Bottle (High Density PolyEthylene) with 200 millilitre(s)) (1200.0 Dose, 2400.0 millilitre(s))

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

19/08/2005

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Vetmedica GmbH

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1624

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**Date of authorisation status change:**

19/08/2005

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0236/001

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**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)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

English (PDF)

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### Labelling

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.