

# Lincocin Soluble Powder, 400 mg/g powder for use in drinking water

Authorised

- Lincomycin hydrochloride

## Product identification

### **Medicine name:**

Lincocin Soluble Powder, 400 mg/g powder for use in drinking water  
Lincocin Soluble Powder, 400 mg/g powder for use in drinking water

### **Active substance:**

Lincomycin hydrochloride

### **Target species:**

Chicken

Pig

### **Route of administration:**

Oral use

## Product details

### **Active substance and strength:**

Lincomycin hydrochloride  
453.63 milligram(s) / 1.00 gram(s)

**Pharmaceutical form:**

Powder for use in drinking water

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**Withdrawal period by route of administration:****Oral use:**

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**Chicken**

- Meat and offal. 5 day

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**Pig**

- Meat and offal. 24 hour

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

QJ01FF02

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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:**

Ireland

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**Available in:**

Ireland

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**Package description:**

White high density polyethylene (HDPE) bottle containing 150 g powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid with an aluminium cap. Pack sizes: Bottle of 150 g

White high density polyethylene (HDPE) bottle containing 1.5 kg powder for use in drinking water with a white tamper evident low density polyethylene (LDPE) lid. Pack sizes: Bottle of 1.5 kg

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Phibro Animal Health (Poland) Sp. z o.o.

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**Marketing authorisation date:**

26/09/2014

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**Manufacturing sites for batch release:**

Zoetis Belgium

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**Responsible authority:**

Health Products Regulatory Authority

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**Authorisation number:**

VPA25497/003/001

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**Date of authorisation status change:**

26/09/2014

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0410/001

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**Concerned member states:**

Belgium France Germany Luxembourg Poland

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