

# GIROLAN/APRALAN

Not authorised

This information is not available for this product.

## Product identification

**Medicine name:**

GIROLAN/APRALAN  
APRALAN SOLUBILE

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**Active substance:**

This information is not available for this product.

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**Target species:**

Pig (weaned piglet)  
Rabbit  
Cattle (pre-ruminant)  
Chicken (broiler)

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**Route of administration:**

In drinking water/milk use

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## Product details

**Active substance and strength:**

This information is not available for this product.

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**Pharmaceutical form:**

Powder for use in drinking water/milk

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**Withdrawal period by route of administration:**

**In drinking water/milk use:**

- **Pig (weaned piglet)**

- Meat and offal. 0 day

- **Rabbit**

- Meat and offal. 0 day

- **Cattle (pre-ruminant)**

- Meat and offal. 28 day

- **Chicken (broiler)**

- Meat and offal. 0 day

- Eggs. no withdrawal period

Eggs: Not authorised for use in laying birds producing eggs for human consumption.  
Do not use within 4 weeks of onset of the laying period.

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

QJ01GB90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Italy

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**Package description:**

box containing 50 sachets containing 2 x 10e6 UI of apramycin sulfate and 3,6 g of product

box containing 50 sachets containing 1 x 10e6 UI of apramycin sulfate and 1,8 g of product

bag containing 1.000 x 10e6 UI of apramycin sulfate and 1.812 g of product

bottle containing 50 x 10e6 UI of apramycin sulfate and 91 g of product

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## Additional information

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**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Elanco GmbH

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**Marketing authorisation date:**

10/11/1997

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**Manufacturing sites for batch release:**

Elanco France S.A.S

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**Responsible authority:**

Ministry Of Health

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**Authorisation number:**

102442

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**Date of authorisation status change:**

2/02/2022

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**Reference member state:**

Spain

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**Procedure number:**

ES/V/0321/001

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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

Combined File of all Documents

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