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# Spizobactin 750,000 IU / 125 mg chewable tablets for dogs

Authorised

- Metronidazole
- Spiramycin

## Product identification

**Medicine name:**

Spizobactin 750,000 IU / 125 mg chewable tablets for dogs

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**Active substance:**

Metronidazole

Spiramycin

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**Target species:**

Dog

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**Route of administration:**

Oral use

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## Product details

**Active substance and strength:**

Metronidazole

125.00 milligram(s) / 1.00 Tablet

Spiramycin

178.57 milligram(s) / 1.00 Tablet

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**Pharmaceutical form:**

Chewable tablet

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

QJ01RA04

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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:**

Germany

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**Available in:**

Germany

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**Package description:**

(ID11) 100 Chewable tablet: Box (cardboard) with 10 Blister (aluminium) each with 10 Chewable tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

(ID1) 10 Chewable tablet: Box (cardboard) with 1 Blister (aluminium) with 10 Chewable tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

(ID10) 100 Chewable tablet: Box (cardboard) with 10 Box (cardboard) each with 1 Blister (aluminium) with 10 Chewable tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

(ID3) 30 Chewable tablet: Box (cardboard) with 3 Blister (aluminium) each with 10 Chewable tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

(ID2) 20 Chewable tablet: Box (cardboard) with 2 Blister (aluminium) each with 10 Chewable tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Le Vet. Beheer B.V.

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**Marketing authorisation date:**

16/08/2017

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**Manufacturing sites for batch release:**

Lelypharma B.V.

Genera d.d.

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**Responsible authority:**

Federal Office Of Consumer Protection And Food Safety

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**Authorisation number:**

402378.00.00

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**Date of authorisation status change:**

18/09/2022

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**Reference member state:**

Germany

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**Procedure number:**

DE/V/0171/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland

France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg

Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain

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

Combined File of all Documents

English (PDF)

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