

# NexGard Spectra 37.5 mg + 7.5 mg - Chewable tablet

Authorised

- Afoxolaner
- Milbemyacin oxime

## Product identification

**Medicine name:**

NexGard Spectra 37.5 mg + 7.5 mg - Chewable tablet

---

**Active substance:**

Afoxolaner

Milbemyacin oxime

---

**Target species:**

Dog

---

**Route of administration:**

Oral use

---

## Product details

**Active substance and strength:**

Afoxolaner

Presentation\_strength:37.5 mg Reference:In house Index:0

Milbemyacin oxime

Presentation\_strength:7.5 mg Reference:In house Index:1

---

**Pharmaceutical form:**

Chewable tablet

---

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

QP54AB51

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

---

**Available in:**

Belgium , Czechia , Estonia , France , Latvia , Lithuania , Luxembourg , Poland , Slovakia , Spain

---

**Package description:**

Packaging:Blister (PVC/Alu), Package\_size:15 tablets

Packaging:Blister (PVC/Alu), Package\_size:3 tablets

Packaging:Blister (PVC/Alu), Package\_size:6 tablets (1 blister of 6 tablets)

Packaging:Blister (PVC/Alu), Package\_size:1 tablet

Packaging:Blister (PVC/Alu), Package\_size:6 tablets (2 blisters of 3 tablets)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

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

---

**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

---

**Marketing authorisation date:**

15/01/2015

---

**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

---

**Responsible authority:**

European Commission

---

**Authorisation number:**

This information is not available for this product.

---

**Date of authorisation status change:**

15/12/2022

---

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)

Published on: 6/02/2026

[Download](#)

ema-puar-v3842-nexgardspectra-ws-vra0000245082-en.pdf

ema-puar-v3842-nexgard-spectra-ws2280-g-vra-0034-g-en.pdf

ema-puar-nexgard-spectra-nex-spe-var-ws1338-en.pdf

ema-puar-nexgard-spectra-v-3842-par-en.pdf

ema-puar-nexgard-spectra-v-3842-var-ii-0008-en.pdf

ema-puar-nexgard-spectra-v-3842-var-ii-0019-en.pdf

ema-puar-nexgard-spectra-nex-spe-var-ws1559-en.pdf