# Exzolt 10 mg/ml - Solution for use in drinking water

Authorised

- Fluralaner

## Product identification

**Medicine name:**
Exzolt 10 mg/ml - Solution for use in drinking water

**Active substance:**
Fluralaner

**Target species:**
Chicken

**Route of administration:**
In drinking water use

## Product details

**Active substance and strength:**
Fluralaner
10.00 milligram(s) / 1.00 Bottle

**Pharmaceutical form:**
Solution for use in drinking water

**Withdrawal period by route of administration:**

**In drinking water use:**

- **Chicken**
  - Meat and offal. 14 day 14 days
  - Egg. 0 day Zero days

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

QP53BE02

**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:**

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

**Package description:**

Packaging:Bottle (glass type III), Package_size:1 bottle, Content:50 ml

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:4 litres

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:1 litre

Packaging:Bottle (glass), Package_size:1 bottle, Content:4 ml

## Additional information

**Entitlement type:**

Marketing Authorisation

**Legal basis of product authorisation:**
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Intervet International B.V.

**Marketing authorisation date:**
18/08/2017

**Manufacturing sites for batch release:**
Intervet Productions S.A.

**Responsible authority:**
European Commission

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
25/10/2022

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents

English (PDF)
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