# Spotinor 10 mg/ml Spot-on Solution for cattle and sheep

Authorised

- Deltamethrin

## Product identification

**Medicine name:**
Spotinor 10 mg/ml Spot-on Solution for cattle and sheep

**Active substance:**
Deltamethrin

**Target species:**
Cattle
Sheep

**Route of administration:**
Spot-on use

## Product details

**Active substance and strength:**
Deltamethrin
10.00 milligram(s) / 1.00 millilitre(s)

**Pharmaceutical form:**
Spot-on solution

**Withdrawal period by route of administration:**

**Spot-on use:**

- **Cattle**
  - Meat and offal. 17 day
  - Milk. 0 hour
- **Sheep**
  - Meat and offal. 35 day

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

QP53AC11

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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:**

Italy

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**Package description:**

250 ml clear high-density polyethylene bottle with internal graduated calibration chamber and awhite screw polypropylene cap in a cardboard box.
1 x 500 ml clear high-density polyethylene bottle with internal graduated calibration chamber and awhite screw polypropylene cap in a cardboard box.
2 x 500 ml clear high-density polyethylene bottle with internal graduated calibration chamber and awhite screw polypropylene cap in a cardboard box.
1 litre white high density polyethylene back pack for use with a suitable dosing device and a white screwpolypropylene cap in a cardboard box.
2.5 litre white high density polyethylene back pack for use with a suitable dosing device and a white screwpolypropylene cap in a cardboard box.

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## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Generic application (Article 13(1) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Norbrook Laboratories (Ireland) Limited

**Marketing authorisation date:**
12/12/2014

**Manufacturing sites for batch release:**
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

**Responsible authority:**
Ministry Of Health

**Authorisation number:**
104632

**Date of authorisation status change:**
18/06/2019

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0544/001

**Concerned member states:**
Austria Belgium Bulgaria Cyprus Denmark Estonia Finland France Greece Hungary Italy Latvia Lithuania Luxembourg Malta Poland Portugal Romania Spain Sweden United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

Combined File of all Documents

English (PDF)
Published on: 2/03/2025
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