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Proactive 1.52 mg/mL Teat Dip/Spray Solution

Authorised

- Iodine (125I)

Product identification

Medicine name:
Proactive 1.52 mg/mL Teat Dip/Spray Solution

Active substance:
Iodine (125I)

Target species:
Cattle

Route of administration:
Teat use

Product details

Active substance and strength:
Iodine (125I)
1.52 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Teat dip/spray solution

Withdrawal period by route of administration:

Teat use:

- **Cattle**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD08AG03

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Greece

Package description:
A dark liquid contained in 200 litre, opaque high-density polyethylene drums with screw closures and seals.
A dark liquid contained in 60 litre, opaque high-density polyethylene drums with screw closures and seals.
A dark liquid contained in 20 litre, opaque high-density polyethylene drums with screw closures and seals.
A dark liquid contained in 10 litre, opaque high-density polyethylene drums with screw closures and seals.
A dark liquid contained in 5 litre, opaque high-density polyethylene drums with screw closures and seals.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Delaval

Marketing authorisation date:
6/07/2009

Manufacturing sites for batch release:
Delaval

Responsible authority:
National Organization For Medicines

Authorisation number:
37383/24-03-2026/K-0271301

Date of authorisation status change:
23/03/2026

Reference member state:
Ireland

Procedure number:
IE/V/0483/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Germany Greece Hungary Iceland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia 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

English (PDF)
Published on: 16/01/2026
Download

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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