

Trodax 340 mg/ml Solution for Injection

Not
authorised

- Nitroxinil

Product identification

Medicine name:

Trodax 340 mg/ml Solution for Injection

Active substance:

Nitroxinil

Target species:

Cattle

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Nitroxinil

340.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:**Subcutaneous use:**

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Cattle

- Meat and offal. 60 day

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Sheep

- Meat and offal. 60 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG08

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Ireland

Package description:

1 litre container of natural polypropylene with chlorobutyl rubber stopper.

500ml container of natural polypropylene with chlorobutyl rubber stopper.

250 ml container of natural polypropylene with chlorobutyl rubber stopper.

100 ml DIN bottle with chlorobutyl rubber stopper.

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:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

1/10/1999

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10454/077/001

Date of authorisation status change:

20/02/2025

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

Documents

Summary of Product Characteristics