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Nanotrim 464.2 mg/g + 100 mg/g powder for use in drinking water/milk for chickens, turkeys, pigs and cattle

Authorised

- Sulfachlorpyridazine sodium
- Trimethoprim

Product identification

Medicine name:
Nanotrim 464.2 mg/g + 100 mg/g powder for use in drinking water/milk for chickens, turkeys, pigs and cattle

Active substance:
Sulfachlorpyridazine sodium
Trimethoprim

Target species:
Pig
Chicken
Turkey
Cattle (pre-ruminant)

Route of administration:
In drinking water/milk use

Product details

Active substance and strength:

Sulfachlorpyridazine sodium

500.00 milligram(s)/gram / 1.00 gram(s)

Trimethoprim

100.00 milligram(s)/gram / 1.00 gram(s)

Pharmaceutical form:

Powder for use in drinking water/milk

Withdrawal period by route of administration:

In drinking water/milk use:

- **Pig**
  - Meat and offal. 7 day
- **Chicken**
  - Meat and offal. 3 day
- **Turkey**
  - Meat and offal. 9 day
- **Cattle (pre-ruminant)**
  - Meat and offal. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW12

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:
Poland

Package description:
Pillow sachet with 100 g made of polyethylene/aluminium/polyethylene terephthalate laminate
1k g resealable block-bottom zipped sachet made of polyethylene/aluminium/polyethylene terephthalate laminate

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 19(1) of Regulation (EU) 2019/6)

Marketing authorisation holder:
HuVepharma

Marketing authorisation date:
21/03/2025

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
3398

Date of authorisation status change:
21/03/2025

Reference member state:
Netherlands

Procedure number:

NL/V/0420/001

Concerned member states:
Austria Bulgaria Croatia Cyprus Czechia Denmark Estonia Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Malta Poland Portugal Romania Slovakia Slovenia Spain

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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