

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

NANOTRIM 464,2 mg/g + 100 mg/g POLVO PARA ADMINISTRACION EN AGUA DE BEBIDA O EN LECHE PARA POLLOS, PAVOS, PORCINO Y BOVINO

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

Authorisation status:

Valid

Authorised in:

Spain

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:

27/02/2025

Manufacturing sites for batch release:

Biovet AD

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

4379 ESP

Date of authorisation status change:

28/02/2025

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

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.