# AVEMIX N° 150

Authorised

- Sulfamethoxypyridazine
- Trimethoprim

## Product identification

**Medicine name:**
AVEMIX N° 150

**Active substance:**
Sulfamethoxypyridazine
Trimethoprim

**Target species:**
Cattle
Horse (foal)
Pig (piglet)
Rabbit
Horse (mare)
Sheep (lamb)
Sheep
Cattle (calf)
Poultry

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Sulfamethoxypyridazine
116.20 milligram(s) / 1.00 gram(s)

Trimethoprim
25.00 milligram(s) / 1.00 gram(s)

---

**Pharmaceutical form:**
Oral powder

---

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

- **Cattle**
  - Milk. no withdrawal period

La spécialité n'est pas destinée aux femelles laitières productrices de lait de consommation.

- **Horse (foal)**
  - Meat and offal. 12 day

- **Pig (piglet)**
  - Meat and offal. 12 day

- **Rabbit**
  - Meat and offal. 12 day

- **Horse (mare)**

- Milk. no withdrawal period

En l'absence de LMR pour le lait, ne pas utiliser chez les femelles productrices de lait de consommation, en lactation ou en période de tarissement ni chez les futures productrices de lait de consommation.

- **Sheep (lamb)**
  - Meat and offal. 12 day

- **Sheep**
  - Milk. no withdrawal period

La spécialité n'est pas destinée aux femelles laitières productrices de lait de consommation.

- **Cattle (calf)**
  - Meat and offal. 12 day

- **Poultry**
  - Meat and offal. 12 day
  - Eggs. no withdrawal period

En l'absence de LMR pour les oeufs, ne pas utiliser chez les espèces pondeuses productrices d'oeuf de consommation.

---

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QJ01EW15

---

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

---

**Authorisation status:**
Valid

---

**Authorised in:**

France

**Package description:**
Available only in French
Available only in French
Available only in French
Available only in French

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Legal basis not covered by Directive 2001/82/EC

**Marketing authorisation holder:**
Vetoquinol S.A.

**Marketing authorisation date:**
10/07/1992

**Manufacturing sites for batch release:**
Vetoquinol Biowet Sp. z o.o.
Vetoquinol S.A.

**Responsible authority:**
French Agency For Food, Environmental And Occupational Health & Safety

**Authorisation number:**
FR/V/1003139 1/1992

**Date of authorisation status change:**
10/07/2012

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 and Labelling

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

---

**Source URL:** *https://medicines.health.europa.eu/veterinary/600000035501*