

# Danilon equidos 1.5 g Granules for horses and ponies

Authorised

- Suxibuzone

## Product identification

**Medicine name:**

Danilon equidos 1.5 g Granules for horses and ponies

---

**Active substance:**

Suxibuzone

---

**Target species:**

Horse

Horse (pony)

---

**Route of administration:**

In-feed use

---

## Product details

**Active substance and strength:**

Suxibuzone

1.50 gram(s) / 10.00 gram(s)

---

**Pharmaceutical form:**

Granules

---

**Withdrawal period by route of administration:****In-feed use:**

- 

**Horse**

- Meat and offal. no withdrawal period

Not to be used in animals intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

- Milk. no withdrawal period

Not to be used in animals intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

---

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

QM01AA90

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Norway

---

**Package description:**

(ID2) 600 gram(s): Box with 60 Bag (paper; aluminium; polyethylene) each with 10 gram(s)

(ID1) 180 gram(s): Box with 18 Bag (paper; aluminium; polyethylene) each with 10 gram(s)

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Generic application (Article 13(1) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Ecuphar Veterinaria S.L.U.

---

**Marketing authorisation date:**

7/10/2011

---

**Manufacturing sites for batch release:**

Esteve Pharmaceuticals S.A.

---

**Responsible authority:**

Norwegian Medical Products Agency

---

**Authorisation number:**

10-7651

---

**Date of authorisation status change:**

18/05/2016

---

**Reference member state:**

Germany

---

**Procedure number:**

DE/V/0192/001

---

**Concerned member states:**

Austria Belgium Czechia Denmark Estonia Hungary Iceland Latvia Lithuania  
Norway Poland Romania Slovakia Slovenia United Kingdom (Northern Ireland)

---

To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

English (PDF)

Published on: 17/02/2026

[Download](#)

### Package Leaflet

English (PDF)

Published on: 17/02/2026

[Download](#)

2401451-paren-20180315.rtf