

# Equipalazone 1 g Oral powder

Authorised

- Phenylbutazone

## Product identification

**Medicine name:**

Fenylbutazon Dechra Vet 1 g pulver til hest og ponni

Equipalazone 1 g Oral powder

**Active substance:**

Phenylbutazone

**Target species:**

Horse (pony)

Horse

**Route of administration:**

Oral use

## Product details

**Active substance and strength:**

Phenylbutazone

1.00 gram(s) / 1.00 Sachet

**Pharmaceutical form:**

Oral powder

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

- 

**Horse (pony)**

- Meat and offal. no withdrawal period

Not for use in horses 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.

- 

**Horse**

- Meat and offal. no withdrawal period

Not for use in horses 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:**

QM01AA01

---

**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

---

**Authorisation status:**

Valid

---

**Authorised in:**

Norway

---

**Package description:**

Equipalazone 1 g or. pwdr. sachet 100

Equipalazone 1 g or. pwdr. sachet 32

---

## Additional information

**Entitlement type:**

Marketing Authorisation

---

**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

---

**Marketing authorisation holder:**

Dechra Regulatory B.V.

---

**Marketing authorisation date:**

1/09/2017

---

**Manufacturing sites for batch release:**

Genera d.d.

---

**Responsible authority:**

Norwegian Medical Products Agency

---

**Authorisation number:**

16-11233

---

**Date of authorisation status change:**

1/09/2017

---

**Reference member state:**

Belgium

---

**Procedure number:**

BE/V/0037/001

---

**Concerned member states:**

Austria Norway Poland Portugal Slovakia 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

### Package Leaflet

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

### Summary of Product Characteristics

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