**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000086080

# Vulketan 2.5 mg/g Gel

Authorised

Ketanserin tartrate

# **Product identification**

#### **Medicine name:**

Vulketan 2.5 mg/g Gel

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#### **Active substance:**

Ketanserin tartrate

#### **Target species:**

Horse

#### **Route of administration:**

Cutaneous use

# **Product details**

# **Active substance and strength:**

Ketanserin tartrate
3.45 milligram(s) / 1.00 gram(s)

#### **Pharmaceutical form:**

Gel

# Withdrawal period by route of administration: Cutaneous use:

Horse

- Meat and offal. no withdrawal period 0 days
- Milk. no withdrawal period 0 days

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

OD03AX90

#### Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

#### **Authorisation status:**

Valid

#### **Authorised in:**

Belgium

# Package description:

Vulketan Tube with 75 g Gel

# Additional information

## **Entitlement type:**

Marketing Authorisation

# Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

# Marketing authorisation holder:

Audevard

## Marketing authorisation date:

9/09/1991

Manufacturing sites for batch release: Sanochemia Pharmazeutika AG
Responsible authority: Federal Agency For Medicines And Health Products
<b>Authorisation number:</b> BE-V155242
Date of authorisation status change: 18/12/2019
To consult adverse reactions on veterinary medicinal products please go to <a href="https://www.adrreports.eu/vet">www.adrreports.eu/vet</a>
Documents
Package Leaflet
This document does not exist in this language (English). You can find it in another language below.
Summary of Product Characteristics
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Labelling

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