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# Vulketan 2.5 mg/g gel for horses

Authorised

- Ketanserin tartrate

## Product identification

**Medicine name:**

Vulketan 2.5 mg/g gel for horses

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**Active substance:**

Ketanserin tartrate

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**Target species:**

Horse

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Ketanserin tartrate

3.45 milligram(s) / 1.00 gram(s)

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**Pharmaceutical form:**

Gel

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**Withdrawal period by route of administration:**

**Cutaneous use:**

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**Horse**

- Meat and offal. 0 day

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

QD03AX90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria

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**Package description:**

The product is packed in 75 g aluminum tubes with HDPE screw cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Audevard

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**Marketing authorisation date:**

31/08/2011

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**Manufacturing sites for batch release:**

Sanochemia Pharmazeutika AG

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**Responsible authority:**

Austrian Agency For Health And Food Safety

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**Authorisation number:**

8-00990

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**Date of authorisation status change:**

31/08/2011

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0265/001

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**Concerned member states:**

Austria Denmark Finland France Germany Iceland Italy Luxembourg  
Netherlands Portugal Sweden United Kingdom (Northern Ireland)

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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: 3/05/2024

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Package Leaflet

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Labelling

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