

Vulketan 2.5 mg/g gel for horses

Authorised

- Ketanserin tartrate

Product identification

Medicine name:

Vulketan 2.5 mg/g gel for horses

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. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD03AX90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

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

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:

Audevard

Marketing authorisation date:

2/08/2011

Manufacturing sites for batch release:

Sanochemia Pharmazeutika AG

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 108178

Date of authorisation status change:

26/01/2022

Reference member state:

Ireland

Procedure number:

IE/V/0265/001

Concerned member states:

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

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Combined File of all Documents

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

Summary of Product Characteristics