Vulketan 2.5 mg/g gel for horses

Authorised

- Ketanserin tartrate

Product identification

Medicine name:
Vulketan 2.5 mg/g gel for horses
Vulketan 2,5 mg/g gel voor paarden

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