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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

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:

Italy

Package description:

Available only in Italian

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:

13/07/2012

Manufacturing sites for batch release:

Sanochemia Pharmazeutika AG

Responsible authority:

Ministry Of Health

Authorisation number:

104327

Date of authorisation status change:

13/07/2012

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

Summary of Product Characteristics