

Zoletil 50, (25 mg/ml+25 mg/ml) lyophilisate and solvent for solution for injection for dogs and cats

Authorised

- Zolazepam hydrochloride
- Tiletamine hydrochloride

Product identification

Medicine name:

Zoletil 50, (25 mg/ml+25 mg/ml) lyophilisate and solvent for solution for injection for dogs and cats

Active substance:

Zolazepam hydrochloride

Tiletamine hydrochloride

Target species:

Dog

Cat

Route of administration:

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Zolazepam hydrochloride
125.00 milligram(s) / 1.00 Vial
Tiletamine hydrochloride
125.00 milligram(s) / 1.00 Vial

Pharmaceutical form:

Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN01AX99

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Available in:

Bulgaria

Package description:

Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Virbac

Marketing authorisation date:

17/06/2007

Manufacturing sites for batch release:

Virbac

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-2046

Date of authorisation status change:

17/06/2007

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

Documents

Summary of Product Characteristics

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Package Leaflet

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

Labelling

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