ZOLETIL 50

Authorised

- Tiletamine hydrochloride
- Zolazepam hydrochloride

Product identification

Medicine name:
ZOLETIL 50

Active substance:
Tiletamine hydrochloride

Zolazepam hydrochloride

Target species:
Dog

Route of administration:
Intravenous use
Intramuscular use

Product details

Active substance and strength:
Tiletamine hydrochloride
125.00 milligram(s) / 1.00 Vial

Zolazepam hydrochloride
125.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Powder and solution for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN01AX99

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Package description:
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:
20/06/2002

Manufacturing sites for batch release:
Virbac

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

110236

Date of authorisation status change:
27/03/2023

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

Documents

Summary of Product Characteristics

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