

ZOLETIL 100 (50 mg/mL + 50 mg/mL)

Authorised

- Tiletamine hydrochloride
- Zolazepam hydrochloride

Product identification

Medicine name:

ZOLETIL 100 (50 mg/mL + 50 mg/mL)

Active substance:

Tiletamine hydrochloride

Zolazepam hydrochloride

Target species:

Cat

Dog

Route of administration:

Intravenous use

Intramuscular use

Product details

Active substance and strength:

Tiletamine hydrochloride

250.00 milligram(s) / 1.00 Vial

Zolazepam hydrochloride
250.00 milligram(s) / 1.00 Vial

Pharmaceutical form:

Lyophilisate 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:

Croatia

Available in:

Croatia

Package description:

Available only in [Croatian](#)

Available only in [Croatian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:

Virbac

Marketing authorisation date:

27/01/2021

Manufacturing sites for batch release:

Valdepharm

Responsible authority:

Ministry Of Agriculture Veterinary And Food Safety Directorate

Authorisation number:

UP/I-322-05/21-01/76

Date of authorisation status change:

22/12/2025

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

Documents

Summary of Product Characteristics

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