Zoletil 100, süstelahuse pulber ja lahusti, koertele ja kassidele

Authorised

- Tiletamine
- Zolazepam

Product identification

Medicine name:
Zoletil 100, süstelahuse pulber ja lahusti, koertele ja kassidele

Active substance:
Tiletamine
Zolazepam

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use

Product details

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

Zolazepam
250.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Powder 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:
Estonia

Available in:
Estonia

Package description:
Available only in Estonian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
7/06/2006

Manufacturing sites for batch release:
Virbac

Responsible authority:
State Agency Of Medicines

Authorisation number:
1408

Date of authorisation status change:
7/06/2006

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

Documents

Summary of Product Characteristics

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