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Zoletil 50 Vet. pulver og solvens til injektionsvæske, opløsning 25 + 25 mg/ml

Authorised

- Tiletamine hydrochloride
- Zolazepam hydrochloride

Product identification

Medicine name:
Zoletil 50 Vet. pulver og solvens til injektionsvæske, opløsning 25 + 25 mg/ml

Active substance:
Tiletamine hydrochloride
Zolazepam hydrochloride

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:

Tiletamine hydrochloride
145.50 milligram(s) / 1.00 Vial

Zolazepam hydrochloride
141.00 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:
Denmark

Available in:
Denmark

Package description:
Available only in Danish
Available only in Danish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Virbac

Marketing authorisation date:

30/10/1991

Manufacturing sites for batch release:
Virbac

Responsible authority:
Danish Medicines Agency

Authorisation number:
14169

Date of authorisation status change:
30/10/1991

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

Documents

Summary of Product Characteristics

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