SEDEDORM 1 MG/ML SOLUTION FOR INJECTION FOR DOGS AND CATS

Authorised

- Medetomidine hydrochloride

Product identification

Medicine name:
SEDEDORM 1 MG/ML SOLUTION FOR INJECTION FOR DOGS AND CATS

Active substance:
Medetomidine hydrochloride

Target species:
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use
Intravenous use

Product details

Active substance and strength:
Medetomidine hydrochloride

1.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN05CM91

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Hungary

Available in:
Hungary

Package description:
Type I clear glass vials of 10 ml capacity. Vials are fitted with a bromobutyl stopper and sealed with an aluminium cap. Vials are packed in a cardboard box. Pack sizes: Box with 1 vial

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Vetpharma Animal Health S.L.

Marketing authorisation date:
17/04/2009

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Responsible authority:
Directorate Of Veterinary Medicinal Products

Authorisation number:
2519/X/09 MgSzH ÁTI

Date of authorisation status change:
17/04/2009

Reference member state:
France

Procedure number:
FR/V/0192/001

Concerned member states:
Belgium Germany Hungary Netherlands Poland Portugal Spain United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents