

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:

United Kingdom (Northern Ireland)

Package description:

Available only in French

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:

31/03/2009

Manufacturing sites for batch release:

Industrial Veterinaria S.A.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 32509/4000

Date of authorisation status change:

27/10/2024

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