

SEDATOR, 1.0 mg/ml, solution for injection for cats and dogs

Authorised

- Medetomidine hydrochloride

Product identification

Medicine name:

SEDATOR, 1.0 mg/ml, solution for injection for cats and dogs

Active substance:

Medetomidine hydrochloride

Target species:

Dog

Cat

Route of administration:

Intramuscular use

Intravenous use

Subcutaneous 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:

Ireland

Available in:

Ireland

Package description:

Clear colourless, sterile aqueous solution and presented in a Type I clear glass vial of 5 ml capacity, each packed in a cardboard box. Vials are fitted with a teflon coated halogenated rubber stopper and sealed with an aluminium cap.

Clear colourless, sterile aqueous solution and presented in a Type I clear glass vial of 10 ml capacity, each packed in a cardboard box. Vials are fitted with a teflon coated halogenated rubber stopper and sealed with an aluminium cap.

Clear colourless, sterile aqueous solution and presented in a Type I clear glass vial of 20 ml capacity, each packed in a cardboard box. Vials are fitted with a teflon coated halogenated rubber stopper and sealed with an aluminium cap.

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:

Eurovet Animal Health B.V.

Marketing authorisation date:

26/09/2008

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10989/057/001

Date of authorisation status change:

26/09/2008

Reference member state:

Ireland

Procedure number:

IE/V/0476/001

Concerned member states:

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics