Atipam 5.0 mg/ml solution for injection for Cats and Dogs - atipamezole hydrochloride

Authorised

Atipamezole hydrochloride

Product identification

Medicine name:

Atipam 5.0 mg/ml solution for injection for Cats and Dogs - atipamezole hydrochloride Atipam 5 mg/ml Oplossing voor injectie

Atipam 5 mg/ml Solution injectable

Atipam 5 mg/ml Injektionslösung

Active substance:

Atipamezole hydrochloride

Target species:

Dog

Cat

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Atipamezole hydrochloride 5.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Dog

•

Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QV03AB90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Package description:

Cardboard box with 1 clear glass type I vial of 20 ml, with a teflon coated halogenated rubber stopper and aluminium cap

Cardboard box with 1 clear glass type I vial of 10 ml, with a teflon coated halogenated rubber stopper and aluminium cap

Cardboard box with 1 clear glass type I vial of 5 ml, with a teflon coated halogenated rubber stopper and 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:

Dechra Regulatory B.V.

Marketing authorisation date:

24/03/2008

Manufacturing sites for batch release:

Eurovet Animal Health B.V.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V315542

Date of authorisation status change:

24/03/2008

Reference member state:

Netherlands

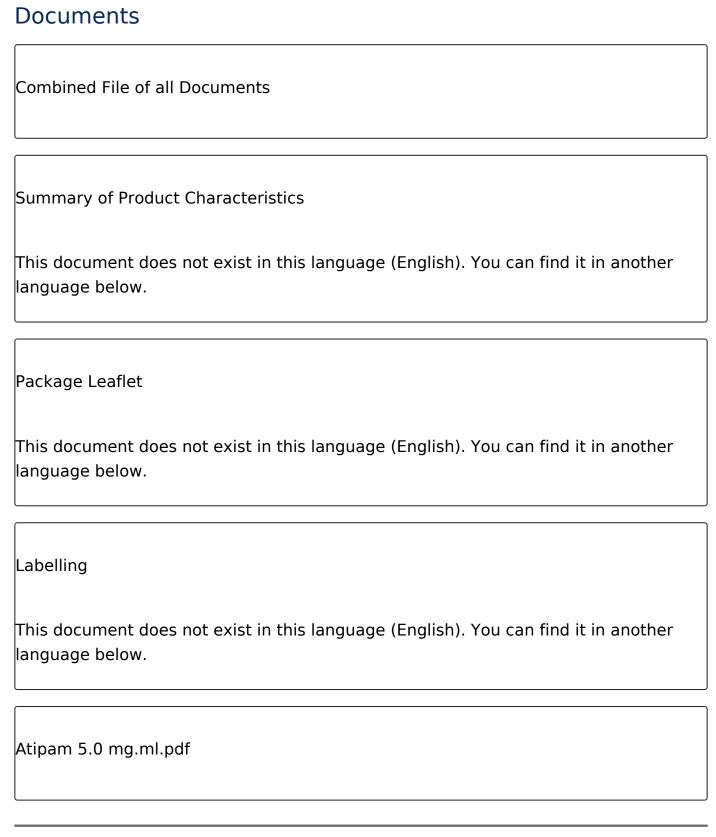
Procedure number:

NL/V/0279/001

Concerned member states:

Austria Belgium Czechia Denmark Finland France Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Norway Poland Portugal Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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



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