CALMIVET SOLUTION INJECTABLE



• Acepromazine maleate

Product identification

Medicine name:

CALMIVET SOLUTION INJECTABLE

Active substance:

Acepromazine maleate

Target species:

Dog

Cat

Horse

Route of administration:

Intramuscular use Intravenous use

Product details

Active substance and strength:

Acepromazine maleate 6.80 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- . Dog
- . Cat
- . Horse
 - Milk. no withdrawal period No withdrawal period
 - Meat and offal. no withdrawal period $$\operatorname{\textsc{No}}$$ No withdrawal period

Intravenous use:

- . Dog
- . Cat
- Horse
 - Milk. no withdrawal period No withdrawal period
 - Meat and offal. no withdrawal period No withdrawal period

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05AA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in French

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation date: 22/06/1992 Manufacturing sites for batch release: Vetoquinol Responsible authority: National Veterinary Medicines Agency Authorisation number: FR/V/0433192 6/1992 Date of authorisation status change: 22/06/2012 To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below. Package Leaflet	Marketing authorisation holder: Vetoquinol S.A.
Vetoquinol Responsible authority: National Veterinary Medicines Agency Authorisation number: FR/V/0433192 6/1992 Date of authorisation status change: 22/06/2012 To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	_
National Veterinary Medicines Agency Authorisation number: FR/V/0433192 6/1992 Date of authorisation status change: 22/06/2012 To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	
Date of authorisation status change: 22/06/2012 To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	
Documents Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	-
Summary of Product Characteristics This document does not exist in this language (English). You can find it in another language below.	
This document does not exist in this language (English). You can find it in another language below.	Documents
language below.	Summary of Product Characteristics
Package Leaflet	
	Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.

Source URL: https://medicines.health.europa.eu/veterinary/600000035503