

MEDERANTIL

Authorised

- Brotizolam

Product identification

Medicine name:

MEDERANTIL

Active substance:

Brotizolam

Target species:

Cattle

Route of administration:

Intravenous use

Product details

Active substance and strength:

Brotizolam

0.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Cattle

- Milk. 0 day
- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN05CD09

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

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 holder:

Boehringer Ingelheim Animal Health France

Marketing authorisation date:

15/06/1987

Manufacturing sites for batch release:

Labiana Life Sciences S.A.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/5798068 5/1987

Date of authorisation status change:

15/06/2012

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.