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GLUCANTIME SOLUTION INJECTABLE

Authorised

- Meglumine antimonate

Product identification

Medicine name:
GLUCANTIME SOLUTION INJECTABLE

Active substance:
Meglumine antimonate

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Meglumine antimonate
300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51DX01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
20/07/1992

Manufacturing sites for batch release:
Haupt Pharma Livron
Boehringer Ingelheim Animal Health France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/7396221 8/1992

Date of authorisation status change:
20/07/2012

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 and Labelling

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