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

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Package Leaflet and Labelling

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