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GLUCANTIME

Authorised

- Meglumine antimonate

Product identification

Medicine name:

GLUCANTIME

Active substance:

Meglumine antimonate

Target species:

Dog

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Meglumine antimonate

300.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Dog

- Unspecified. 0 day

Intravenous use:

-

Dog

- Unspecified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP51DX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in [Italian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Marketing authorisation date:

18/12/1995

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

18/12/1995

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

Documents

Combined File of all Documents

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