

Danilon 150 mg/g, granulato per uso orale per equidi non dpa

Authorised

- Suxibuzone

Product identification

Medicine name:

Danilon 150 mg/g, granulato per uso orale per equidi non dpa

Active substance:

Suxibuzone

Target species:

Horse (non food-producing)

Route of administration:

Oral use

Product details

Active substance and strength:

Suxibuzone

150.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Granules

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM01AA90

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in [Italian](#)

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:

Ecuphar Veterinaria S.L.U.

Marketing authorisation date:

15/03/1999

Manufacturing sites for batch release:

Esteve Pharmaceuticals S.A.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

16/09/2008

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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