DANILON 500 mg/g, granulato per uso orale per equidi non dpa

Authorised

- Suxibuzone

Product identification

Medicine name:
DANILON 500 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
500.00 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

Available 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:
11/11/2020

Manufacturing sites for batch release:
Meribel Pharma Parets S.L.

Responsible authority:
Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:
11/11/2020

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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