Danilon equidos 1.5 g Granules for horses and ponies

Not authorised

- Suxibuzone

Product identification

Medicine name:
Danilon equidos 1.5 g Granules for horses and ponies
Danilon equidos 1,5 g kyrni fyrir hesta og smáhesta (pony).

Active substance:
Suxibuzone

Target species:
Horse
Horse (pony)

Route of administration:
In-feed use

Product details

Active substance and strength:
Suxibuzone
1.50 gram(s) / 10.00 gram(s)

Pharmaceutical form:

Granules

Withdrawal period by route of administration:
In-feed use:

- **Horse**

  - Meat and offal. no withdrawal period

Not to be used in animals intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

  - Milk. no withdrawal period

Not to be used in animals intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AA90

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

Authorisation status:
Surrendered

Authorised in:
Iceland

Package description:
(ID2): 1 unspecified outer container with 60 Sachet (Paper, Aluminium, PolyEthylene) with 10 gram(s) (600 gram(s))
(ID1): 1 unspecified outer container with 18 Sachet (Paper, Aluminium, PolyEthylene) with 10 gram(s) (180 gram(s))

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Ecuphar Veterinaria S.L.U.

Marketing authorisation date:
27/05/2011

Manufacturing sites for batch release:
Esteve Pharmaceuticals S.A.

Responsible authority:
Icelandic Medicines Agency

Authorisation number:
IS/2/11/016/01

Date of authorisation status change:
1/05/2022

Reference member state:
Germany

Procedure number:
DE/V/0192/001

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

Documents

Summary of Product Characteristics

Combined File of all Documents

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