Meranox 25 mg/ml Oral Suspension for Pigeons

Authorised

- Fenbendazole

Product identification

Medicine name:
Meranox 25 mg/ml Oral Suspension for Pigeons

Active substance:
Fenbendazole

Target species:
Homing pigeon

Route of administration:
Oral use

Product details

Active substance and strength:
Fenbendazole
25.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AC13

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Type III amber glass bottles of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, including a 1 ml and a 5 ml dosing syringe. Carton box with 10 x 50 ml.
Type III amber glass bottles of 10 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, includng a 1 ml dosing syringe. Carton box with 1 x 10 ml.
Type III amber glass bottles of 10 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, includng a 1 ml dosing syringe. Carton box with 10 x 10 ml.
Type III amber glass bottles of 30 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, including a 1 ml and a 5 ml dosing syringe. Carton box with 1 x 30 ml.
Type III amber glass bottles of 30 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, including a 1 ml and a 5 ml dosing syringe. Carton box with 10 x 30 ml.
Type III amber glass bottles of 50 ml closed with tamper-evident HDPE screw caps with ring and colourless LDPE syringe insert, including a 1 ml and a 5 ml dosing syringe. Carton box with 1 x 50 ml.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:

Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Avimedical B.V.

Marketing authorisation date:
11/05/2017

Manufacturing sites for batch release:
Floris Veterinaire Producten B.V.

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402271.00.00

Date of authorisation status change:
23/07/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0299/001

Concerned member states:
Belgium Germany Luxembourg United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Combined File of all Documents