

Zerofen 222.2 mg/g Granules.

Authorised

- Fenbendazole

Product identification

Medicine name:

Zerofen 222.2 mg/g Granules.

Active substance:

Fenbendazole

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Fenbendazole

22.22 gram(s) / 100.00 gram(s)

Pharmaceutical form:

Granules

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AC13

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Package description:

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 1 g sachets. Sachets containing 1 g are available in the following presentations: 100 x 1 g

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 5 g sachets.

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 4 g sachets. Sachets containing 4 g are available in the following presentations: 4 x 4 g.

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 2 g sachets. Sachets containing 2 g are available in the following presentations: 120 x 2 g .

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 2 g sachets. Sachets containing 2 g are available in the following presentations: 70 x 2 g

Foil paper sachets composed of 45 or 50 gsm paper/10 gsm polyethylene/8 um foil/25 gsm polyethylene. The product is packed in 1 g sachets. Sachets containing 1 g are available in the following presentations: 120 x 1 g

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Informed consent (abridged application) - Council Directive 81/851/EEC

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

15/03/1996

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10987/140/001

Date of authorisation status change:

15/03/1996

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

Documents

Summary of Product Characteristics

ie-puar-np-600000064247-zerofen-22--granules-en.pdf