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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