Troscan 100 mg film coated tablet for Dogs

Authorised

- Nitroscanate

Product identification

Medicine name:
Troscan 100 mg film coated tablet for Dogs

Active substance:
Nitroscanate

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Nitroscanate
100.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP52AX01

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
1. Aluminium foil strips in outer carton.or2. Aluminium foil blister containing:- Lidding foil: 20 micron hard tempered aluminium foil - one side coated with heatseal lacquer and one side primed forprinting.- Blister film: Cold formable Aluminium Bottom foil oPA/Alu/PVC - 25/45/60 micron.100 tablets (for veterinary surgeons only)
1. Aluminium foil strips in outer carton.or2. Aluminium foil blister containing:- Lidding foil: 20 micron hard tempered aluminium foil - one side coated with heatseal lacquer and one side primed forprinting.- Blister film: Cold formable Aluminium Bottom foil oPA/Alu/PVC - 25/45/60 micron.1 x 6 tablets
1. Aluminium foil strips in outer carton.or2. Aluminium foil blister containing:- Lidding foil: 20 micron hard tempered aluminium foil - one side coated with heatseal lacquer and one side primed forprinting.- Blister film: Cold formable Aluminium Bottom foil oPA/Alu/PVC - 25/45/60 micron.1 x 4 tablets

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:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:

10/02/1994

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10987/141/001

Date of authorisation status change:
10/02/1994

Reference member state:
Ireland

Procedure number:
IE/V/0147/001

Concerned member states:
France

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