Clavucill 40mg/10mg, tablets for dogs and cats

Authorised

- Potassium clavulanate
- Amoxicillin trihydrate

Product identification

Medicine name:
Clavucill 40mg/10mg, tablets for dogs and cats

Active substance:
Potassium clavulanate
Amoxicillin trihydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium clavulanate
18.35 milligram(s) / 1.00 Tablet

Amoxicillin trihydrate
45.92 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CR02

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

Authorisation status:
Valid

Authorised in:
Ireland

Package description:
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 10 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 250 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 100 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 80 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 50 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 30 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 20 Tablets.
The tablets are packed in aluminium foil strips consisting of polyester/Alu/low density polyethylene.Pack sizes: Tablet strips, packed in carton boxes of 500 Tablets.

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:
V.M.D.

Marketing authorisation date:
22/05/2015

Manufacturing sites for batch release:
V.M.D.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10817/002/001

Date of authorisation status change:
22/05/2015

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

Documents

Summary of Product Characteristics