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Epityl 60mg Flavoured Tablets for Dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Epityl 60mg Flavoured Tablets for Dogs

Active substance:
Phenobarbital

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Phenobarbital
60.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Chewable tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN03AA02

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

Authorisation status:
Valid

Authorised in:
Germany

Package description:
White HDPE containers with a polypropylene child resistant cap containing 500 tablets.
White HDPE containers with a polypropylene child resistant cap containing 100 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 1000 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 500 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 100 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 90 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 80 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 70 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 60 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 50 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 40 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 30 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 20 tablets.
Blister strips (PVC/Aluminium) containing 10 tablets in cartons of 10 tablets.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Chanelle Pharmaceuticals Manufacturing Limited

Marketing authorisation date:
13/08/2013

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
401670.00.00

Date of authorisation status change:
12/12/2018

Reference member state:
Ireland

Procedure number:
IE/V/0625/001

Concerned member states:
Austria Belgium Bulgaria Cyprus Czechia Denmark Finland France Germany Greece Italy Netherlands Poland Portugal Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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Summary of Product Characteristics

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