Epityl 60mg Flavoured Tablets for Dogs

Authorised

- Phenobarbital

Product identification

Medicine name:
Epityl 60mg Flavoured Tablets for Dogs
Epityl 60 mg okusne tablete za pse

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:
Slovenia

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:
8/09/2023

Manufacturing sites for batch release:
Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:
Agency For Medicinal Products And Medical Devices Of The Republic Of Slovenia

Authorisation number:
MR/V/0788/001

Date of authorisation status change:
8/09/2023

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)

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 26/01/2026
Download

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

Labelling

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

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