Carpcoat 40 mg film-coated tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Carpcoat 40 mg film-coated tablets for dogs

Active substance:
Carprofen

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Carprofen
40.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AE91

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

Authorisation status:
Valid

Authorised in:
Romania

Package description:
(ID12) 250 Film-coated tablet: Box (cardboard) with 25 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID8) 80 Film-coated tablet: Box (cardboard) with 8 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID6) 60 Film-coated tablet: Box (cardboard) with 6 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID2) 20 Film-coated tablet: Box (cardboard) with 2 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID11) 120 Film-coated tablet: Box (cardboard) with 12 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID9) 90 Film-coated tablet: Box (cardboard) with 9 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID10) 100 Film-coated tablet: Box (cardboard) with 10 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID5) 50 Film-coated tablet: Box (cardboard) with 5 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)

(ID1) 10 Film-coated tablet: Box (cardboard) with 1 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) with 10 Film-coated tablet, closed with Foil (aluminium)
(ID4) 40 Film-coated tablet: Box (cardboard) with 4 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID3) 30 Film-coated tablet: Box (cardboard) with 3 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID7) 70 Film-coated tablet: Box (cardboard) with 7 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
7/04/2026

Manufacturing sites for batch release:
Alfasan Nederland B.V.
Lelypharma B.V.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
260060

Date of authorisation status change:

7/04/2026

Reference member state:
Germany

Procedure number:
DE/V/0351/003

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland France Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway Poland Portugal Romania Slovakia Slovenia Spain Sweden United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

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