Carpcoat 20 mg film-coated tablets for dogs

Authorised

- Carprofen

Product identification

Medicine name:
Carpcoat 20 mg film-coated tablets for dogs
Carpcoat 20 mg comprimidos revestidos por película para cães

Active substance:
Carprofen

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Carprofen
20.00 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:
Portugal

Package description:
(ID2) 20 Film-coated tablet: Box (board) with 2 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID1) 10 Film-coated tablet: Box (board) with 1 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) with 10 Film-coated tablet, closed with Foil (aluminium)
(ID3) 30 Film-coated tablet: Box (board) with 3 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID11) 120 Film-coated tablet: Box (board) with 12 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID12) 250 Film-coated tablet: Box (board) with 25 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID9) 90 Film-coated tablet: Box (board) with 9 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID8) 80 Film-coated tablet: Box (board) with 8 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID5) 50 Film-coated tablet: Box (board) with 5 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)

(ID4) 40 Film-coated tablet: Box (board) with 4 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID6) 60 Film-coated tablet: Box (board) with 6 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID7) 70 Film-coated tablet: Box (board) with 7 Blister (polyvinyl chloride; polyethylene; polyvinylidene chloride) each with 10 Film-coated tablet, closed with Foil (aluminium)
(ID10) 100 Film-coated tablet: Box (board) with 10 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:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
27/10/2025

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

Responsible authority:
Directorate General For Food And Veterinary

Authorisation number:
1745/02/25DFVPT

Date of authorisation status change:

27/10/2025

Reference member state:
Germany

Procedure number:
DE/V/0351/002

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)

Generic of:
600000073243

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Documents

Combined File of all Documents

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