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Amoxitab 50 mg tablets for dogs and cats

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:
Amoxitab 50 mg tablets for dogs and cats

Active substance:
Amoxicillin trihydrate

Target species:
Dog
Cat

Route of administration:
Oral use

Product details

Active substance and strength:
Amoxicillin trihydrate
57.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01CA04

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID9) 90 Tablet: unspecified outer container with 9 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID8) 80 Tablet: unspecified outer container with 8 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID7) 70 Tablet: unspecified outer container with 7 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID6) 60 Tablet: unspecified outer container with 6 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID5) 50 Tablet: unspecified outer container with 5 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID4) 40 Tablet: unspecified outer container with 4 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID3) 30 Tablet: unspecified outer container with 3 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID2) 20 Tablet: unspecified outer container with 2 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID13) 100 Tablet: unspecified outer container with 10 unspecified outer container each with 1 Blister (aluminium) with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID12) 500 Tablet: unspecified outer container with 50 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID11) 250 Tablet: unspecified outer container with 25 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

(ID10) 100 Tablet: unspecified outer container with 10 Blister (aluminium) each with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)
(ID1) 10 Tablet: unspecified outer container with 1 Blister (aluminium) with 10 Tablet, closed with Foil (polyvinyl chloride, polyethylene, polyvinylidene chloride)

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:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
12/12/2016

Manufacturing sites for batch release:
Lelypharma B.V.
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
402365.00.00

Date of authorisation status change:
19/03/2021

Reference member state:
Germany

Procedure number:
DE/V/0177/001

Concerned member states:

Sweden

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

Documents

Combined File of all Documents

English (PDF)
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