

Amoxibactin 250 mg tablets for dogs

Authorised

- Amoxicillin trihydrate

Product identification

Medicine name:

Amoxibactin 250 mg tablets for dogs
Amoxibactin 250 mg Comprimé

Active substance:

Amoxicillin trihydrate

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Amoxicillin trihydrate
287.50 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:

Oral use:

- Dog

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Luxembourg

Package description:

Cardboard box of 9 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 8 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 7 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 6 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 25 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets.
Cardboard box of 50 Aluminium - PVC/PE/PVDC blisters of 10 tablets
Cardboard box of 5 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box containing 10 separate cardboard boxes, each containing 1 blister of 10 tablets

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:

Le Vet. Beheer B.V.

Marketing authorisation date:

9/12/2015

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Ministere De La Sante Division De La Pharmacie Et Des Medicaments

Authorisation number:

V/333/16/01/1489

Date of authorisation status change:

9/12/2015

Reference member state:

Netherlands

Procedure number:

NL/V/0186/002

Concerned member states:

Austria Belgium Cyprus Czechia Denmark Estonia Finland France Germany
Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Norway
Poland Portugal Slovakia Spain United Kingdom (Northern Ireland)

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Documents

Summary of Product Characteristics

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