

Tsefalen 500 mg film-coated tablets for dogs

Authorised

- Cefalexin

Product identification

Medicine name:

Tsefalen 500 mg film-coated tablets for dogs

Tsefalen 500 mg compresse rivestite con film per cani

Active substance:

Cefalexin

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Cefalexin

500.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Carton box containing 9 PVC/Aluminium blister pack of 12 tablets, with a total of 108 tablets

Carton box containing 3 PVC/Aluminium blister pack of 12 tablets, with a total of 36 tablets

Carton box containing 1 PVC/Aluminium blister pack of 12 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:

Nextmune Italy S.r.l.

Marketing authorisation date:

14/12/2012

Manufacturing sites for batch release:

Acs Dobfar S.p.A.

Responsible authority:

Ministry Of Health

Authorisation number:

104441

Date of authorisation status change:

1/06/2017

Reference member state:

Italy

Procedure number:

IT/V/0125/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Denmark Estonia Finland France
Germany Greece Hungary Ireland Latvia Liechtenstein Lithuania
Luxembourg Malta Netherlands Norway Poland Portugal Romania Slovakia
Slovenia Spain Sweden United Kingdom (Northern Ireland)

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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