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ARIXIL VET 20 MG FILM-COATED TABLET FOR DOGS

Authorised

- Benazepril hydrochloride

Product identification

Medicine name:

ARIXIL VET 20 MG FILM-COATED TABLET FOR DOGS

Active substance:

Benazepril hydrochloride

Target species:

Dog

Route of administration:

Oral use

Product details

Active substance and strength:

Benazepril hydrochloride

20.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Film-coated tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QC09AA07

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Greece

Package description:

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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:

Industrial Veterinaria S.A.

Marketing authorisation date:

27/01/2019

Manufacturing sites for batch release:

aniMedica GmbH

Industrial Veterinaria S.A.

Responsible authority:

National Organization For Medicines

Authorisation number:

8945/28-01-2019/K-0230802

Date of authorisation status change:

27/01/2019

Reference member state:

France

Procedure number:

FR/V/0330/002

Concerned member states:

Austria Finland Germany Greece Ireland Italy Poland Portugal Spain
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

eu-puar-frv0330002-mr-rpe469-en.pdf