АЛЕРГОВЕТ 10% ИНЖЕКЦИОНЕН РАЗТВОР

Authorised

- Diphenhydramine hydrochloride

Product identification

Medicine name:
АЛЕРГОВЕТ 10% ИНЖЕКЦИОНЕН РАЗТВОР

Active substance:
Diphenhydramine hydrochloride

Target species:
Dog
Horse

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Diphenhydramine hydrochloride
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QR06AA02

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Biovet AD

Marketing authorisation date:
10/12/2008

Manufacturing sites for batch release:
Biovet AD

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2151

Date of authorisation status change:
6/03/2023

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.