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Anarthron 100 mg/ml Solution for Injection

Authorised

- Pentosan polysulfate sodium

Product identification

Medicine name:
Anarthron 100 mg/ml Solution for Injection

Active substance:
Pentosan polysulfate sodium

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Pentosan polysulfate sodium
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AX90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
A 10 ml Ph.Eur. Type 1 clear glass vial fitted with a 20 mm rubber stopper Ph.Eur and closed by a plastic flip off seal attached to an aluminium seal.

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Maperath Herbal Limited

Marketing authorisation date:
20/04/2007

Manufacturing sites for batch release:
Eurovet Animal Health B.V.
Forte Healthcare Limited

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 10420

Date of authorisation status change:
29/03/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0355/001

Concerned member states:
Austria Belgium Cyprus Czechia France Greece Hungary Iceland Italy Norway Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

English (PDF)
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Updated on: 13/03/2026
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