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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:
Norway

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:
5/04/2011

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

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
10-7650

Date of authorisation status change:
1/05/2012

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)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

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