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Oapent 100 mg/ml Solution for injection for dogs

Authorised

- Pentosan polysulfate sodium

Product identification

Medicine name:
Oapent 100 mg/ml Solution for injection for dogs

Active substance:
Pentosan polysulfate sodium

Target species:
Dog

Route of administration:
Solution for injection

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:
Ireland

Package description:
20 ml Clear glass vial (Ph. Eur. type I) closed with rubber stopper. The vial is sealed with aluminium cap equipped with a polypropylene “flip-off” seal, packed in a cardboard box
10 ml Clear glass vial (Ph. Eur. type I) closed with rubber stopper. The vial is sealed with aluminium cap equipped with a polypropylene “flip-off” seal, packed in a cardboard box

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Vetpharma Animal Health S.L.

Marketing authorisation date:
5/06/2026

Manufacturing sites for batch release:
Labiana Life Sciences S.A.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10516/026/001

Date of authorisation status change:
5/06/2026

Reference member state:
Netherlands

Procedure number:
NL/V/0403/001

Concerned member states:
Denmark Germany Greece Ireland Italy Portugal Spain United Kingdom (Northern Ireland)

Generic of:
600000090038

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

Documents

Combined File of all Documents

NLV0403001DC Oapent PUAR -final.pdf