Cartrophen Vet 100 mg/ml Solution for Injection.

Authorised

- Pentosan polysulfate sodium

Product identification

Medicine name:
Cartrophen Vet 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

Withdrawal period by route of administration:

Subcutaneous use:

- **Dog**

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:
A clear, colourless to slightly yellow, aqueous solution contained in a 10 ml Ph.Eur. Type I clear glass vial fitted with a 20 mmbromobutyl rubber stopper 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 (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Maperath Herbal Limited

Marketing authorisation date:
1/10/1991

Manufacturing sites for batch release:
Arthropharm (Ireland) Limited
Eurovet Animal Health B.V.

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22748/001/001

Date of authorisation status change:
1/10/1991

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

Documents

Summary of Product Characteristics
