

Clevor 30 mg/ml - Eye drops, solution

Authorised

- Ropinirole hydrochloride

Product identification

Medicine name:

Clevor 30 mg/ml - Eye drops, solution

Active substance:

Ropinirole hydrochloride

Target species:

Dog

Route of administration:

Ocular use

Ocular use

Product details

Active substance and strength:

Ropinirole hydrochloride

34.20 milligram(s) / 1.00 Container

Pharmaceutical form:

Eye drops, solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN04BC04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Hungary , Ireland , Italy , Latvia , Lithuania , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden

Package description:

Packaging:Single-dose container (LDPE), Package_size:3 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:6 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:5 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:4 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:2 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:1 single-dose container,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:10 single-dose containers,
Content:0.6 ml

Packaging:Single-dose container (LDPE), Package_size:8 single-dose containers,
Content:0.6 ml

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:

Orion Corporation

Marketing authorisation date:

13/04/2018

Manufacturing sites for batch release:

Orion Corporation

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

4/02/2020

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

Documents

Combined File of all Documents

English (PDF)

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