Neptra 16.7 mg/ml florfenicol + 16.7 mg/ml terbinafine hydrochloride + 2.2 mg/ml mometasone furoate - Ear drops, solution in single-dose container

Authorised

- Florfenicol
- Terbinafine hydrochloride
- Mometasone furoate

Product identification

Medicine name:
Neptra 16.7 mg/ml florfenicol + 16.7 mg/ml terbinafine hydrochloride + 2.2 mg/ml mometasone furoate - Ear drops, solution in single-dose container

Active substance:
Florfenicol

Terbinafine hydrochloride

Mometasone furoate

Target species:
Dog

Route of administration:
Auricular use

Product details

Active substance and strength:

Florfenicol
16.70 milligram(s) / 1.00 millilitre(s)

Terbinafine hydrochloride
16.70 milligram(s) / 1.00 millilitre(s)

Mometasone furoate
2.20 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Ear drops, solution in single-dose container

Withdrawal period by route of administration:

Auricular use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QS02CA91

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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

Package description:
Packaging:Tube (LDPE/Alu/LDPE), Package_size:1 tube, Content:1 ml
Packaging:Tube (LDPE/Alu/LDPE), Package_size:20 tube, Content:1 ml
Packaging:Tube (LDPE/Alu/LDPE), Package_size:10 tube, Content:1 ml
Packaging:Tube (LDPE/Alu/LDPE), Package_size:2 tube, Content:1 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Elanco Animal Health GmbH

Marketing authorisation date:
10/12/2019

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
26/04/2021

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

Documents

Combined File of all Documents

English (PDF)
Published on: 18/09/2024
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