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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

- Florfenicol
- Terbinafine hydrochloride
- Mometasone furoate

Authorised

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

- QS02CA91

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

- 10/12/2019

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

Documents

Product information

Combined File of all Documents

English (PDF)
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