Easotic 1.11 mg/ml + 1505 IU/ml + 15.1 mg/ml - Ear drops, suspension

Authorised

- Hydrocortisone aceponate
- Miconazole nitrate
- Gentamicin sulfate

Product identification

Medicine name:
Easotic 1.11 mg/ml + 1505 IU/ml + 15.1 mg/ml - Ear drops, suspension

Active substance:
Hydrocortisone aceponate

Miconazole nitrate

Gentamicin sulfate

Target species:
Dog

Route of administration:
Auricular use

Product details

Active substance and strength:
Hydrocortisone aceponate

Presentation_strength:1.11 mg Reference:Hse Index:0

Miconazole nitrate

Presentation_strength:15.1 mg Reference:Ph Eur Index:1

Gentamicin sulfate

Presentation_strength:1505 IU Reference:Ph Eur Index:2

Pharmaceutical form:

Ear drops, suspension

Withdrawal period by route of administration:

Auricular use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QS02CA03

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:

Cyprus , Denmark , Estonia , Hungary , Iceland , Ireland , Italy , Latvia , Lithuania , Norway , Portugal , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Pipette (HDPE), Package_size:200 pipettes, Content:1 ml

Packaging:Pipette (HDPE), Package_size:100 pipettes, Content:1 ml

Packaging:Pipette (HDPE), Package_size:50 pipettes, Content:1 ml

Packaging:Pipette (HDPE), Package_size:10 pipettes, Content:1 ml

Packaging:Pipette (HDPE), Package_size:5 pipettes, Content:1 ml

Packaging:Container (PP), Package_size:1 container, Content:10 ml

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:
Virbac S.A.

Marketing authorisation date:
20/11/2008

Manufacturing sites for batch release:
VIRBAC

Responsible authority:
European Commission

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

Date of authorisation status change:
20/11/2008

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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