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Posatex 8.5 mg/ml orbifloxacin + 0.9 mg/ml mometasone furoate + 0.9 mg/ml posaconazole - Ear drops, suspension

Authorised

- Mometasone furoate
- Posaconazole
- Orbifloxacin

Product identification

Medicine name:
Posatex 8.5 mg/ml orbifloxacin + 0.9 mg/ml mometasone furoate + 0.9 mg/ml posaconazole - Ear drops, suspension

Active substance:
Mometasone furoate

Posaconazole

Orbifloxacin

Target species:
Dog

Route of administration:
Auricular use

Product details

Active substance and strength:

Mometasone furoate
0.86 milligram(s) / 1.00 Bottle

Posaconazole
0.86 milligram(s) / 1.00 Bottle

Orbifloxacin
8.55 milligram(s) / 1.00 Bottle

Pharmaceutical form:

Ear drops, suspension

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 , Bulgaria , Cyprus , Czechia , France , Germany , Greece , Hungary , Ireland , Italy , Luxembourg , Norway , Poland , Portugal , Romania , Slovakia , Spain , United Kingdom (Northern Ireland)

Package description:

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:35.1 ml
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:17.5 ml
Packaging:Bottle (HDPE), Package_size:1 bottle, Content:8.8 ml

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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:
Intervet International B.V.

Marketing authorisation date:
23/06/2008

Manufacturing sites for batch release:
Vet Pharma Friesoythe GmbH

Responsible authority:
European Commission

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

Date of authorisation status change:
23/06/2008

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: 14/03/2026

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