

Imoxat 80 mg + 8 mg - Spot-on solution

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Imoxat 80 mg + 8 mg - Spot-on solution

Active substance:

Moxidectin

Imidacloprid

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Moxidectin

Presentation_strength:8 mg Reference:Hse Index:0

Imidacloprid

Presentation_strength:80 mg Reference:Hse Index:1

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB52

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)

Package description:

Packaging:Pipette (PP/COC/EVOH/PP), Package_size:3 pipettes, Content:0.8 ml

Packaging:Pipette (PP/COC/EVOH/PP), Package_size:1 pipette, Content:0.8 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Chanelle Pharmaceuticals Manufacturing Ltd

Marketing authorisation date:

7/12/2021

Manufacturing sites for batch release:

Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

26/01/2023

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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