Imoxat 400 mg + 100 mg - Spoton solution

Authorised

- Moxidectin
- Imidacloprid

Product identification

Medicine name:

Imoxat 400 mg + 100 mg - Spot-on solution

Active substance:

Moxidectin

Imidacloprid

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Moxidectin

Presentation strength:100 mg Reference:Hse Index:0

Imidacloprid

Presentation strength:400 mg Reference:Hse Index:1

Pharmaceutical form:

Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

. Dog

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:4 ml Packaging:Pipette (PP/COC/EVOH/PP), Package size:1 pipette, Content:4 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:
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Chanelle Pharmaceuticals Manufacturing Limited

Responsible authority:

European Commission

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

7/12/2021

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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