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Ataxxa 2000 mg/400 mg spot-on solution for dogs over 25 kg

Authorised

- Permethrin
- Imidacloprid

Product identification

Medicine name:
Ataxxa 2000 mg/400 mg spot-on solution for dogs over 25 kg

Active substance:
Permethrin

Imidacloprid

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Permethrin
2000.00 milligram(s) / 1.00 Pipette

Imidacloprid

400.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC54

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Latvia

Available in:
Latvia

Package description:
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
Available only in Latvian
White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 4 ml of solutionBox containing 1 pipettes.
White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 4 ml of solutionBox containing 3 pipettes.
White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 4 ml of solutionBox containing 4 pipettes.
White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 4 ml of solutionBox containing 6 pipettes.
White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex

bag.6 ml pipette containing 4 ml of solutionBox containing 10 pipettes.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
29/09/2015

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Food And Veterinary Service

Authorisation number:
V/DCP/15/0038

Date of authorisation status change:
29/09/2015

Reference member state:
Ireland

Procedure number:
IE/V/0439/004

Concerned member states:
Austria Belgium Bulgaria Croatia Czechia Estonia France Germany Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Slovenia Spain United Kingdom (Northern Ireland)

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Documents

Combined File of all Documents

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Summary of Product Characteristics