

Ataxxa 500 mg/100 mg spot-on solution for dogs over 4 kg up to 10 kg

Authorised

- Permethrin
- Imidacloprid

Product identification

Medicine name:

Ataxxa 500 mg/100 mg spot-on solution for dogs over 4 kg up to 10 kg

Ataxxa 500 mg - 100 mg Spot-on oplossing

Ataxxa 500 mg - 100 mg Solution pour spot-on

Ataxxa 500 mg - 100 mg Lösung zum Auftropfen

Active substance:

Permethrin

Imidacloprid

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Permethrin

500.00 milligram(s) / 1.00 Pipette

Imidacloprid

100.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 10 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 6 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 4 pipettes.

White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex

bag.3 ml pipette containing 1 ml of solution.Box containing 3 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 1 pipette.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 10 pipettes.

White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 6 pipettes.

White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 3 ml pipette containing 1 ml of solution. Box containing 4 pipettes.

White polypropylene pipette closed with a polyoxymethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.3 ml pipette containing 1 ml of solution.Box containing 1 pipette.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 3 ml pipette containing 1 ml of solution. Box containing 3 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:

9/03/2016

Manufacturing sites for batch release:

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V490986

Date of authorisation status change:

9/03/2016

Reference member state:

Ireland

Procedure number:

IE/V/0439/002

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)

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)

Published on: 30/06/2024

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Package Leaflet

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Labelling

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