

Ataxxa 1250 mg/250 mg spot-on solution for dogs over 10 kg up to 25 kg

Authorised

- Permethrin
- Imidacloprid

Product identification

Medicine name:

Ataxxa 1250 mg/250 mg spot-on solution for dogs over 10 kg up to 25 kg

Active substance:

Permethrin

Imidacloprid

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Permethrin

1250.00 milligram(s) / 1.00 Pipette

Imidacloprid
250.00 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:

France

Available in:

France

Package description:

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 2.5 ml of solutionBox containing 1 pipette.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 2.5 ml of solutionBox containing 3 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 2.5 ml of solutionBox containing 4 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 2.5 ml of solutionBox containing 6 pipettes.

White polypropylene pipette closed with a polyethylene cap. Each pipette is packed in apolyethylene terephthalate/aluminium/low density polyethylene triplex bag.6 ml pipette containing 2.5 ml of solutionBox containing 10 pipettes.

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

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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:

9/09/2015

Manufacturing sites for batch release:

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/7319388 1/2015

Date of authorisation status change:

23/09/2020

Reference member state:

Ireland

Procedure number:

IE/V/0439/003

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

Updated on: 13/03/2026

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Package Leaflet and Labelling

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