

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

Authorised

- Imidacloprid
- Permethrin

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 voor honden met een gewicht van 4 kg tot 10 kg

Active substance:

Imidacloprid

Permethrin

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Imidacloprid

100.00 milligram(s) / 1.00 Pipette

Permethrin

500.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Withdrawal period by route of administration:

Spot-on use:

- Dog
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Package description:

White polypropylene pipette closed with either a polyethylene or 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 either a polyethylene or 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 either a polyethylene or 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 4 pipettes.

White polypropylene pipette closed with either a polyethylene or 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 either a polyethylene or 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 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:

7/10/2015

Manufacturing sites for batch release:

Krka d.d. Novo Mesto

Tad Pharma GmbH

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 115887

Date of authorisation status change:

1/02/2022

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

Combined File of all Documents

This document does not exist in this language (English). You can find it in another language below.

Summary of Product Characteristics

Source URL: <https://medicines.health.europa.eu/veterinary/600000046792>