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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:
Italy

Available in:
Italy

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

bag.6 ml pipette containing 2.5 ml of solutionBox containing 1 pipette.
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 2.5 ml of solutionBox containing 3 pipettes.
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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 2.5 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 2.5 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:
22/12/2015

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

Responsible authority:
Ministry Of Health

Authorisation number:
104800

Date of authorisation status change:
22/07/2020

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)

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Documents

Combined File of all Documents

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Summary of Product Characteristics