

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 šķīdums pilināšanai uz ādas suņiem no 4 kg līdz 10 kg

Active substance:

Permethrin

Imidacloprid

Target species:

Dog

Route of administration:

Topical 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

Withdrawal period by route of administration:

Topical 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:

Latvia

Package description:

Available only in [Latvian](#)

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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:

29/09/2015

Manufacturing sites for batch release:

Tad Pharma GmbH
Krka d.d. Novo Mesto

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/15/0036

Date of authorisation status change:

29/09/2015

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

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