

# Damtix 200 mg/40 mg spot-on solution for dogs up to 4 kg

Authorised

- Permethrin (40:60)
- Imidacloprid

## Product identification

**Medicine name:**

Damtix 200 mg/40 mg spot-on solution for dogs up to 4 kg

DAMTIX (200+40)MG/0.4ML ΔΙΑΛΥΜΑ ΓΙΑ ΕΠΙΧΥΣΗ ΣΕ ΣΗΜΕΙΟ

**Active substance:**

Permethrin (40:60)

Imidacloprid

**Target species:**

Dog

**Route of administration:**

Spot-on use

## Product details

**Active substance and strength:**

Permethrin (40:60)

200.00 milligram(s) / 1.00 Pipette

Imidacloprid  
40.00 milligram(s) / 1.00 Pipette

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**Pharmaceutical form:**

Spot-on solution

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QP53AC54

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Greece

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**Available in:**

Greece

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**Package description:**

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 1 ml pipette containing 0.4 ml of solution Box containing 10 pipettes

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 1 ml pipette containing 0.4 ml of solution Box containing 6 pipettes

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 1 ml pipette containing 0.4 ml of solution Box containing 4 pipettes

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 1 ml pipette containing 0.4 ml of solution Box containing 3 pipettes

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 1 ml pipette containing 0.4 ml of solution Box containing 1 pipette

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

KRKA tovarna zdravil d.d. Novo mesto

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**Marketing authorisation date:**

21/06/2022

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**Manufacturing sites for batch release:**

KRKA tovarna zdravil d.d. Novo mesto

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**Responsible authority:**

National Organization For Medicines

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**Authorisation number:**

67541/22-06-2022/K-0248901

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**Date of authorisation status change:**

21/06/2022

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0663/001

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**Concerned member states:**

Finland France Germany Greece Italy Portugal Spain

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Summary of Product Characteristics

Combined File of all Documents