

Damtix 2000 mg/400 mg spot-on solution for dogs over 25 kg

Authorised

- Permethrin (40:60)
- Imidacloprid

Product identification

Medicine name:

Damtix 2000 mg/400 mg spot-on solution for dogs over 25 kg

Active substance:

Permethrin (40:60)

Imidacloprid

Target species:

Dog

Route of administration:

Spot-on use

Product details

Active substance and strength:

Permethrin (40:60)

2000.00 milligram(s) / 1.00 Pipette

Imidacloprid

400.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP53AC54

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

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. 6 ml pipette containing 4.0 ml of solution Box containing 1 pipette

White polypropylene pipette closed with a polyethylene (HDPE) cap. Each pipette is packed in a polyethylene terephthalate/aluminium/low density polyethylene triplex bag. 6 ml pipette containing 4.0 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. 6 ml pipette containing 4.0 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. 6 ml pipette containing 4.0 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. 6 ml pipette containing 4.0 ml of solution Box 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:

1/04/2022

Manufacturing sites for batch release:

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10774/073/004

Date of authorisation status change:

1/04/2022

Reference member state:

Ireland

Procedure number:

IE/V/0663/004

Concerned member states:

Finland France Germany Greece Italy Portugal Spain

To consult adverse reactions on veterinary medicinal products please go to
www.adrreports.eu/vet

Documents

Summary of Product Characteristics

Combined File of all Documents