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

Authorised

This information is not available for this product.

Product identification

Medicine name:
Damtix 200 mg/40 mg spot-on solution for dogs up to 4 kg
DAMTIX 200 mg/40 mg Lösung zum Auftropfen für Hunde bis 4 kg

Active substance:
This information is not available for this product.

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
This information is not available for this product.

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:
Germany

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

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:
22/03/2022

Manufacturing sites for batch release:
Krka d.d. Novo Mesto

Responsible authority:
Federal Office Of Consumer Protection And Food Safety

Authorisation number:
V7004634.00.00

Date of authorisation status change:
22/03/2022

Reference member state:
Ireland

Procedure number:
IE/V/0663/001

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

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