

Selehold 60 mg spot-on solution for cats 7.6↕10.0 kg

Authorised

This information is not available for this product.

Product identification

Medicine name:

Selehold 60 mg spot-on solution for cats 7.6↕10.0 kg
SELEHOLD 60 mg solutie spot-on pentru pisici 7.6-10.0 kg

Active substance:

This information is not available for this product.

Target species:

Cat

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:

- Cat

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Package description:

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 3 ml pipette containing 1.0 ml of solution. Cardboard box containing 1 pipette.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 3 ml pipette containing 1.0 ml of solution. Cardboard box containing 3 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 3 ml pipette containing 1.0 ml of solution. Cardboard box containing 6 pipettes.

Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spike packaged into a laminated triplex bag composed of polyester, aluminium and polyethylene. 3 ml pipette containing 1.0 ml of solution. Cardboard box containing 15 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:

23/10/2018

Manufacturing sites for batch release:

Krka d.d. Novo Mesto

Tad Pharma GmbH

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

230160

Date of authorisation status change:

23/10/2023

Reference member state:

Ireland

Procedure number:

IE/V/0394/003

Concerned member states:

Belgium Bulgaria Croatia Czechia Estonia France Germany Greece Hungary
Italy Latvia Lithuania 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

Summary of Product Characteristics

English (PDF)

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