

File downloaded on 2026-07-10

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000048790>

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

Authorised

- Selamectin

Product identification

Medicine name:

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

Active substance:

Selamectin

Target species:

Cat

Route of administration:

Spot-on use

Product details

Active substance and strength:

Selamectin

60.00 milligram(s) / 1.00 Pipette

Pharmaceutical form:

Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AA05

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Available in:

Hungary

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 15 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 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 1 pipette.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:

16/10/2018

Manufacturing sites for batch release:

TAD Pharma GmbH

KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

4003/X/18 NÉBIH ÁTI

Date of authorisation status change:

16/10/2018

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 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

Summary of Product Characteristics