Selehold 120 mg spot-on solution for dogs 10.1–20.0 kg

Authorised

- Selamectin

Product identification

Medicine name:
Selehold 120 mg spot-on solution for dogs 10.1–20.0 kg
Selehold 120 mg разтвор за прилагане върху ограничен участък за кучета 10,1–20 kg

Active substance:
Selamectin

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Selamectin
120.00 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:
Bulgaria

Available in:
Bulgaria

Package description:
Translucent polypropylene unit-dose pipette with polyethylene or polyoxymethylene or polypropylene closure with spikepackaged 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 spikepackaged 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 spikepackaged 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 spikepackaged 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:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
KRKA tovarna zdravil d.d. Novo mesto

Marketing authorisation date:
7/11/2018

Manufacturing sites for batch release:
TAD Pharma GmbH
KRKA tovarna zdravil d.d. Novo mesto

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-2850

Date of authorisation status change:
7/11/2018

Reference member state:
Ireland

Procedure number:
IE/V/0395/003

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)

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Documents

Summary of Product Characteristics

English (PDF)
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