File downloaded on 2026-04-25
Source URL: https://medicines.health.europa.eu/veterinary/en/600000051764

SELAPRO 30 mg spot-on solution for very small dogs (2.6–5.0 kg)

Authorised

- Selamectin

Product identification

Medicine name:
SELAPRO 30 mg spot-on solution for very small dogs (2.6–5.0 kg)

Active substance:
Selamectin

Target species:
Dog

Route of administration:
Spot-on use

Product details

Active substance and strength:
Selamectin
30.00 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:
Ireland

Package description:
The veterinary medicinal product is supplied in single dose pipettes which are moulded from a film composed of 3 layers: apolypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. Thepipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film andpresented in an outer box.The products are supplied in boxes of 1 unit dose pipettes.
The veterinary medicinal product is supplied in single dose pipettes which are moulded from a film composed of 3 layers: apolypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. Thepipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film andpresented in an outer box.The products are supplied in boxes of 3 unit dose pipettes.
The veterinary medicinal product is supplied in single dose pipettes which are moulded from a film composed of 3 layers: apolypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. Thepipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film andpresented in an outer box.The products are supplied in boxes of 4 unit dose pipettes.
The veterinary medicinal product is supplied in single dose pipettes which are moulded from a film composed of 3 layers: apolypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. Thepipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film andpresented in an outer box.The products are supplied in boxes of 6 unit dose pipettes.

The veterinary medicinal product is supplied in single dose pipettes which are moulded from a film composed of 3 layers: apolypropylene/COC/polypropylene, solvent free lacquer laminate and a copolymer of polyethylene/EVOH/polyethylene. Thepipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film andpresented in an outer box.The products are supplied in boxes of 24 unit dose 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:
Norbrook Laboratories (Ireland) Limited

Marketing authorisation date:
26/10/2018

Manufacturing sites for batch release:
Norbrook Manufacturing Limited
Norbrook Laboratories Limited

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA22664/144/004

Date of authorisation status change:
26/10/2018

Reference member state:
Ireland

Procedure number:
IE/V/0391/004

Concerned member states:
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics