

Selapro 60 mg Spot-on Solution for Large Cats (7.6–10.0 kg)

Authorised

- Selamectin

Product identification

Medicine name:

SELAPRO 60 mg spot-on solution for large cats (7.6–10.0 kg)

Selapro 60 mg Spot-on Solution for Large 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:

United Kingdom (Northern 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. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented 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. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented 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. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented 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. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented 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. The pipettes are sealed within a child resistant 4-ply foil sachet composed of LDPE/nylon/aluminium foil/polyester film and presented 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 Limited

Marketing authorisation date:

23/10/2018

Manufacturing sites for batch release:

Norbrook Manufacturing Limited

Norbrook Laboratories Limited

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 02000/4429

Date of authorisation status change:

23/01/2024

Reference member state:

Ireland

Procedure number:

IE/V/0391/003

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