FRONTLINE TRI-ACT 270.4 MG / 2019.2 MG SPOT-ON SOLUTION FOR DOGS 20-40 KG

Authorised

- Fipronil
- Permethrin

Product identification

Medicine name:
FRONTLINE TRI-ACT 270.4 MG / 2019.2 MG SPOT-ON SOLUTION FOR DOGS 20-40 KG

Active substance:
Fipronil

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Fipronil
270.40 milligram(s) / 1.00 Pipette

Permethrin
2019.20 milligram(s) / 1.00 Pipette

Pharmaceutical form:
Spot-on solution

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AX65

Legal status of supply:
Veterinary medicinal product not subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Malta

Package description:
Plastic card of 1 pipette containing 4 ml
Carton box of 6 pipettes containing 4 ml each
Carton box of 3 pipettes containing 4 ml each

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
17/12/2014

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Ministry For Agriculture Fisheries And Animal Rights

Authorisation number:
VMA40

Date of authorisation status change:
17/12/2014

Reference member state:
France

Procedure number:
FR/V/0266/004

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia Germany Greece Hungary Italy Latvia Lithuania Luxembourg Malta 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

Combined File of all Documents

eu-puar-frv0266004-mr-rpe962-en.pdf