FRONTLINE TRI-ACT 33.8 MG / 252.4 MG SPOT-ON SOLUTION FOR DOGS 2- 5 KG

Authorised

- Fipronil
- Permethrin

Product identification

Medicine name:
FRONTLINE TRI-ACT 33.8 MG / 252.4 MG SPOT-ON SOLUTION FOR DOGS 2- 5 KG

Active substance:
Fipronil

Permethrin

Target species:
Dog

Route of administration:
Cutaneous use

Product details

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

Permethrin
252.40 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:
Netherlands

Package description:
Plastic card of 1 pipette containing 0,5 ml
Carton box of 6 pipettes containing 0,5 ml each
Carton box of 3 pipettes containing 0,5 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 Netherlands B.V.

Marketing authorisation date:
27/08/2014

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 113896

Date of authorisation status change:
29/03/2022

Reference member state:
France

Procedure number:
FR/V/0266/001

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

English (PDF)
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