# Galliprant 20 mg - Tablet

Authorised

- Grapiprant

## Product identification

**Medicine name:**
Galliprant 20 mg - Tablet

**Active substance:**
Grapiprant

**Target species:**
Dog

**Route of administration:**
Oral use

## Product details

**Active substance and strength:**
Grapiprant
20.00 milligram(s) / 1.00 Tablet

**Pharmaceutical form:**
Tablet

**Anatomical therapeutic chemical veterinary (ATCvet) codes:**
QM01AX

**Legal status of supply:**
Veterinary medicinal product subject to veterinary prescription

**Authorisation status:**
Valid

**Authorised in:**
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

**Available in:**
Austria , Belgium , Czechia , Denmark , Finland , France , Germany , Hungary , Ireland , Italy , Luxembourg , Netherlands , Norway , Poland , Portugal , Romania , Spain , Sweden , United Kingdom (Northern Ireland)

**Package description:**
Packaging:Bottle (HDPE), Package_size:30 tablets
Packaging:Bottle (HDPE), Package_size:7 tablets

## Additional information

**Entitlement type:**
Marketing Authorisation

**Legal basis of product authorisation:**
Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

**Marketing authorisation holder:**
Elanco GmbH

**Marketing authorisation date:**
9/01/2018

**Manufacturing sites for batch release:**
Elanco France S.A.S.

**Responsible authority:**
European Commission

**Authorisation number:**
This information is not available for this product.

**Date of authorisation status change:**
9/01/2018

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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