

# DOMOSEDAN GEL 7.6 mg/ml oromucosal gel

Authorised

- DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

## Product identification

**Medicine name:**

DOMOSEDAN GEL 7.6 mg/ml oromucosal gel

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**Active substance:**

DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

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**Target species:**

Horse

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**Route of administration:**

Sublingual use

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## Product details

**Active substance and strength:**

DETOMIDINE HYDROCHLORIDE FOR VETERINARY USE

7.60 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Oromucosal gel

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**Withdrawal period by route of administration:****Sublingual use:**

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**Horse**

- Meat and offal. 0 day
- Milk. 0 hour

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

QN05CM90

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Poland

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**Available in:**

Poland

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**Package description:**

Pre-filled, single-dose syringe enabling doses from 1.0 ml to 3.0 ml packed in outer carton. Pre-filled syringes consist of a syringe barrel (HDPE), cap (LDPE), plunger (HDPE) and a locking ring. Package sizes: 1 x 3.0 ml (1 syringe per carton)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Orion Corporation

**Marketing authorisation date:**

10/07/2009

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**Manufacturing sites for batch release:**

Orion Corporation

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**Responsible authority:**

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

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**Authorisation number:**

1907

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**Date of authorisation status change:**

10/07/2009

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0218/001

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**Concerned member states:**

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

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To consult adverse reactions on veterinary medicinal products please go to

[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Package Leaflet

This document does not exist in this language (English). You can find it in another language below.

## Labelling

This document does not exist in this language (English). You can find it in another language below.

## Summary of Product Characteristics

English (PDF)

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## Combined File of all Documents