File downloaded on 2026-11-24
Source URL: https://medicines.health.europa.eu/veterinary/en/600000030207

EMEDOG, 1 MG/ML, SOLUTION FOR INJECTION FOR DOGS

Authorised

- Apomorphine hydrochloride hemihydrate

Product identification

Medicine name:
EMEDOG, 1 MG/ML, SOLUTION FOR INJECTION FOR DOGS

Active substance:
Apomorphine hydrochloride hemihydrate

Target species:
Dog

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Apomorphine hydrochloride hemihydrate
1.17 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN04BC07

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Domes Pharma

Marketing authorisation date:
31/07/2015

Manufacturing sites for batch release:
Haupt Pharma Livron

Responsible authority:
Medicines Evaluation Board

Authorisation number:

REG NL 115812

Date of authorisation status change:
5/04/2022

Reference member state:
France

Procedure number:
FR/V/0281/001

Concerned member states:
Austria Belgium Denmark Finland Germany Greece Ireland Italy Luxembourg Netherlands Norway Poland Portugal Romania Spain Sweden 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)
Published on: 14/03/2026
Download

eu-puar-frv0281001-mr-rpe760-en.pdf