File downloaded on 2026-11-08
Source URL: https://medicines.health.europa.eu/veterinary/en/600000036533

Gujatal 100 mg/ml, solution for infusion for horses

Authorised

- Guaifenesin

Product identification

Medicine name:
Gujatal 100 mg/ml, solution for infusion for horses

Active substance:
Guaifenesin

Target species:
Horse

Route of administration:
Intravenous use

Product details

Active substance and strength:
Guaifenesin
100.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for infusion

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM03BX90

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
500 ml polypropylene bottle with a bromobutyl rubber stop and aluminium cap

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Eurovet Animal Health B.V.

Marketing authorisation date:
28/09/2017

Manufacturing sites for batch release:
Eurovet Animal Health B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 110606

Date of authorisation status change:
25/01/2022

Reference member state:
Netherlands

Procedure number:
NL/V/0162/001

Concerned member states:
France Germany 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

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