Triquest Vet, 333 mg/ml + 67 mg/ml, oral suspension for horses

Authorised

- Sulfadiazine
- Trimethoprim

Product identification

Medicine name:
Triquest Vet, 333 mg/ml + 67 mg/ml, oral suspension for horses

Active substance:
Sulfadiazine
Trimethoprim

Target species:
Horse

Route of administration:
Oral use
Oral use

Product details

Active substance and strength:
Sulfadiazine
333.00 milligram(s) / 1.00 millilitre(s)

Trimethoprim
67.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral suspension

Withdrawal period by route of administration:
Oral use:

- **Horse**
  - Meat and offal. 20 day

Oral use:

- **Horse**
  - Milk. no withdrawal period

Milk: Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01EW10

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

Authorisation status:
Valid

Authorised in:
France

Available in:
France

Package description:
225 ml: bottle (HDPE), closed with a screw cap (PP)
450 ml: bottle (HDPE), closed with a screw cap (PP)
90 ml: bottle (HDPE), closed with a screw cap (PP)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – change in strength (Article 19(1)(a) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Alfasan Nederland B.V.

Marketing authorisation date:
11/10/2024

Manufacturing sites for batch release:
Alfasan Nederland B.V.
Lelypharma B.V.

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6335206 9/2024

Date of authorisation status change:
11/10/2024

Reference member state:
Sweden

Procedure number:
SE/V/0125/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Estonia France Germany Greece Hungary Iceland Ireland Italy Latvia Lithuania Luxembourg Netherlands Norway 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

Package Leaflet and Labelling

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

Summary of Product Characteristics

English (PDF)
Published on: 13/05/2026
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Combined File of all Documents

Package Leaflet

Labelling

eu-puar-sev-0125001-dc-triquest-vet-horse-en.pdf