

Cubarmix Equi 400 mg/g and 80 mg/g oral powder for horses

Authorised

- Sulfadiazine
- Trimethoprim

Product identification

Medicine name:

Cubarmix Equi 400 mg/g and 80 mg/g oral powder for horses

Active substance:

Sulfadiazine

Trimethoprim

Target species:

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Sulfadiazine

400.00 milligram(s) / 1.00 gram(s)

Trimethoprim

80.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Oral powder

Withdrawal period by route of administration:

Oral use:

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Horse

- Meat and offal. 6 month

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Romania

Package description:

White cylindrical polypropylene container, covered with a low density polyethylene lid containing 500g of oral powder.

White cylindrical polypropylene container, covered with a low density polyethylene lid containing 800g of oral powder.

Heat-sealed, white coloured, 4-layer sachet containing 12.5 g of oral powder. 15 sachets per pack.

Heat-sealed, white coloured, 4-layer sachet containing 12.5 g of oral powder. 100 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 37.5 g of oral powder. 5 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 37.5 g of oral powder. 10 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 37.5 g of oral powder. 100 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 50 g of oral powder. 5 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 50 g of oral powder. 10 sachets per pack

Heat-sealed, white coloured, 4-layer sachet containing 50 g of oral powder. 100 sachets per pack

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:

Dopharma Research B.V.

Marketing authorisation date:

25/01/2026

Manufacturing sites for batch release:

Dopharma B.V.

Dopharma France

Responsible authority:

Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:

260020

Date of authorisation status change:

25/01/2026

Reference member state:

Ireland

Procedure number:

IE/V/0673/001

Concerned member states:

Austria Belgium Denmark France Germany Hungary Italy Netherlands
Poland 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)

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Summary of Product Characteristics

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