Source URL: https://medicines.health.europa.eu/veterinary/en/600000062692

Ditrivet 120 100 mg + 20 mg Tabletka

Authorised

- Trimethoprim
- Sulfadiazine

Product identification

Medicine name:

Ditrivet 120 100 mg + 20 mg Tabletka

Active substance:

Trimethoprim

Sulfadiazine

Target species:

Dog

Fox

Mink

Cattle

Sheep

Horse

Cat

Pig

Nutria

Route of administration:

Oral use

Product details

Active substance and strength:

Trimethoprim

20.00 milligram(s) / 1.00 Tablet

Sulfadiazine

100.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:

Tablet

Withdrawal period by route of administration:

Oral use:

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Fox

- All relevant tissues. no withdrawal period Not applicable.

•

Mink

- All relevant tissues. no withdrawal period Not applicable.

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Cattle

- Meat and offal. 14 day

•

Sheep

- Meat and offal. 14 day

•

Horse

- Meat and offal. 14 day

•

Pig

- Meat and offal. 14 day

•

Nutria

- All relevant tissues. no withdrawal period Not applicable.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01EW10

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in **Polish**

Available only in Polish

Available only in Polish

Available only in Polish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:

Biofaktor Sp. z o.o.

Marketing authorisation date:

18/03/1999

Manufacturing sites for batch release: Biofaktor Sp. z o.o.
Responsible authority: Office For Registration Of Medicinal Products Medical Devices And Biocidal Products
Authorisation number: 0911
Date of authorisation status change: 18/03/1999
To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet
Documents
Summary of Product Characteristics
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