

PARTICULARS TO APPEAR ON THE OUTER PACKAGE – CARTON

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Marbocyl P 5 mg tablet

2. STATEMENT OF ACTIVE SUBSTANCES

Each tablet contains:

Active substance:

Marbofloxacin 5mg

3. PACKAGE SIZE

Carton contains

10 tablets

20 tablets

30 tablets

40 tablets

50 tablets

100 tablets

250 tablets

4. TARGET SPECIES

Dogs and cats



5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Oral use.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

Vetoquinol UK Limited

14. MARKETING AUTHORISATION NUMBERS

Vm 08007/5024

15. BATCH NUMBER

Lot

Vetoquinol logo

**MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE
PACKAGING UNITS - BLISTERS**

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Marbocyl P 5 mg



2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Marbofloxacin 5mg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

Vetoquinol logo

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Marbocyl P 5 mg tablet for dogs and cats

2. Composition

Each tablet contains:

Active substance:

Marbofloxacin 5mg

Beige brown spotted round tablet.

3. Target species

Dogs and cats.



4. Indications for use

Infections caused by susceptible strains of organisms.

Dogs: The veterinary medicinal product is indicated in the treatment of:

- Skin and soft tissue infections (skinfold pyoderma, impetigo, folliculitis, furunculosis, cellulitis).
- Urinary tract infections (UTI) associated or not with prostatitis.
- Respiratory tract infections.

Cats: The veterinary medicinal product is indicated in the treatment of:

- Skin and soft tissue infections (wounds, abscesses).
- Upper respiratory tract infections.

5. Contraindications

Do not use in dogs aged less than 12 months, or less than 18 months for exceptionally large breeds of dogs, such as Great Danes, Briard, Bernese, Bouvier and Mastiffs, with a longer growth period.

Do not use in cats aged less than 16 weeks.

Do not use in cases of hypersensitivity to fluoroquinolones or any of the excipients.

Do not use in cases of resistance against quinolones since (almost) complete cross-resistance exists against other fluoroquinolones.

Do not use for infections resulting from strict anaerobes, yeast or fungi.

6. Special warnings

None.

Special precautions for safe use in the target species:

Fluoroquinolones have been shown to induce erosion of articular cartilage in juvenile dogs and care should be taken to dose accurately especially in young animals.

Fluoroquinolones are also known for their potential neurological side effects. Cautious use is recommended in dogs and cats diagnosed as suffering from epilepsy.

A low urinary pH could have an inhibitory effect on the activity of marbofloxacin.

Fluoroquinolones should be reserved for the treatment of clinical conditions which have responded poorly or are expected to respond poorly to other classes of antimicrobials. Whenever possible, use of fluoroquinolones should be based on susceptibility testing. Use of the product deviating from the instructions given in the SPC may increase the prevalence of bacteria resistant to the fluoroquinolones and may decrease the effectiveness of treatment with other quinolones due to the potential for cross resistance.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

People with known hypersensitivity to fluoroquinolones should avoid using this product.

In case of accidental ingestion seek medical attention and show the package leaflet or the label to the physician. Wear gloves when handling or dividing tablets. Wash hands after use.

Pregnancy:

Studies in pregnant rats and rabbits showed no side effects on pregnancy. However no specific studies have been carried out in pregnant cats and dogs.

Interaction with other medicinal products and other forms of interaction:

Fluoroquinolones are known to interact with orally administered cations (Aluminium, Calcium, Magnesium, Iron). In such cases, the bioavailability may be reduced.

Overdose:

Overdosage may cause acute signs in the form of neurological disorders, which should be treated symptomatically.

Major incompatibilities:

Not applicable.

7. Adverse events

Dogs and cats:

Rare (1 to 10 animals / 10 000 animals treated):	Vomiting ^{1,2} , soft stool ^{1,2} Hyperactivity ^{1,2,3} Excessive thirst ^{1,2} Allergic skin reaction ^{3,4} Neurological disorder (ataxia, convulsion) Joint pain Aggression Depression
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At the therapeutic recommended dosage, no severe side-effects are to be expected in dogs and cats. In particular, no lesions of the articular joints were encountered in clinical studies at the recommended dose rate

¹ Mild

² These signs cease spontaneously and do not require discontinuation of treatment

³ Transient

⁴ Due to the release of histamine

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation using the contact details at the end of this leaflet, or via your national reporting system at:

Website: <https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine>

e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Recommended dose rate:

To ensure a correct dosage body weight should be determined as accurately as possible to avoid underdosing.

2 mg/kg/day in dogs and cats in a single daily administration by oral route. The daily dose is achieved as follows:

Cats and small dogs: MARBOCYL®P 5 mg - 1 tablet per 2.5 kg bw

Duration of treatment:

Dogs:

- In skin and soft tissue infections, treatment duration is at least 5 days. Depending on the course of the disease, it may be extended up to 40 days.
- In urinary tract infections, treatment duration is at least 10 days. Depending on the course of the disease, it may be extended up to 28 days.
- In respiratory infections, treatment duration is at least 7 days and, depending on the course of the disease, it may be extended up to 21 days.

Cats:

- For skin and soft tissue infections (wounds, abscesses, phlegmons) treatment duration is 3 to 5 days.
- For upper respiratory infections treatment duration is 5 days. To ensure a correct dosage body weight should be determined as accurately as possible to avoid underdosing.

9. Advice on correct administration

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Keep out of the sight and reach of children.

This veterinary medicinal product does not require any special storage conditions.

Do not use this veterinary medicinal product after the expiry date which is stated on the label or carton after Exp. The expiry date refers to the last day of that month.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local

requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon how to dispose of medicines no longer required.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Vm 08007/5024

Aluminium /aluminium blister strips containing 10 tablets/blister.

Cardboard box:

Pack sizes of 10, 20, 30, 40, 50, 100 and 250 tablets.

Not all pack sizes may be marketed.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder and contact details to report suspected adverse reactions:

Vetoquinol UK Limited
Steadings Barn
Pury Hill Business Park
Nr. Alderton
Towcester
Northamptonshire
NN12 7LS
Tel +44 (0)1280 814 500

Manufacturer responsible for batch release:

Vetoquinol SA
Magny-Vernois
70200 Lure
France

17. Other information

POM-V

Vetoquinol logo

Gavin Hall
Approved: 21 April 2026