

## 1 NAME OF THE VETERINARY MEDICINAL PRODUCT

Milbactor 2.5 mg/25 mg tablets for small dogs and puppies weighing at least 0.5 kg

## 2 QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains:

### Active substances

Milbemycin Oxime	2.5	mg
Praziquantel	25.0	mg

### Excipients

For the full list of excipients, see section 6.1.

## 3 PHARMACEUTICAL FORM

Tablet.

Yellowish-white with brown spots, oval, biconvex tablets scored on one side.

The tablets can be divided into equal halves.

## 4 CLINICAL PARTICULARS

### 4.1 Target Species

Dogs (small dogs and puppies).

### 4.2 Indications for use, specifying the target species

In dogs : treatment of mixed infections by adult cestodes and nematodes of the following species:

- Cestodes:

*Dipylidium caninum*

*Taenia* spp.

*Echinococcus* spp.

*Mesocestoides* spp.

- Nematodes:

*Ancylostoma caninum*

*Toxocara canis*

*Toxascaris leonina*

*Trichuris vulpis*

*Crenosoma vulpis* (Reduction of the level of infection)

*Angiostrongylus vasorum* (Reduction of the level of infection by immature adult (L5) and adult parasite stages; see specific treatment and disease prevention schedules)

under section 4.9 "Amounts to be administered and administration route").

*Thelazia callipaeda* (see specific treatment schedule under section 4.9 "Amounts to be administered and administration route").

The product can also be used in the prevention of heartworm disease (*Dirofilaria immitis*), if concomitant treatment against cestodes is indicated.

### **4.3 Contraindications**

Do not use in puppies of less than 2 weeks of age and/or weighing less than 0.5 kg. Do not use in case of hypersensitivity to the active substances or to any of the excipients.

See also point Special precautions for use.

### **4.4 Special warnings for each target species**

Parasite resistance to any particular class of anthelmintic may develop following frequent, repeated use of an anthelmintic of that class.

### **4.5 Special precautions for use**

#### Special precautions for use in animals

Studies with milbemycin oxime indicate that the margin of safety in certain dogs of Collie or related breeds is less than in other breeds. In these dogs, the recommended dose should be strictly observed.

The tolerance of the product in young puppies from these breeds has not been investigated.

Clinical signs in Collies are similar to those seen in the general dog population when overdosed (see in point 4.10).

Treatment of dogs with a high number of circulating microfilariae can sometimes lead to the appearance of hypersensitivity reactions, such as pale mucous membranes, vomiting, trembling, laboured breathing or excessive salivation. These reactions are associated with the release of proteins from dead or dying microfilariae and are not a direct toxic effect of the product. The use in dogs suffering from microfilaremia is thus not recommended.

In heartworm risk-areas, or in the case it is known that a dog has been travelling to and from heartworm risk regions, before using the product, a veterinary consultation is advised to exclude the presence of any concurrent infestation of *Dirofilaria immitis*. In the case of a positive diagnosis, adulticidal therapy is indicated before administering the product.

Echinococcosis represents a hazard for humans. In case of Echinococcosis, specific guidelines on the treatment and follow up and on the safeguard of persons have to be followed. Experts or institutes of parasitology should be consulted.

No studies have been performed with severely debilitated dogs or individuals with seriously compromised kidney or liver function. The product is not recommended for such animals or only according to a benefit/risk assessment by the responsible veterinarian.

In dogs less than 4 weeks old, tape worm infection is unusual. Treatment of animals less than 4 weeks old with a combination product may therefore not be necessary.

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

Wash hands after use.

In the event of accidental ingestion of the tablets, particularly by a child, seek medical advice immediately and show the package leaflet or the label to the doctor. Part tablets should be returned to the open blister pocket and inserted into the outer carton.

#### **4.6 Adverse reactions (frequency and seriousness)**

On very rare occasions, systemic signs (such as lethargy), neurological signs (such as muscle tremors and ataxia) and/or gastrointestinal signs (such as emesis, diarrhoea, anorexia and drooling) have been observed in dogs after administration of the combination of milbemycin oxime and praziquantel.

#### **4.7 Use during pregnancy, lactation or lay**

The product may be used in breeding dogs including pregnant and lactating bitches.

#### **4.8 Interaction with other medicinal products and other forms of interaction**

No interactions were observed when the recommended dose of the macrocyclic lactone selamectin was administered during treatment with the combination of milbemycin oxime and praziquantel at the recommended dose. In the absence of further studies, caution should be taken in the case of concurrent use of the product and other macrocyclic lactones. Also, no such studies have been performed with reproducing animals.

#### 4.9 Amounts to be administered and administration route

Oral use.

Dogs should be weighed to ensure accurate dosing.

Minimum recommended dose rate: 0.5 mg of milbemycin oxime and 5 mg of praziquantel per kg are given once orally.

The product should be administered with or after some food.

Depending on the bodyweight of the dog, the practical dosing is as follows:

Body weight	Tablets
0.5 – 1 kg	1/2 tablet
>1 – 5 kg	1 tablet
>5 – 10 kg	2 tablets

In cases when heartworm disease prevention is used and at the same time treatment against tapeworm is required, the product can replace the monovalent product for the prevention of heartworm disease.

For treatment of *Angiostrongylus vasorum* infections, milbemycin oxime should be given four times at weekly intervals. It is recommended, where concomitant treatment against cestodes is indicated, to treat once with the product and continue with the monovalent product containing milbemycin oxime alone, for the remaining three weekly treatments.

In endemic areas administration of the product every four weeks will prevent angiostrongylosis by reducing immature adult (L5) and adult parasite burden, where concomitant treatment against cestodes is indicated.

For the treatment of *Thelazia callipaeda*, milbemycin oxime should be given in 2 treatments, seven days apart. Where concomitant treatment against cestodes is indicated, the product can replace the monovalent product containing milbemycin oxime alone.

#### 4.10 Overdose (symptoms, emergency procedures, antidotes), if necessary

No data available.

#### 4.11 Withdrawal period(s)

Not applicable.

## 5 PHARMACOLOGICAL or IMMUNOLOGICAL PROPERTIES

Pharmacotherapeutic group: Endectocides, Macrocyclic lactones  
ATCvet code: QP54AB51

### 5.1 Pharmacodynamic properties

Milbemycin oxime belongs to the group of macrocyclic lactones, isolated from the fermentation of *Streptomyces hygroscopicus* var. *aureolacrimosus*. It is active against mites, against larval and adult stages of nematodes as well as against larvae of *Dirofilaria immitis*.

The activity of milbemycin is related to its action on invertebrate neurotransmission: Milbemycin oxime, like avermectins and other milbemycins, increases nematode and insect membrane permeability to chloride ions via glutamate-gated chloride ion channels (related to vertebrate GABA<sub>A</sub> and glycine receptors). This leads to hyperpolarisation of the neuromuscular membrane and flaccid paralysis and death of the parasite.

Praziquantel is an acylated pyrazino-isoquinoline derivative. Praziquantel is active against cestodes and trematodes. It modifies the permeability for calcium (influx of Ca<sup>2+</sup>) in the membranes of the parasite inducing an imbalance in the membrane structures, leading to membrane depolarisation and almost instantaneous contraction of the musculature (tetany), rapid vacuolization of the syncytial tegument and subsequent tegumental disintegration (blebbing), resulting in easier expulsion from the gastrointestinal tract or death of the parasite.

### 5.2 Pharmacokinetic properties

After oral administration of praziquantel in the dog, after a small amount of food, peak serum levels of parent are rapidly attained (T<sub>max</sub> approximately 0.25-2.5 hours) and decline quickly (t<sub>1/2</sub> approximately 1 hour); there is a substantial hepatic first-pass effect, with very rapid and almost complete hepatic biotransformation, principally to monohydroxylated (also some di- and tri-hydroxylated) derivatives, which are mostly glucuronide and/or sulfate conjugated before excretion. Plasma binding is about 80%. Excretion is fast and complete (about 90% in 2 days); the principal route of elimination is renal.

After oral administration of milbemycin oxime in dogs, after a small amount of food, peak plasma levels occur at about 0.75-3.5 hours, and decline with a half-life of the unmetabolised milbemycin oxime of 1-4 days. Bioavailability is about 80%.

In the rat, metabolism appears to be complete although slow, since unchanged milbemycin oxime has not been found in urine or faeces. Main metabolites in the rat are monohydroxylated derivatives, attributable to hepatic biotransformation. In addition to relatively high liver concentrations, there is some concentration in fat, reflecting its lipophilicity.

## **6 PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

Cellulose, microcrystalline  
Lactose monohydrate  
Povidone  
Croscarmellose sodium  
Silica, colloidal anhydrous  
Meat Flavour  
Yeast powder  
Magnesium stearate

### **6.2 Incompatibilities**

Not applicable.

### **6.3 Shelf-life**

Shelf life of the veterinary medicinal product as packaged for sale: 3 years  
Shelf life for halved tablets after first opening the immediate packaging: 6 months.

### **6.4 Special precautions for storage**

Store in the original package in order to protect from moisture. This veterinary medicinal product does not require any special temperature storage conditions. Halved tablets should be stored below 25°C in the original blister and be used for the next administration.  
Keep the blister in the outer carton.

### **6.5 Nature and composition of immediate packaging**

Blister packs consisting of cold formed OPA/Al/PVC foil and aluminium foil.  
Box with 1 blister of 4 tablets.  
Box with 12 blisters, each blister contains 4 tablets.  
Not all pack sizes may be marketed.

## **6.6 Special precautions for the disposal of unused veterinary medicinal product or waste materials derived from the use of such products**

Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.

The product should not enter water courses as this may be dangerous for fish and other aquatic organisms.

## **7 MARKETING AUTHORISATION HOLDER**

Krka, d.d., Novo mesto  
Šmarješka cesta 6,  
8501 Novo mesto  
Slovenia

## **8 MARKETING AUTHORISATION NUMBER(S)**

VPA10774/030/001

## **9 DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION**

Date of first authorisation: 9<sup>th</sup> January 2015

## **10 DATE OF REVISION OF THE TEXT**

October 2017