ANNEX III

LABELLING AND PACKAGE LEAFLET

A. LABELLING

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

box with 1 blister of 2 tablets
box with 2 blisters of 2 tablets
box with 12 blisters of 2 tablets
box with 24 blisters of 2 tablets

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

MILPRO 16 mg/40 mg film-coated tablets for cats

Milbemycin oxime/Praziquantel

Broad spectrum wormer.

Cat ≥ 2 kg

2. STATEMENT OF ACTIVE AND OTHER SUBSTANCES

Each tablet contains:
milbemycin oxime 16 mg
praziquantel 40 mg

3. PHARMACEUTICAL FORM

Film-coated tablets

4. PACKAGE SIZE

2 tablets
4 tablets
24 tablets
48 tablets

5. TARGET SPECIES

Cats (weighing at least 2 kg).

6. INDICATION(S)

Read the package leaflet before use.

[optional]
[This pictogram will be removed during the national phase in Sweden]

*****Only for those countries where the product is available without prescription:*****

In cats: treatment of mixed infections by immature and adult cestodes (tapeworms) and adult nematodes (roundworms) of the following species:
Cestodes:
Echinococcus multilocularis,
Dipylidium caninum,
Taenia spp.,
Nematodes:
Ancylostoma tubaeforme,
Toxocara cati

Prevention of heartworm disease (*Dirofilaria immitis*), if concomitant treatment against cestodes is indicated.

7. METHOD AND ROUTE(S) OF ADMINISTRATION

Oral use

	
2 – 4 kg	x1/2
>4 – 8 kg	x1
>8 – 12 kg	x1½

Read the package leaflet before use.

8. WITHDRAWAL PERIOD(S)

9. SPECIAL WARNINGS, IF NECESSARY

Read the package leaflet before use.

10. EXPIRY DATE

EXP {month/year}

11. SPECIAL STORAGE CONDITIONS

Keep the blister in the outer carton.
Read the package leaflet before use.

12. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCTS OR WASTE MATERIALS, IF ANY

Disposal: Read the package leaflet.

13. THE WORDS "FOR ANIMAL TREATMENT ONLY" AND CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE, IF APPLICABLE

For animal treatment only.

14. THE WORDS "KEEP OUT OF THE SIGHT AND REACH OF CHILDREN"

Keep out of the sight and reach of children.

15. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

VIRBAC
1ère avenue 2065m LID
06516 Carros
France

16. MARKETING AUTHORISATION NUMBER(S)

Vm 05653/4184

17. MANUFACTURER'S BATCH NUMBER

Lot {number}

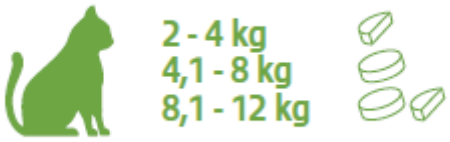
MINIMUM PARTICULARS TO APPEAR ON BLISTERS

blister of 2 tablets

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

MILPRO 16 mg/40 mg tablets for cats

Milbemycin oxime/Praziquantel



2. NAME OF THE MARKETING AUTHORISATION HOLDER

VIRBAC

3. EXPIRY DATE

EXP {month/year}

4. BATCH NUMBER

Lot {number}

5. THE WORDS "FOR ANIMAL TREATMENT ONLY"

For animal treatment only.

B. PACKAGE LEAFLET

PACKAGE LEAFLET FOR:
Milpro 16 mg/40 mg film-coated tablets for cats

1. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER AND OF THE MANUFACTURING AUTHORISATION HOLDER RESPONSIBLE FOR BATCH RELEASE, IF DIFFERENT

Marketing authorisation holder and manufacturer responsible for batch release:
VIRBAC
1ère avenue 2065m LID
06516 Carros
France

2. NAME OF THE VETERINARY MEDICINAL PRODUCT

Milpro 16 mg/40 mg film-coated tablets for cats

Milbemycin oxime, Praziquantel

3. STATEMENT OF THE ACTIVE SUBSTANCE(S) AND OTHER INGREDIENTS

Each tablet contains:
Active substances:

	Appearance	Milbemycin oxime	Praziquantel
Milpro 16 mg/40 mg film-coated tablets for cats	Oval shaped, red to pink, meat flavoured tablets with a score on both sides.	16 mg	40 mg

Excipients:

	Excipient	Quantity
Milpro 16 mg/40 mg film-coated tablets for cats	Allura red AC (E129)	0,1 mg
	Titanium Dioxide (E171)	0,5 mg

The tablets can be divided into halves.

4. INDICATION(S)

In cats: treatment of mixed infections by immature and adult cestodes (tapeworms) and adult nematodes (roundworms) of the following species:
Cestodes:

Echinococcus multilocularis,
Dipylidium caninum,
Taenia spp.,

Nematodes:
Ancylostoma tubaeforme,
Toxocara cati

Prevention of heartworm disease (*Dirofilaria immitis*) if concomitant treatment against cestodes is indicated.

5. CONTRAINDICATIONS

Milpro 16 mg/40 mg film-coated tablets for cats
Do not use in cats weighing less than 2 kg

Do not use in cases of hypersensitivity to the active substances or to any of the other ingredients.
See also point "SPECIAL WARNINGS".

6. ADVERSE REACTIONS

In very rare occasions, especially in young cats, hypersensitivity reactions, systemic signs (such as lethargy), neurological signs (such as ataxia and muscle tremors) and/or gastrointestinal signs (such as emesis and diarrhoea) may be observed after administration of the veterinary medicinal product.

The frequency of adverse reactions is defined using the following convention:
- very common (more than 1 in 10 animals treated displaying adverse reaction(s))
- common (more than 1 but less than 10 animals in 100 animals treated)
- uncommon (more than 1 but less than 10 animals in 1,000 animals treated)
- rare (more than 1 but less than 10 animals in 10,000 animals treated)
- very rare (less than 1 animal in 10,000 animals treated, including isolated reports).

If you notice any serious effects or other effects not mentioned in this package leaflet, please inform your veterinary surgeon.

Alternatively you can report via your national reporting system {national system details}.

7. TARGET SPECIES

Cats.

8. DOSAGE FOR EACH SPECIES, ROUTE(S) AND METHOD OF ADMINISTRATION

As per good veterinary practice, animals should be weighed to ensure accurate dosing.

Oral use.

Minimum recommended dose rate: 2 mg of milbemycin oxime and 5 mg of praziquantel per kg are given once orally as a single dose.

The product should be administered with or after some food.
The product is a small size tablet.
To aid with administration, the product has been coated with a meat flavour.

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

Weight	**Milpro 16 mg/40 mg film-coated tablets for cats**
0.5 - 1 kg	
> 1 – 2 kg	
2 – 4 kg	1/2 tablet
>4 – 8 kg	1 tablet
>8 – 12 kg	1 + 1/2 tablets

The product can be inserted into a programme for prevention of heartworm disease if at the same time treatment against tapeworms is indicated. The product has a duration of heartworm prevention of one month. For prevention of heartworm disease the use of a monosubstance is preferred.

9. ADVICE ON CORRECT ADMINISTRATION

Not applicable.

10. WITHDRAWAL PERIOD(S)

Not applicable.

11. SPECIAL STORAGE PRECAUTIONS

Keep out of the sight and reach of children.
This veterinary medicinal product does not require any special temperature storage conditions.
Half tablets should be stored in the original blister and be used for the next administration.
Keep the blister in the outer carton.

Do not use this veterinary medicinal product after the expiry date which is stated on the carton and blister after “EXP”. The expiry date refers to the last day of that month.

Shelf life after first opening the immediate packaging: 6 months.

12. SPECIAL WARNINGS

Special warnings for each target species:

It is recommended to treat all the animals living in the same household concomitantly.

In order to develop an effective worm control programme local epidemiological information and the living conditions of the cat should be taken into account and therefore it is recommended to seek professional advice.

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

When *D. caninum* infection is present, concomitant treatment against intermediate hosts, such as fleas and lice, should be considered to prevent re-infection.

Special precautions for use in animals:

No studies have been performed with severely debilitated cats 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.

Studies have shown that 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 the absence of data on cats with microfilaraemia, its use should be according to a benefit risk assessment by the attending veterinarian.

As the tablets are flavoured, they should be stored in a safe place out of the reach of animals.

As per good veterinary practice, animals should be weighed to ensure accurate dosing.
Ensure cats and kittens weighing between 0.5 kg and ≤2 kg receive the appropriate tablet strength (4 mg MBO/10 mg praziquantel) and the appropriate dose (1/2 or 1 tablet) for the corresponding weight band (1/2 tablet for cats weighing 0.5 to 1 kg ; 1 tablet for cats weighing >1 to 2 kg – 1 tablet).

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

Wash hands after use.

Part tablets should be returned to the open blister pack and stored in the carton.

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.

Echinococcosis represents a hazard for humans. As Echinococcosis is a notifiable disease to the World Organisation for Animal Health (OIE), specific guidelines on the treatment and follow-up, and on the safeguard of persons, need to be obtained from the relevant competent authority.

Pregnancy and lactation:
In a study, this combination of active substances was demonstrated to be well tolerated in breeding queens, including during pregnancy and lactation. As a specific study with this product has not been performed, use during pregnancy and lactation only according to a benefit/risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction:
The concurrent use of the combination praziquantel/milbemycin oxime with selamectin is well tolerated. No interactions were observed when the recommended dose of the macrocyclic lactone selamectin was administered during treatment with the combination 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.

Overdose (symptoms, emergency procedures, antidotes):
In a study conducted with the product administered at 1X, 3X and 5X the therapeutic dose, and for a duration which exceed the therapeutic indication, i.e. 3 times at 15 day-intervals, signs uncommonly reported at the recommended dose (see section 'ADVERSE REACTIONS') have been observed at 5-fold the therapeutic dose after the second and third treatments. These signs disappeared spontaneously within a day.

13. SPECIAL PRECAUTIONS FOR THE DISPOSAL OF UNUSED PRODUCT OR WASTE MATERIALS, IF ANY

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.

14. DATE ON WHICH THE PACKAGE LEAFLET WAS LAST APPROVED

April 2021

15. OTHER INFORMATION

Available pack sizes:

Milpro 16 mg/40 mg film-coated tablets for cats
Carboard box of 2 tablets containing 1 blister of 2 tablets
Carboard box of 4 tablets containing 2 blisters of 2 tablets
Carboard box of 24 tablets containing 12 blisters of 2 tablets
Carboard box of 48 tablets containing 24 blisters of 2 tablets

Not all pack sizes may be marketed.

Approved: 23/09/21