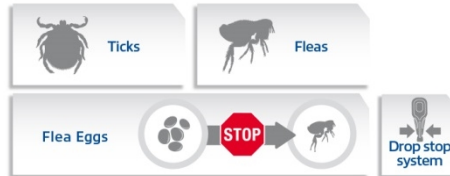


LABELLING

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Box containing 1 individual pipette placed in overblister
Box containing 4 individual pipettes placed in 2 overblisters
Box containing 24 individual pipettes placed in 12 overblisters
Box containing 60 individual pipettes placed in 30 overblisters



1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Effipro Duo 402 mg/120 mg Spot-on Solution

2. STATEMENT OF ACTIVE SUBSTANCES

Each 4.02 ml pipette contains:

Fipronil 402 mg
Pyriproxifen 120.6 mg

3. PACKAGE SIZE



1 pipette
4.02 ml



4 pipettes
4.02 ml



24 pipettes
4.02 ml



60 pipettes
4.02 ml

4. TARGET SPECIES

Dogs 40-60 kg

5. INDICATIONS

To be used against infestations with fleas alone or in association with ticks.

6. ROUTES OF ADMINISTRATION

Spot-on use.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

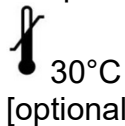
Exp. {mm/yyyy}

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 30°C.

Store in a dry place.

Keep the blister pack in the outer carton in order to protect from light.



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

Read the package leaflet before use.

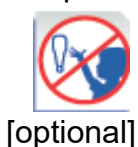


11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

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

VIRBAC

14. MARKETING AUTHORISATION NUMBERS

Vm 05653/5095

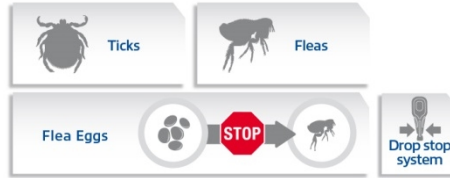
Vm 05653/3063

15. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE OUTER PACKAGE

Dispensing envelope with a capacity of either one or two 2-pipette blister(s) (to be included in large boxes only)




1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Effipro Duo 402 mg/120 mg Spot-on Solution

2. STATEMENT OF ACTIVE SUBSTANCES

Each pipette contains:

| |  [optional] | Dog | Fipronil | Pyriproxyfen |
|------------|---|----------|----------|--------------|
| [tick box] | 0.67 ml | 2-10 kg | 67 mg | 20.1 mg |
| [tick box] | 1.34 ml | 10-20 kg | 134 mg | 40.2 mg |
| [tick box] | 2.68 ml | 20-40 kg | 268 mg | 80.4 mg |
| [tick box] | 4.02 ml | 40-60 kg | 402 mg | 120.6 mg |

3. PACKAGE SIZE

This intermediate package is intended to contain at least one 2-pipette blister (and up to two 2-pipette blisters).

4. TARGET SPECIES

Dogs

5. INDICATIONS

To be used against infestations with fleas alone or in association with ticks.

6. ROUTES OF ADMINISTRATION

Spot-on use.

For dogs over 60 kg the appropriate combination of pipettes should be used.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

For lot number and expiry date, please refer to the overblister or pipette.

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 30°C.

Store in a dry place.

Keep the blister pack in the outer carton in order to protect from light.



30°C

[optional]

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

Read the package leaflet before use.



[optional]

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.



[optional]

13. NAME OF THE MARKETING AUTHORISATION HOLDER

VIRBAC

1ère avenue 2065m LID

06516 Carros

France

14. MARKETING AUTHORISATION NUMBER

Vm 05653/5095

Vm 05653/3063

15. BATCH NUMBER

For lot number and expiry date, please refer to the overblister or pipette.

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

Overblister packed in 1 pipette blister or 2 pipette blisters divisible per pipette

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

EFFIPRO DUO



2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

402 mg/120 mg
40-60 kg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS

Individual Pipette

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

EFFIPRO DUO



2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

402 mg/120 mg
40-60 kg

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp.{mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Effipro Duo 67 mg/20 mg Spot-on Solution for Small Dogs
Effipro Duo 134 mg/40 mg Spot-on Solution for Medium Dogs
Effipro Duo 268 mg/80 mg Spot-on Solution for Large Dogs
Effipro Duo 402 mg/120 mg Spot-on Solution for Very large Dogs

2. Composition

| Each pipette contains: | Active substances | | Excipients | |
|-----------------------------------|-------------------|--------------|------------|----------|
| | Fipronil | Pyriproxyfen | BHA* | BHT** |
| Pipette volume (single dose unit) | | | | |
| 0.67 ml | 67 mg | 20.1 mg | 0.134 mg | 0.067 mg |
| 1.34 ml | 134 mg | 40.2 mg | 0.268 mg | 0.134 mg |
| 2.68 ml | 268 mg | 80.4 mg | 0.536 mg | 0.268 mg |
| 4.02 ml | 402 mg | 120.6 mg | 0.804 mg | 0.402 mg |

*: Butylhydroxyanisole E320, **: Butylhydroxytoluene E321

Clear, colourless to yellowish solution.

3. Target species

Dogs

4. Indications for use

To be used against infestations with fleas alone or in association with ticks.

Against fleas:

Treatment and prevention of infestations by fleas (*Ctenocephalides felis*). One treatment prevents further infestations for 7 weeks.

Prevention of the multiplication of fleas preventing flea eggs developing into adult fleas for 12 weeks after application.

The veterinary medicinal product can be used as part of a treatment strategy for the control of Flea Allergy Dermatitis (FAD) where this has been previously diagnosed by a veterinary surgeon.

Against ticks:

Treatment of infestations by ticks (*Ixodes ricinus*).

One treatment provides persistent acaricidal efficacy for 2 weeks against *Ixodes ricinus*, and for 4 weeks against *Dermacentor reticulatus* and *Rhipicephalus sanguineus*.

If ticks of some species (*Dermacentor reticulatus*, *Rhipicephalus sanguineus*) are present at the time of application, not all ticks may be killed within 48 hours.

5. Contraindications

Do not use in rabbits, as adverse reactions and even death could occur.

Do not use in cases of known hypersensitivity to the active substances or to any of the excipients.

6. Special warnings

Special warnings:

Shampooing or immersion of the animal in water directly after treatment may reduce the duration of activity. The veterinary medicinal product remains effective against fleas for 5 weeks when the dog is shampooed at monthly intervals after treatment. If the dog requires shampooing, it is better to do so before treatment.

Water immersion repeated on two occasions post treatment did not affect adulticidal efficacy against fleas nor the efficacy related to the prevention of the development of flea eggs into adult fleas.

The influence of water immersion or shampooing of the dog on the efficacy of the veterinary medicinal product against ticks has not been evaluated.

At the beginning of the control measures, in the case of an infestation, the animal's basket, bedding and regular resting areas such as carpets and soft furnishings should be treated, with a suitable insecticide and vacuumed regularly.

To reduce environmental flea challenge, all animals living in the same household should also be treated with a suitable flea control veterinary medicinal product.

The veterinary medicinal product does not prevent ticks from attaching to animals. Transmission of infectious disease by ticks cannot be completely excluded if conditions are unfavourable. Immediate efficacy has been demonstrated against *Ixodes ricinus*, indicating that ticks of this species are likely to be killed within 48 hours of veterinary medicinal product application. If *Dermacentor reticulatus* or *Rhipicephalus sanguineus* ticks are present when the veterinary medicinal product is applied, these ticks may not be killed within the first 48 hours.

Once dead, ticks will often drop off the animal. Any remaining ticks should be carefully removed, ensuring that their mouth parts are not left within the skin.

Special precautions for safe use in the target species:

For external use only.

Animals should be weighed accurately prior to treatment.

In absence of safety data, the veterinary medicinal product should not be used in puppies less than 10 weeks old and/or weighing less than 2 kg.

Care should be taken to avoid the content of the pipette coming into contact with the eyes or mouth of the recipient dogs. In particular oral uptake due to the licking of the application site by treated or in-contact animals should be avoided.

Do not apply the veterinary medicinal product on wounds or damaged skin.

In the absence of additional safety studies, do not repeat the treatment at intervals of less than 4 weeks.

The use of the veterinary medicinal product has not been studied in sick and debilitated dogs.

Consult your veterinary surgeon before using the veterinary medicinal product if your dog is unwell or currently receiving any other veterinary treatment.

Inform your veterinary surgeon that you are using this veterinary medicinal product if s/he provides your dog with any other medication.

Seek veterinary advice if the veterinary medicinal product is accidentally ingested or comes into contact with your animal's eyes.

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

The veterinary medicinal product may cause neurotoxicity.

The veterinary medicinal product may be harmful if swallowed.

Avoid contact with skin and mouth.

Do not smoke, drink or eat during application.

In the event of accidental ingestion, seek medical advice immediately and show the package leaflet or the label to the physician.

This veterinary medicinal product can cause eye and mucous membrane irritation.

Avoid contact with skin, eye and mouth, including hand to eye contact.

In the case of accidental skin or eye contact, immediately and thoroughly flush with water.

If skin or eye irritation persists, seek medical advice and show the package leaflet or the label to the physician.

Wash hands after use.

Treated animals should not be handled until the application site is dry, and children should not be allowed to play with treated animals until the application site is dry. It is therefore recommended that animals are not treated during the day, but should be treated during the early evening, and that recently treated animals should not be allowed to sleep with owners, especially children.

Keep the pipettes in the original packaging until ready for use and dispose of used pipettes immediately.

For animal treatment only.

Special precautions for the protection of the environment:

Fipronil and pyriproxyfen may adversely affect aquatic organisms. Dogs should be prevented from accessing streams and rivers for 48-hours following treatment.

Other precautions:

The veterinary medicinal product may have adverse effects on painted, varnished or other household surfaces or furnishings. Allow the application site to dry before permitting contact with such materials.

Pregnancy and lactation:

Laboratory studies in dogs have not produced any evidence of teratogenic or embryotoxic effects. The safety of the veterinary medicinal product has not been established during pregnancy and lactation. Use only according to the benefit-risk assessment by the responsible veterinarian.

Interaction with other medicinal products and other forms of interaction:

Do not use simultaneously with other flea or tick products which are applied directly on to the animal.

Overdose:

No serious adverse effects were observed in a safety study in 10-week old puppies treated with up to 5 times the maximum recommended dose 3 times at intervals of 4 weeks and with the maximum recommended dose 6 times at intervals of 4 weeks.

The risk of experiencing adverse reactions (see section Adverse events) may however increase with overdosing, so animals should always be treated with correct pipette size according to bodyweight.

Major incompatibilities:

None known.

7. Adverse events

Dogs:

| |
|--|
| Very rare (<1 animal / 10,000 animals treated, including isolated reports): |
| Application site reaction ¹ (e.g. Application site greasy fur ² , Application site skin scaling ^{2,3} , Application site skin squamosis, Application site alopecia, Application site pruritus, Application site erythema, Application site skin discolouration) |
| Generalised itching, Alopecia |
| Hypersalivation, Vomiting |
| Neurological disorder ⁴ (e.g. Hyperaesthesia, Central nervous system depression, Neurological symptoms) |
| Respiratory signs |

¹ Transient

² Cosmetic effect

³ Slight

⁴ Reversible

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 holder or the local representative of the marketing authorisation holder 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

Dosage:

| Dog weight | Pipette volume (single dose unit) | Fipronil (mg) | Pyriproxyfen (mg) |
|------------|-----------------------------------|---------------|-------------------|
| 2-10 kg | 0.67 ml | 67 | 20.1 |
| 10-20 kg | 1.34 ml | 134 | 40.2 |
| 20-40 kg | 2.68 ml | 268 | 80.4 |
| 40-60 kg | 4.02 ml | 402 | 120.6 |

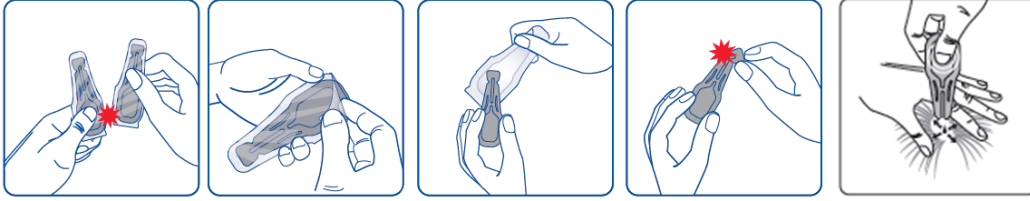
For dogs over 60 kg the appropriate combination of pipettes should be used.

Method of administration:

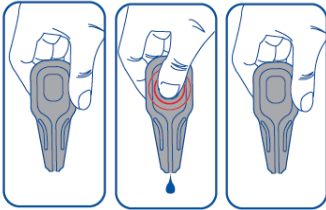
Remove the pipette from the overblister. Hold the pipette upright. Tap the narrow part of the pipette to ensure that the contents are within the main body of the pipette. Break the snap-off top of the spot-on pipette along the scored line.

Part the pet's coat on base of the neck before the shoulder blades until the skin is visible. Place the tip of the pipette directly against the skin and squeeze gently several times to

empty the contents. If necessary the contents of the pipette can be administered at one or two additional point(s) along the pet's back to avoid run-off or more superficial application to the hair coat, particularly in large dogs.



Drop stop system (the veterinary medicinal product is released only by pressing the body of the pipette).



9. Advice on correct administration

One pipette provides a single treatment, with the possibility to repeat administrations on a monthly basis.

Consult your veterinary surgeon if the veterinary medicinal product fails to control the flea and tick infestation.

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Keep out of the sight and reach of children.

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

Do not store above 30°C.

Store in a dry place.

Keep the blister pack in the outer carton in order to protect from light.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.

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

Do not contaminate ponds, waterways or ditches with the veterinary medicinal product or empty container.

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.

13. Classification of veterinary medicinal products

Veterinary medicinal product not subject to prescription.

14. MARKETING AUTHORISATION NUMBERS AND PACK SIZES

Effipro Duo 67 mg/20 mg Spot-on Solution for Small Dogs Vm 05653/5092 + 05653/3060
Effipro Duo 134 mg/40 mg Spot-on Solution for Medium Dogs Vm 05653/5093 + 05653/3061
Effipro Duo 268 mg/80 mg Spot-on Solution for Large Dogs Vm 05653/5094 + 05653/3062
Effipro Duo 402 mg/120 mg Spot-on Solution for Very Large Dogs Vm 05653/5095 + 05653/3063

Boxes of 1, 4, 24 and 60 pipettes (large boxes including envelopes intended for dispensing a reduced number of pipettes).

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 manufacturer responsible for batch release:

VIRBAC
1ère avenue 2065m LID
06516 Carros
France

Local representatives and contact details to report suspected adverse reactions:

Virbac Ltd
Suffolk, IP30 9UP –UK
Tel: +44 (0)-1359 243243

17. Other information

NFA-VPS

Pyriproxyfen is an insect growth regulator (IGR) of the class of compounds known as juvenile hormone analogues. It prevents, by contact, the emergence of adult insects by blocking the development of eggs (ovicidal effect), larvae and pupae (larvicidal effect), which are subsequently eliminated.

Combination of fipronil and pyriproxyfen provides an insecticidal and acaricidal activity against fleas (*Ctenocephalides felis*), ticks (*Rhipicephalus sanguineus*, *Dermacentor reticulatus*, *Ixodes ricinus*) in addition to preventing flea eggs developing into adult fleas.

Gavin Hall

Approved: 12 September 2025